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Kenneth Söderhäll

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Invertebrate Immunity

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DEDICATION

This volume is dedicated to all my present and former PhD students, post-docs and fellow scientists who worked in my laboratory and who have contributed to our research on invertebrate immunity.

PREFACE

The total number of animal species range up to around 1,000,000 species, of which invertebrates constitute more than 95%, including a vast diversity of organisms from unicellular protozoans to the much more complex echinoderms and protochordates. Insects are by far the largest group of animals within the invertebrates and have received a large amount of research interest due to their importance as vectors for human and animal diseases, such as malaria, and the serious harm done by insects to crops and food.

Thus a long-standing research interest in diseases, pathogens and immune responses of insects there has been. The finding by the late Professor Hans G. Boman and colleagues that *Drosophila* responded to a challenge with dead or live bacteria by the synthesis of antibacterial peptides initiated an intense research interest on the mechanism of this induction. In 1996 Jules Hoffmann, Bruno Lemaitre and colleagues published a paper in which they for the first time showed that Toll was involved in the production of the antifungal peptide drosomycin. Subsequently, this led to the discovery of Toll-like receptors (TLRs) in vertebrates and their importance in immune responses, particularly in mammals. Research on the role of TLRs in immune responses is very intense and the importance of these receptors is probably greater in mammals than in most invertebrates. The finding that Toll was involved in induction of an antimicrobial peptide opened the possibility of performing detailed genetic studies of the signaling pathways involved in the production of antimicrobial peptides. The completion of the *Drosophila* genome in 2000 made it possible to carry out powerful molecular genetic analysis of the immune system of this insect. For the past 10 years several genomes of insects and other invertebrates have been sequenced, making comparisons between invertebrates possible, not only between *Drosophila* and mammals. Several chapters in this book deal with immune responses in different groups of insects. It is evident that the immune responses are very similar between insects, but that there are also differences (Chapters 8-12). Interesting research is now being performed on mosquitoes because they are vectors for many human diseases. One important aspect, for example, is how malaria avoids any immune responses while entering into and being inside the vector mosquitoes (Chapter 12). Interesting studies are now carried out on developmental biology and innate immunity in *Hydra* (Chapter 1), and there is now interest in studying immune reactions in other invertebrate groups,

especially in the model organism *Caenorhabditis elegans* (Chapter 6), but also in leeches and earthworms (Chapters 4 and 5).

For the past 10 years research has increasingly been focused on crustaceans (shrimp, crabs and crayfish) (see Chapter 13) and mollusks (scallop, oysters and mussels) (see Chapters 2-3), mainly because of their importance as farmed species for consumption. In these aquatic animals antibacterial peptides have a greater variation and each peptide appears to be produced in several isoforms induced by different bacterial species, indicating that there seems to be some sort of specific response to different bacterial species. The ways in which hemocytes are synthesized have also been studied in detail in crayfish, and the finding of a family of astakines similar to prokineticins, which are involved in hematopoiesis, show that these animals may be well suited for studies on hemocytes and their synthesis (Chapter 13).

Recent research has shown that there are great differences in immune responses between different invertebrate groups, but of course also that there are many similarities. Toll receptors are present in some invertebrate groups but so far the importance of their function has been mainly studied in insects, while sea urchins, which have more than 200 Toll receptors still await further studies in toll receptor function (Chapter 14). The significance of another group of molecules, the so-called Dscams, which seem to be restricted to insects and crustaceans and are also present in vertebrates, is difficult to define. Their exact role in immunity is unknown at this moment.

To further emphasize the diversity between different invertebrate animal groups, the clotting reaction can serve as a good example and is described in two chapters (Chapters 7 and 13). In crustaceans it comprises a transglutaminase and a clotting protein, whereas in horse-shoe crabs clotting it is induced by microbial polysaccharides (LPS and beta-1,3-glucans) so that a proteolytic cascade is activated and terminates with coagulogen being cleaved by the proclotting enzyme to form the clot (Chapter 7). There is no similarity between the clotting proteins of crustaceans and horse-shoe crabs, the process is totally different. The only similarity is that transglutaminase is involved in both groups of animals.

The melanization reaction is an important innate immune response and which is present in most invertebrates. It was first noted by Söderhäll and Unestam in 1977 that this process, i.e., activation of prophenoloxidase, was induced in the presence of beta-1,3-glucans. Subsequently, this was shown in insects and several other invertebrates. Recently Lee and colleagues managed to show that the prophenoloxidase activating system and the induction of the Toll pathway in an insect share the same proteolytic cascade (Chapter 9). Interestingly, melanization is a highly conserved immune response which is present in nearly all vertebrates where it provides protection against UV-light and other stressors. Melanization in vertebrates is catalysed by tyrosinase, which has no homology with prophenoloxidase except for the copper binding sites. However, the reactions tyrosinase and phenoloxidase catalyze are exactly the same, converting phenols to quinones and subsequently melanin. Melanization is responsible for skin color in humans, and in other animals, and it is also an important component in the brain and eyes.

Another innate immune system which is present in both vertebrates and invertebrates is the complement system, and to date a complement-like system seems to be operable in echinoderms (Chapter 14), horseshoe crabs (Chapter 7) and tunicates (Chapter 15). In insects the so-called thiolester-containing proteins are proposed to be complement-like

and have been shown to function as opsonic proteins in mosquitoes (Chapter 12). It remains to be shown whether true complement proteins are present in other protostome groups and not only in horse-shoe crabs.

A growing interest in invertebrate immunity is the study of the link between physiology and immunity. Surely diet, ageing, reproduction, reproductive behavior, time of day, and use of pathogens are all likely to have an effect on immunity and immune studies. The time of day for immune and challenge studies are important, and both in *Drosophila* and crayfish it has been shown that the immune system varies in efficiency during a day. This means that scientists should be aware of these facts when planning or making experiments.

In this book I have gathered scientists who are working with different invertebrates, and it can be seen that the insects are the still attracting most research and researchers. However, an increasing interest is emerging to study new invertebrate groups, especially those where the genome is known, as seen in Chapters 1, 5, 14 and 15. Even though *Drosophila* has been and still is an excellent model for immune studies, it is now clear that there are great differences between immune responses in *Drosophila* and that of several other invertebrates, which indeed calls for more research on other invertebrates.

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ABOUT THE EDITOR...



KENNETH SÖDERHÄLL studied at Uppsala University and obtained his MSc degree in 1972 and PhD in 1978. He was promoted to associate professor (Docent) in 1980. After receiving his PhD he worked in many different laboratories as a post-doc or as a guest scientist at the University of Montpellier/St.Christol, the Medical Cell Biology department at Tromsø University, the Millport Marine Biological Station in Scotland, University College of Wales in Swansea, the Marine Biological Laboratory at Woods Hole and John Hopkins University. In 1986 he held a Royal Society Fellowship at Swansea. In collaboration with Valerie J. Smith, he developed a new method to isolate and separate blood cells from invertebrates, which was based on using an anticoagulant with a low pH and EDTA, a method now used to isolate most invertebrate blood cells. When he returned to Uppsala he took a position as a researcher at the Swedish Science Research Council in 1987, obtaining a chair and appointment to professor and head of department at Uppsala University in 1989. He continued with research mainly on the proPO-system in arthropods. His group was first to clone proPO from an invertebrate and he has worked with this so-called melanization reaction for several years. His research team has also deciphered the clotting reaction in crustaceans and shown that it consists of a clotting protein present in plasma and a clotting enzyme, a transglutaminase present in the blood cells. This clotting system is distinct from that of a horseshoe crabs' in which a

proteinase cascade demonstrates the great diversity in immune processes in invertebrates. He has published 230 original papers and 65 reviews in journals and books, and he has edited 4 books. He is presently on the editorial board on *Fish and Shellfish Immunology*, and is the editor of *Journal of Experimental Biology*. Dr. Söderhäll previously served on *Journal Experimental Zoology*, *Animal Biology*, *Journal of Invertebrate Pathology* and *Developmental and Comparative Immunology*. Since 2000 he has coedited the journal *Developmental and Comparative Immunology* and beginning in 2009 he is editor-in-chief for this journal. He was president for the International Society of Developmental and Comparative Immunology from 2006-2009. Currently he heads the Department of Comparative Physiology at Uppsala University (<http://www.fu.uu.se/jamfys/is.html>).

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CHAPTER 1

CNIDARIAN IMMUNITY: A Tale of Two Barriers

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Abstract: The phylum Cnidaria is one of the earliest branches in the animal tree of life providing crucial insights into the early evolution of immunity. The diversity in cnidarian life histories and habitats raises several important issues relating to immunity. First, in the absence of specific immune cells, cnidarians must have effective mechanisms to defend against microbial pathogens. Second, to maintain tissue integrity, colonial forms have to rely on their capacity of self/nonself discrimination to rapidly detect approaching allogeneic cells as foreign and to eliminate them. And third, since cnidarians are colonized by complex bacterial communities and in many cases are home to algal symbionts, successful growth means for cnidarians to be able to distinguish between beneficial symbionts and pathogenic intruders. The aim of this chapter is to review the experimental evidence for innate immune reactions in Cnidaria. We show that in these diploblastic animals consisting of only two cell layers; the epithelial cells are able to mediate all innate immune responses. The endodermal epithelium appears as a chemical barrier employing antimicrobial peptides while the ectodermal epithelium is a physicochemical barrier supported by a glycocalyx. Microbial recognition is mediated by pattern recognition receptors such as Toll- and Nod-like receptors. Together, the data support the hypothesis that the establishment of epithelial barriers represents an important step in evolution of host defense in eumetazoan animals more than 600 million years ago.

INTRODUCTION: CNIDARIA PROVIDE INFORMATION ABOUT THE EVOLUTION OF IMMUNITY

Cnidarians are among the earliest known phyletic lineages and are sister taxon to the Bilateria (Fig. 1).¹⁻⁴ Cnidarians possess most of the gene families found in bilaterians^{1,5-7} and

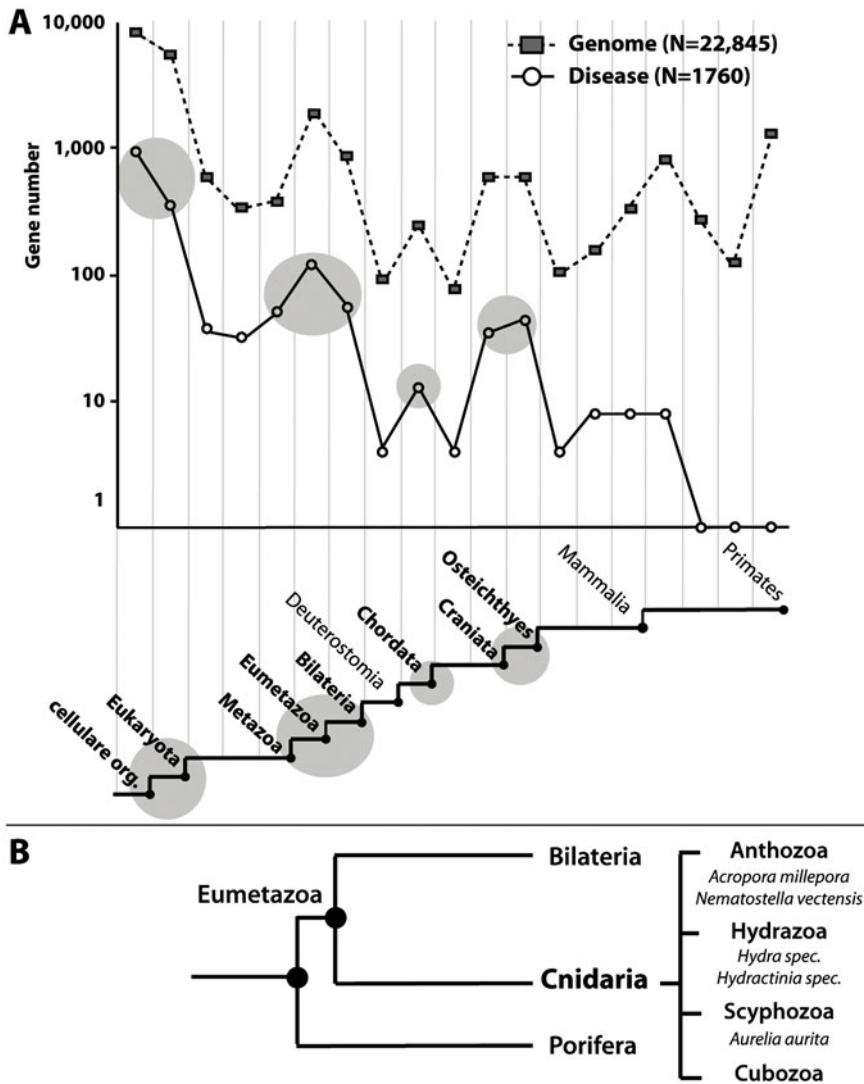


Figure 1. Phylostratigraphy of all human genes and disease-causing genes. A) The total number of human genes found in different phyla is plotted (dashed line—boxed). Distribution respectively first appearance of evaluated human disease genes in animal phylogeny (N = 1,760 black line—open circle) (Panel A modified with permission from Domazet-Lošo et al.¹⁸ ©2008 Oxford University Press). B) Cnidarians are among the earliest known phyletic lineages and are sister taxon to the Bilateria. The four classes of Cnidaria (Hydrozoa, Scyphozoa, Cubozoa, Anthozoa) contain popular model organisms for comparative immunology such as the freshwater polyp *Hydra* or the colonial Hydrozoan *Hydractinia*.

have retained many ancestral genes that have been lost in *D. melanogaster* and *C. elegans*.^{7,8} Since their genome organization and content is remarkably similar to that of morphological complex bilaterians, they provide insight into the content of the “genetic tool kit” present in the Cnidarian–bilaterian ancestor. The four classes of Cnidaria (Hydrozoa, Scyphozoa,

Cubozoa, Anthozoa) contain popular model organisms for comparative immunology such as the freshwater polyp *Hydra* or the colonial Hydrozoan *Hydractinia* (Fig. 1B).

For analytical purposes, an important technical breakthrough in Cnidarian immunology was the development of a transgenic procedure allowing efficient generation of transgenic *Hydra*⁹ and *Nematostella*¹⁰ lines by embryo microinjection. The application of morpholino techniques in *Nematostella* embryonic tissue to knock down gene expression offers a way to analyse the function of novel immuno-candidate genes during development.^{11,12} Furthermore, computational approaches have become increasingly powerful and sophisticated with each basal metazoan genome that has been completely sequenced^{1-4,13} and with each advance in the technologies used for genome analysis.¹⁴

The ultimate goal of using bioinformatics in the comparative immunology field is to be able to identify the genomic basis for key immunological characteristics appearing in our metazoan ancestor. This goal has yet to be reached, but there has been considerable progress in this area. For instance, the origin of major regulators of innate and adaptive immune response, the interferon regulatory factors family and the Rel/NFκB family, was traced back to the appearance of multicellular organisms.¹⁵⁻¹⁷ Moreover, rich sequence data from numerous species of basal metazoans allowed to determine the evolutionary origin of mutations leading to Mendelian genetic diseases. In a phylostratigraphic analysis¹⁸ of about 1700 human genes that have been linked to a heritable genetic disease, their evolutionary origin could be traced back to the origin of the first cells. This study also showed that other large groups of disease genes emerged more than one billion years ago around the first appearance of multicellular animals as well as at the time of origin of bony fishes about 400 million years ago. Surprisingly, almost no disease associated gene emerged after the origin of mammals (Fig. 1A). The study suggests that over 90% of the disease genes have emerged before the bilaterian radiation. The results are important because they show that many diseases have ancient origin and that functional knowledge gained about such disease genes from remote model organisms such as Cnidarians is relevant for understanding the genes in humans. The identification of genes responsible for human diseases affecting biological barriers (e.g., skin or intestinal mucosa) often does not in itself provide a clue to etiopathogenesis or therapeutic targets, as the interaction of a suite of genes in a complex system such as the human is difficult to understand. Likewise the involved pathways that ultimately lead to the development of the disease phenotype are unclear. Searching for the evolutionary origin of the disease-causing genes and characterizing the variation in such genes under known evolutionary pressures may provide insights into the development of diseases in humans and identify new targets for therapy or prevention. *“To understand the context of the biological processes in which a gene is involved, it may be advisable to use model organisms that represent the evolutionary level at which these genes emerged”* (cited from ref. 18, pp 2706).

CNIDARIAN IMMUNE RESPONSES ARE PERFORMED BY TWO EPITHELIAL BARRIERS

Cnidarians are diploblastic consisting of an ectodermal and an endodermal epithelium. While both layers are separated by an extracellular matrix (mesoglea) a true mesoderm is missing. In the freshwater polyp *Hydra* (Fig. 2A) epithelial cells in both layers (see Fig. 2B), are multifunctional having both secretory and phagocytic activity.^{19,20} A combined biochemical and transcriptome analysis approach revealed that in *Hydra* most innate

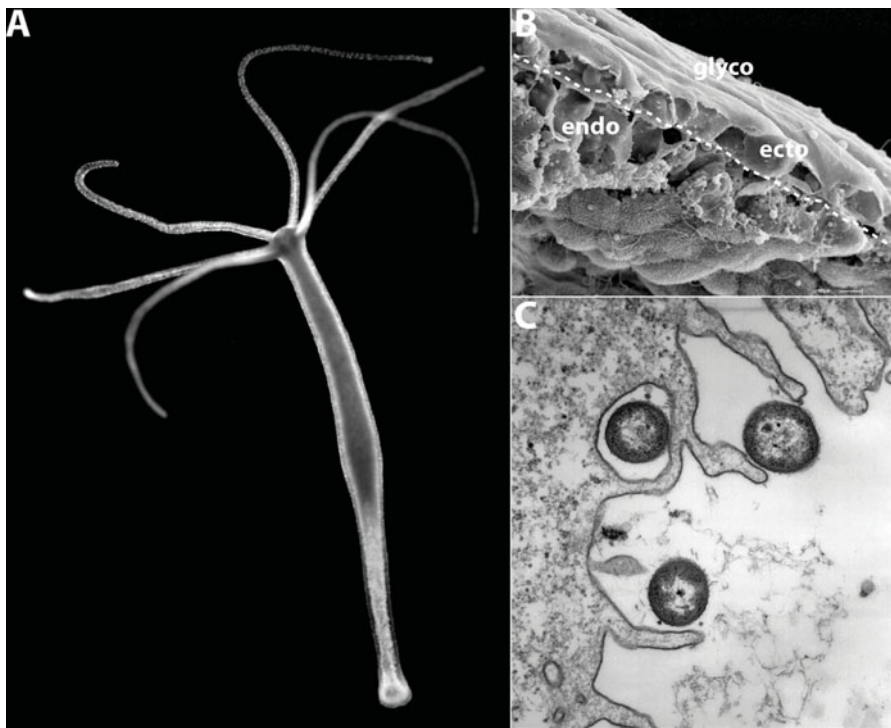


Figure 2. Cnidarians are diploblastic animals. A) Live image of *Hydra oligactis* (photo by S. Fraune). B) Raster electron micrograph showing the ectodermal (ecto) and endodermal epithelium (endo); which are separated by an extracellular matrix (mesoglea—dashed line); a true mesoderm is missing. The apical part of hydra ectodermal epithelial cells is covered by a glycocalyx layer (glyco) (photo by F. Anton-Erxleben). C) Transmission electron micrographic of endodermal epithelial cells phagocyte bacteria from the gastric lumen. (Panel C modified from Bosch et al,²⁰ ©2010, with permission from Elsevier)

immune responses are mediated by epithelial cells.^{20,21} Endodermal epithelial cells in *Hydra* not only contribute to digestion and uptake of food but also are capable to phagocytose and destroy bacteria present in the gastric cavity (Fig. 2C). Within the endodermal layer gland cells in addition to epithelial cells contribute to innate immune reactions by producing potent antimicrobial serine protease inhibitors.²² There are no motile immune effector cells or phagocytes in sensu strictu (i.e., specialized cells that engulf and ingest other cells or particles) known in *Hydra*. In the sea fan coral *Gorgonia ventalina* granular amoebocytes were reported²³ to increase in number after contact with a fungal pathogen (*Aspergillus sydowii*). Interestingly, at the site of infection a high prophenoloxidase (melanisation) activity was observed which seems to have its origin in the surrounding amoebocytes.²³ Thus, this protective strategy seems to protect a whole colony rather than the individual polyp. Some cnidarians have a remarkable capability of regeneration. In *Hydra*, for example, gross damage to the tissue is quickly repaired due to the presence of continuously proliferating stem cells.²⁴ Since cells infected or damaged by pathogens such as bacteria or fungi are quickly removed by apoptosis¹⁹ and replaced by noninfected cells, this enormous regeneration capacity may be considered an additional arm of the innate immune defense.

CNIDARIANS DO NOT LIVE ALONE BUT ARE INTIMATELY ASSOCIATED WITH SYMBIANTS

The principal function of the immune system is to mediate interactions of the host with the associated or colonizing microbiota.²⁵ Cnidarian tissue is loaded with microbes. Phylogenetic analyses as well as molecular techniques including FISH²⁶ and metagenomic approaches have demonstrated that in Cnidarians such as corals the microbial communities are highly complex.^{27,28} The communities are distinct from those in the water column, are coral species-specific, are spatially and temporally stable and most likely have multiple roles in the physiological function of the coral host.^{26,29-34} Disturbances in the balance between corals and colonizing microbiota appear to facilitate emergence of coral disease.³⁵ When investigating the microbiotic world in the freshwater polyp *Hydra*, different species were found to be associated with species-specific microbiota (Fig. 3A).³⁶ Since microbial phylotypes were identical in polyps taken directly from the wild and polyps cultured under constant laboratory conditions, the epithelia appear to select their species-specific microbes. Interestingly, disturbing the cellular composition of the epithelium results in rapid changes of the associated microbiota.³⁷ The molecular basis of this “interkingdom” host-microbe communication is currently under investigation. While the site of residence of most of the polyp-associated microbes is not yet known, in *Hydra oligactis* endodermal epithelial cells were discovered to contain intracellular symbionts belonging to the α -Proteobacteria clade (Fig. 3B).

In addition to being intimately associated with and colonized by microbes, some Cnidarian species contain symbiotic algae. In fact, Cnidaria are the phylogenetically oldest Eumetazoa phylum known to form symbiotic relationships with unicellular algae. In Anthozoans most intracellular symbionts belong to the Zooxanthellae. They are coevolved and important for long term survival of the solitary polyp or the coral colony. Various forms of stress including increase of temperature³⁸ or infection by pathogens cause cnidarian polyps to dispel their symbionts.^{39,40}

To understand and ultimately prevent this “coral bleaching”, the cellular and molecular interactions underlying this interaction are currently investigated in several coral species with particular emphasis on the establishment, maintenance and breakdown of these cooperative partnerships.⁴¹⁻⁴⁴ Most coral species must acquire symbionts anew with each generation and therefore must engage complex recognition and specificity processes, that results in the establishment of a stable symbiosis.

Another model organism for studying the set-up of long-term symbiotic interactions is the freshwater polyp *Hydra*. *Hydra viridis* forms a stable symbiosis with intracellular green algae of the *Chlorella* group.⁴⁵ The symbionts are located in endodermal epithelial cells (Fig. 3C). Each alga is enclosed by an individual vacuolar membrane resembling a plastid of eukaryotic origin at an evolutionary early stage of symbiogenesis.⁴⁶ Proliferation of symbiont and host is tightly correlated. The photosynthetic symbionts provide nutrients to the polyps enabling *Hydra* to survive extended periods of starvation.^{45,47} Symbiotic *Chlorella* is unable to grow outside the host indicating a loss of autonomy during establishment of the intimate symbiotic interactions with *Hydra*. During sexual reproduction of the host, *Chlorella* algae are translocated into the oocyte giving rise to a new symbiont population in the hatching embryo.⁴⁸ How the cnidarian host not only tolerates but also supports the continuous presence of eukaryotic symbionts within its cells is one most of the fascinating questions in biology.

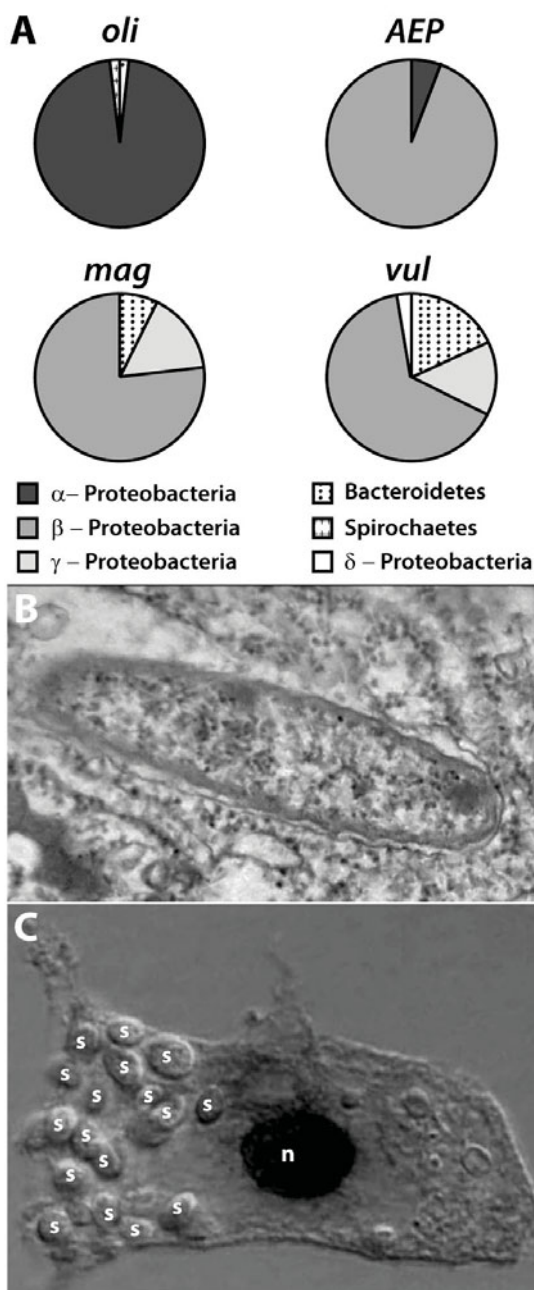


Figure 3. Cnidarians are intimately associated with symbionts. A) Bacterial communities identified from different hydra species. B) Transmission electron micrograph of bacterial endosymbiont in the cytoplasm of an ectodermal epithelial cell of *Hydra oligactis* (Panel B modified from S. Fraune et al.,³⁶ ©2007, with permission from the National Academy of Sciences, USA). C) Phase contrast micrograph of a single mazerated endodermal epithelial cell containing symbiotic algae (s) in the basal part below the nucleus (n) (Panel C modified from Habetha et al⁴⁸).

CNIDARIANS RECOGNIZE MICROBIAL ASSOCIATED MOLECULAR PATTERNS THROUGH GERM LINE ENCODED PATTERN RECOGNITION RECEPTORS (PRRs)

Among the most conserved cell surface receptors for microbial associated molecular patterns (MAMPs) are the Toll-like receptors (TLRs). While recognition of MAMPs by a prototypical TLR is mediated by leucine rich regions (LRRs) in the extracellular part, signal transduction requires the Toll-interleukin-like receptor (TIR) domain in the cytosolic region of the receptor. In vertebrates, different TLRs recognize different MAMPs.⁴⁹ Screening of the *Hydra* transcriptome and genome lead to the unexpected discovery that a bona fide TLR gene is not present.¹⁵ However, data bank searches revealed that in *Hydra* there are two transmembrane proteins with LRRs similar to LRRs present in vertebrate TLRs. These proteins, HyLRR-1 and HyLRR-2, lack any intracellular domain and, therefore, also do not contain the typical TIR domain (Fig. 4A). The screening approach also uncovered two genes, HyTRR-1 and HyTRR-2, encoding for transmembrane proteins with a canonical TIR domain but lacking any conspicuous extracellular domains. Functional assays involving HEK cells co-expressing HyTRR-1 and HyLRR-2 receptors indicated that both transmembrane proteins function together to recognize and transduce the bacterial flagella component flagellin.²⁰ Thus, in *Hydra* the TLR function is present but allocated on two different genes. Since the anthozoan *Nematostella vectensis* contains a prototypical TLR,¹⁵ MAMP recognition by TLRs seems to be a very ancestral feature. This is supported by the finding that in addition to the TLR transmembrane receptors also TLR-specific signal transduction components such as MyD88 and NF κ B are present (Fig. 4A). The findings imply that TLR-mediated innate immune responses are ancient, originating in the common ancestor of bilaterian animals. Besides flagellin which is sensed by HyTRR-1–HyLRR-2 receptor complex also LPS was found to induce innate immune reactions.²⁰ How the LPS signal is transmitted into the cell and if the TLR like pathway is involved, is currently under investigation.

In all organisms there is an elaborate intracellular detection array for internal invasion. Intracellular MAMP recognition is mediated by Nod-like receptors (NLRs). As summarized elsewhere,⁵⁰ NLR genes encode for cytosolic proteins that comprise a trimodular domain structure, characterized by a central NACHT, a N-terminal DEATH-fold-like effector binding domain, e.g., a Pyrin domain (PYD), DEATH or caspase recruitment domain (CARD) and a C-terminal leucine rich repeat region (LRR). Upon ligand sensing via the C-terminal LRRs, which can be regarded as the intracellular counterpart of the ectodomain of the TLRs, the molecules have propensity to form self-oligomers, thereby recruiting proximity activated binding partners. This leads to either activation of proinflammatory signalling pathways such as NF- κ B or direct activation of proinflammatory caspases. How do Cnidarians detect microbial invaders inside their cells? As indicated in Figure 4B, a typical *Hydra* HyNLR consists of a DEATH and a NACHT domain. Interaction with other DEATH domain containing proteins such as HyDODE may result in activation of a caspase cascade leading to programmed cell death (Fig. 4B). Homologs of the NLRs (e.g., R genes) have been discovered throughout the plant and animal kingdoms.⁵⁰ Although the Nod-receptor family has many unresolved features, the high evolutionary conservation of the NLRs underlines their significance in host defense. In contrast to vertebrates, Cnidaria such as *Acropora*, *Nematostella* and *Hydra* have numerous NLR-related receptors of different molecular configuration.⁵¹ Since several NLRs have the ability to activate the inflammasome complex and cell death,⁵² in Cnidarians with their enormous regeneration

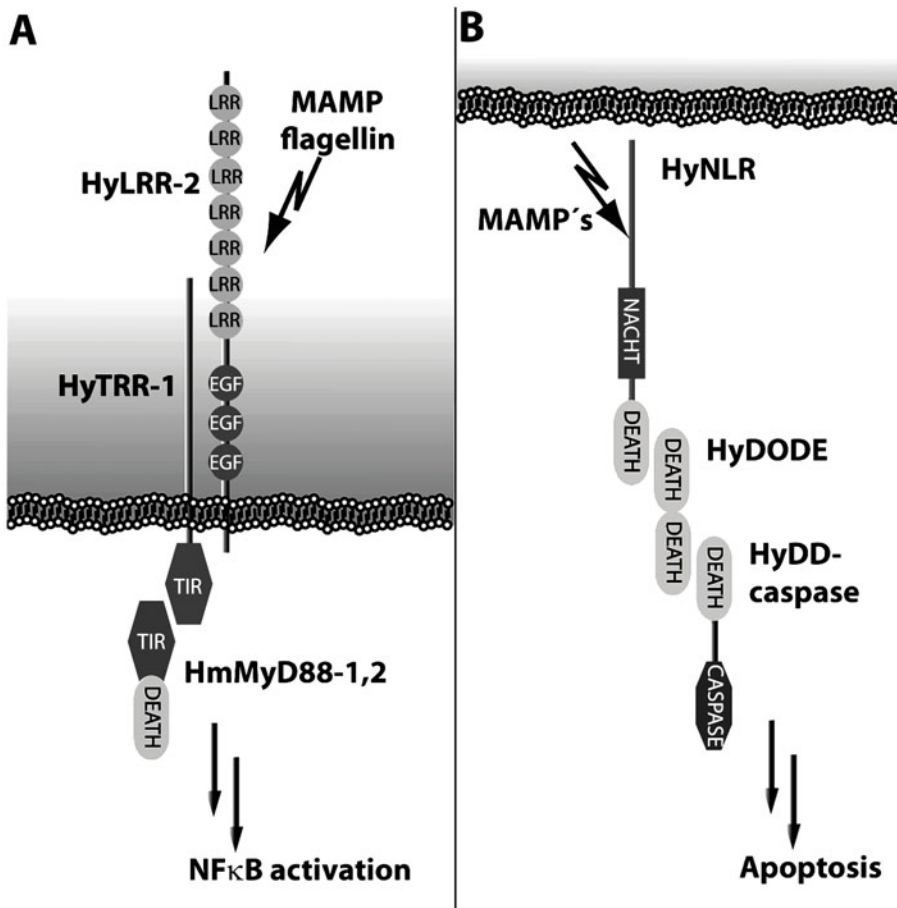


Figure 4. Microbial associated molecular pattern recognition in *Hydra*. Scheme representing the molecules responsible for A) extracellular and B) cytosolic recognition and transmission of MAMPs in *Hydra* A). The MAMP flagellin is detected by HyLRR-2 and via HyTRR-1 transmitted into the cell. The signal is transduced further by conserved TLR-signalling pathway components. B) Cytosolic MAMP's are sensed by HyNLR's and by homotypic interaction of death domain containing proteins programmed cell death is initiated (modified from Lange et al⁵¹).

capacity, apoptosis appears to be an important part of innate immunity. Unravelling the signaling pathways and exact molecular functions of these ancient NLRs will shed light on the primary cellular defense programs directed against invading microbiota.

Interestingly not only MAMPs are able to induce a defence reaction in Cnidarians. A much larger group of inducers are “danger associated molecular patterns” (DAMPs). These are substances normally buried inside the cell and therefore not detectable by neighbouring cells. In case of cell injury and/or cell death, these substances are released and suggested danger to other intact cells within the tissue. Two examples for DAMPs which are easy to apply in experimental procedures are extracellular nucleic acid and monosodium urate. In *Hydra*, increased concentrations of extracellular nucleic acid as

well as monosodium urate stimulate an innate immune response.²⁰ The receptors for these danger signals in *Hydra* or other Cnidarians are currently not known.

CNIDARIANS PRODUCE ANTIMICROBIAL PEPTIDES WHICH ARE EFFECTIVE EVEN AGAINST HUMAN PATHOGENS

Silencing of HyTRR-1 and HyLRR-2 leads to drastic reduction in antimicrobial activity²⁰ and indicates that in Cnidarians—as in other organisms—activation of MAMP receptors induces the production of a plethora of different antimicrobial peptides (AMPs). A detailed biochemical characterization of the antimicrobial activity in *Hydra* revealed the presence of both conserved AMPs such as Hydramacin-1 (Fig. 5A) and novel or taxon-specific AMPs such as Arminin 1a and Periculin-1 which have no counterpart in the transcriptomes of any other organisms. We proposed elsewhere⁵³ that these taxon-specific genes, in combination with rewiring of the genetic networks of conserved regulatory genes, accomplish fine adaptations of Cnidarians to constantly changing ecological conditions. An intriguing example of such a taxonomically restricted antimicrobial peptide is Periculin-1, termed due to its rapid response to a wide variety of bacterial and tissue “danger” signals.²⁰ Analysis of the deduced amino acid sequence of Periculin-1 and the charge distribution within the molecule revealed an anionic N-terminal region and an 8 cysteine residues containing cationic C-terminal region.²⁰ No identifiable orthologues were found in any sequence database outside the genus *Hydra*. Periculin-1 has a bactericidal activity and is expressed in the endodermal epithelium as well as in a subpopulation of ectodermal interstitial cells. Screening other taxa indicates that each animal species contains a significant number of such “orphan” genes encoding potent antimicrobial peptides. The novel antimicrobial peptide Aurelin is another examples of such orphan genes (Fig. 5A) found in *Aurelia aurita*, one of the most common and widely recognized type of jellyfish found near the coasts in the Atlantic, Arctic and Pacific Oceans.⁵⁴ Aurelin, an antimicrobial peptide with a molecular mass of 4.3 kDa was shown to exhibit activity against Gram-positive and Gram-negative bacteria. The primary structure of its molecular precursor consists of a canonical signal peptide, anionic propiece and a mature cationic part.⁵⁴

Functional characterization of Hydramacin-1 indicated a novel mode of antimicrobial action. As shown in Figure 5B, exposure of bacteria to low amounts of Hydramacin-1 caused bacteria to aggregate. Soon thereafter bacteria display typical features of programmed cell death with cytosolic proteolysis but intact cell membranes (Fig. 5B). Determination of the three dimensional structure²¹ revealed that the molecule possesses two short α -helices at the N-terminus which are separated by a long flexible loop (Fig. 5C). The C-terminal region contains two β -strands in an antiparallel arrangement separated by a long flexible loop. This predicted structure (Fig. 5C) indicates that Hydramacin-1 functions as “glue” causing clumping of bacterial cells followed by programmed cell death. Moreover, when used in liquid growth inhibition assays, recombinant Hydramacin-1 was capable of killing a large number of human Gram-negative pathogens, including the extended spectrum beta-lactamase (ESBL) strains of *E.coli*, *Klebsiella oxytoca* and *Klebsiella pneumoniae*, which are resistant to penicillin derivatives. Based on the observation of Hydramacin-induced clumping of bacteria followed by cell death (Fig. 5B) it is tempting to speculate that Hydramacin-1 acts as efficient AMP by interfering with the quorum sensing system of *Hydra*'s pathogenic bacteria.

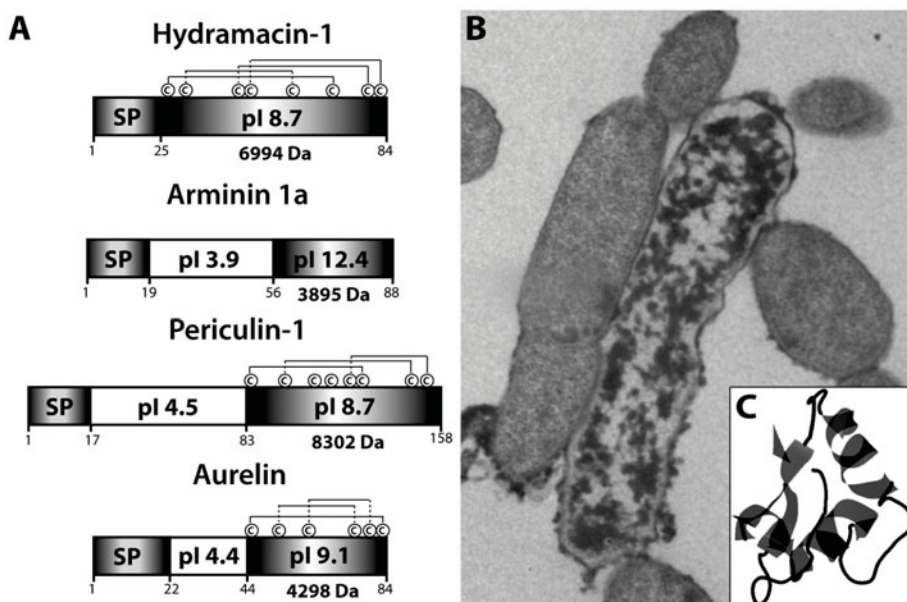


Figure 5. Cnidaria produce highly efficient antimicrobial peptides. A) Schematic representation of four antimicrobial peptides from *Hydra magnipapillata* and *Aurelia aurita*. All peptides contain a signal peptide (SP) and a cationic C-terminal active part (note the pI-value). In addition, Arminin 1a, Periculin-1 and Aurelin are made as precursor peptides, which need to be activated by the cleavage off the anionic N-terminal part. B) Transmission electron micrograph of Hydramacin-1 treated *Escherichia coli*. (photo by F. Anton-Erxleben) C) Ribbon cartoon representation of the average three dimensional structure of Hydramacin-1.

Since the last decade has seen the inexorable proliferation of a host of antibiotic-resistant bacteria including a particularly pernicious strain of bacteria known as methicillin-resistant *Staphylococcus aureus* (MRSA), we started to screen *Hydra* tissue systematically for the presence of novel antibiotics (Augustin and Bosch, unpubl.). Soon after the discovery of novel antimicrobial peptide Arminin 1a⁵⁵ we have developed the molecule “ARM-C31-N”. ARM-C31-N consists of 31 amino acids which are amidated at the C-terminus. The molecule has a calculated mass of 3896 Da and an isoelectric point of 12.1. When used in liquid growth inhibition assays, ARM-C31-N is capable of killing a large number of resistant bacteria including methicillin resistant *S. aureus* and vancomycin resistant strains of *E. faecalis* and *E. faecium*.⁵⁵ More generally speaking, antimicrobial peptides from basal metazoans which are only distantly related to humans may provide interesting leading-structures to design a novel generation of antibiotics since human pathogens share little or no evolutionary history with Cnidarian-associated microbes and, therefore, appear to be particularly vulnerable by Cnidarian-antimicrobial molecules.

Aurelin, Periculin-1 and Arminin 1a are made as precursors. For their activation a negatively charged N-terminal part is cleaved to release a highly positively charged C-terminal part. In some, but not all AMPs this cationic C-terminal region is rich in cysteines indicating that this domain requires a distinct three dimensional structure for activity. The regulatory mechanisms promoting constitutive and inducible antimicrobial peptide (AMP) expression in the epithelial barriers are not yet well understood.

Cnidarian tissue is not only packed with strong antimicrobial molecules but also with serine protease inhibitors with strong antimicrobial activity. For example, when biochemically analyzing *Hydra magnipapillata* tissue for antistaphylococcal activity we discovered a kazal-type serine protease inhibitor, Kazal-2 produced in endodermal gland cells.²² In liquid growth inhibition assays, native Kazal-2 protein has potent antistaphylococcal activity which most likely is due to the inhibition of a bacterial specific serine protease.²²

Interestingly all peptide or protease inhibitors isolated so far in *Hydra*, Hydramacin-1, Arminin 1a, Periculin-1, as well as Kazal 2, are expressed in the endodermal layer.^{20,22} We, therefore, assume that they contribute to the chemical defense properties of this layer.

HOW DO CNIDARIA DISTINGUISH SELF FROM NONSELF?

Alloimmune specificity and histocompatibility, driven by genetic polymorphism, are ancient determinants of self-/nonself-recognition.⁵⁶ *Hydractinia* is a colonial marine Cnidarian composed of a limited number of repeating structural units, polyps and stolons. Stolons are vascular-type canals which join the asexually proliferating polyps. *Hydractinia* maintain self-perpetuating stem cell lineages throughout their life history with the interstitial stem cells giving rise to the germ line and to several other cell types. After metamorphosis the planula larva forms the first polyp of a new *Hydractinia* colony. Whenever two or more planulae recruited to the same shell, the colonies may grow into contact and allorecognition interactions start. Allogeneic contacts have two major classes of outcomes: rejection and fusion.⁵⁷⁻⁵⁹ Lange et al have shown that all contacts follow a similar sequence of events.⁶⁰ When the leading edge of two asexually expanding colonies comes into contact, a large number of nematocysts are transported to the regions in contact (Fig. 6A). Once a threshold number of nematocysts have accumulated, they either disperse in fusion interactions or fire and damage the allogeneic tissue in rejection interactions.⁶⁰ These allorecognition responses play a fundamental role in maintaining the genetic and physiological integrity of the colony⁶¹⁻⁶³ because the germ line is not sequestered and because interstitial cells migrate within *Hydractinia* colonies. Thus, depending on the outcomes of allorecognition reactions, there is the risk of losing access to the germ line through “somatic cell parasitism”.^{59,64} More than 50 years ago Hauenschild postulated a model in which allorecognition and the ability to fuse between stolons of different colonies is under the control of one polymorphic locus.^{58,65,66} Five decades after Hauenschild’s pioneering experiments, an extensive inbreeding program⁶⁷ demonstrated that allorecognition segregate in a single chromosomal region but contains two closely linked loci, *alr1* and *alr2* within 1.7 cM.^{68,69} Individual *Hydractinia* that share at least one allele at both loci undergo fusion. In contrast, if no alleles are shared, a rejection process is initiated.

Analysis of the region of the *Hydractinia* genome corresponding to the *alr2* locus led Nicotra et al⁷⁰ recently to the identification of a transmembrane protein, CDS7, which has three polymorphic domains in the extracellular region (Fig. 6B). Although a functional characterization of this receptor is still missing, Nicotra et al⁷⁰ found that polymorphism in the first domain of CDS7 is an obligatory element of histocompatibility. These observations for the first time have uncovered a likely candidate for allorecognition in a basal metazoan. Since this receptor is unrelated to allorecognition receptors in tunicates⁷¹ or vertebrates (MHC), this report in addition supports the view that mechanisms used

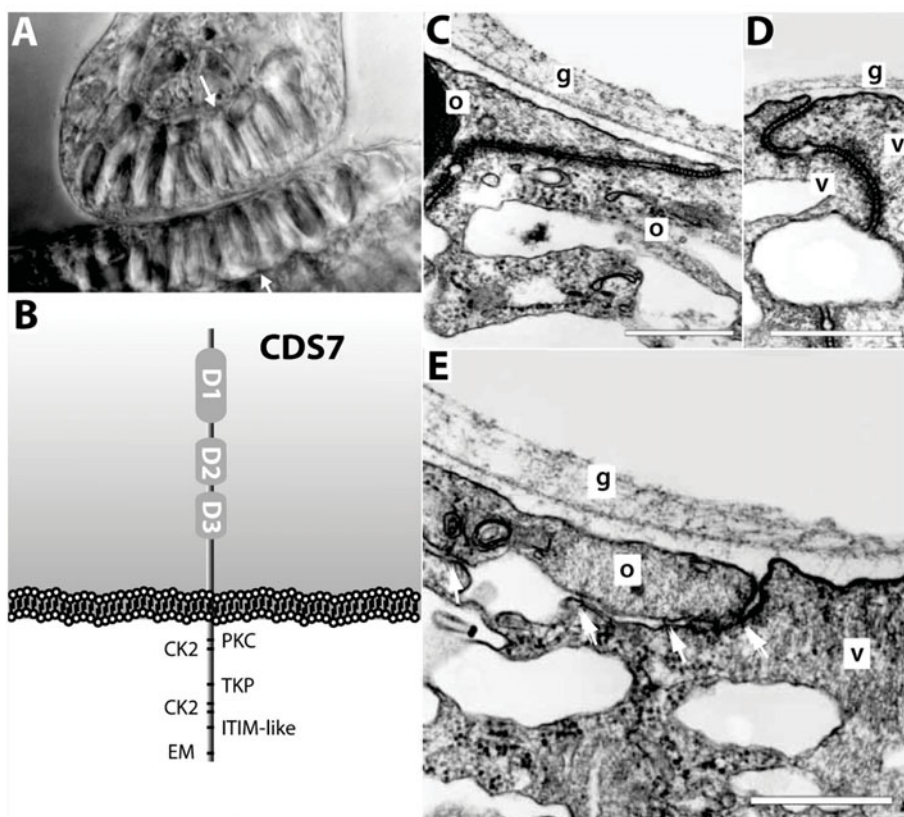


Figure 6. Cnidaria discriminate between self and nonself. A) Magnification of a stolon tip and stolon flank from rejecting *Hydractinia symbiolongicarpus* colonies, showing recruitment of nematocytes (arrowheads) to the contact point (Panel A modified from Lange et al,⁶⁰ ©2010, with permission from Elsevier). B) Schematic representation of a transmembrane receptor (CDS7) with three polymorphic domains (D1-D3) proposed from allorecognition locus *alr2* by Nicotra et al.⁷⁰ C-D) Transmission electron micrographs from ectoderm of grafting experiments between two animals of C) *Hydra oligactis* (o), D) *Hydra vulgaris* (v) and E) *Hydra vulgaris* (v) and *Hydra oligactis* (o). White arrows point to the heterotypic contact with a few irregular septae and spaces between cell membranes (compare septae in C) and D)). Note the species-specific differences in thickness of glycocalyx (g). C-E) Modified from Kuznetsov et al.⁷³

for self/nonself recognition evolved independently in different organisms.⁷² Intensive searches for identifying molecules involved in allorecognition in solitary cnidarians such as *Hydra* were unsuccessful up to now.^{73,74} There is, however, ample evidence for xenorecognition based on grafting experiments involving different Cnidarian species.⁷⁵ Close examination of such xenografts revealed strong histoincompatibility due to the inability to establish functional cell-cell contact structures such as septate junctions (Fig. 6C to E).⁷³

CONCLUSION AND PERSPECTIVE—WHERE WILL THE TALE LEAD US?

In the absence of an adaptive immune system, Cnidarians employ an elaborate innate immune system to detect and eliminate nonself using their two cell layers as efficient defense barriers. In the absence of additional protective structures the endoderm appears to function as a chemical barrier relying exclusively on the activity of endodermally produced antimicrobial substances. In contrast to the endoderm, the ectoderm in *Hydra* is covered by a glycocalyx (Fig. 2B, Fig. 6C-E). The fact that in conditions where the glycocalyx is destroyed, the epithelium is highly vulnerable to pathogenic bacteria and fungi (Augustin and Bosch, pers. observation) strongly indicates that the ectoderm may function as a physicochemical ectodermal barrier protecting against invaders from the environment. In both barriers, multifunctional epithelial cells serve for pathogen recognition and defense. Their unique properties allow each epithelial barrier to deal with specific microenvironments and control the microbiota of each milieu. An important point for understanding the maintenance of homeostasis in our ancient multicellular ancestor is that neither barrier functions alone, but within the context of an organism and thus, the function of each epithelial barrier is likely to influence each other. As researchers look more closely at innate immune responses in a wide spectrum of animal taxa, it becomes clear that in all eumetazoan animals the first responders to pathogens are often the infected host epithelial or endothelial cells, rather than the arsenal of “professional” innate immune cells (macrophages and dendritic cells).⁷⁶ Thus, strict maintenance of epithelial barriers is certainly an ancestral and essential feature of innate immunity.

In future we expect Cnidarians to contribute to at least three different areas of comparative immunology. First, the recent accumulation of genomic, phylogenetic and functional data on components of innate immunity in Cnidarians will allow to uncover the innate immune repertoire in the eumetazoan ancestor. Scientists have traditionally devoted considerably more energy to understanding how immune systems work than to how they have evolved. Devoting energy for unravelling the logic of the immune system in Cnidarians, however, seems to be indispensable to understand how it all started at the dawn of evolution of multicellular animals and—maybe even more important—to reveal the evolutionary conserved mechanisms which allows the adaptation of organismal defense to environmental conditions. Second, Cnidarians are always associated with a complex microbiota. Understanding the Cnidarian host—microbe interactions and the communication molecules involved, will not only contribute to understanding the interactions of Cnidarian epithelia with microbial communities but will also provide a window into the evolution of beneficial microbiomes. Which roles do microbial communities play in the health of Cnidarians? Does the microbial flora associated with Cnidaria contribute to the antimicrobial defense of the animal? What principles govern the assembly and maintenance of the Cnidarian microbiome? Do antimicrobial peptides not only kill bacteria but are also involved in keeping the structure of the microbial community in balance? Answers to these questions will considerably improve our understanding of the role that tissue-associated microbial communities play in health and disease. This type of basic information in simple model systems is badly needed in light of the fact that disturbance of host microbe interactions in man results in severe inflammatory diseases such as bowel diseases. Third, in human medicine, the increasing prevalence of antibiotic-resistant microbes requires the development of new antimicrobial compounds. Antimicrobial peptides of animal origin may be an effective alternative or additive of conventional antibiotics for therapeutic use. The recent characterisation of highly active antimicrobial peptides in *Hydra* and *Aurelia* show

that antimicrobial peptides from marine and freshwater cnidarians may represent a largely unexploited resource to design new antibiotics with broad-spectrum antimicrobial activity.

Much remains to be learned about the mechanisms of microbe recognition and defence in Cnidarians. It is clear that we are only beginning to understand the diversity and details of the mechanisms that are used to regulate their interaction with the microbial world, as well as the factors involved. Recent technological advances, particular in genomic analysis, however, suggest that the next few years are likely to be filled with many exciting and unanticipated discoveries that could rapidly reveal the mysteries of the two barriers.

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CHAPTER 2

GASTROPOD IMMUNOBIOLOGY

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Abstract: Over their 500 million year history, gastropods have radiated into marine, freshwater and terrestrial environments and adopted life styles ranging from herbivory to carnivory to endoparasitism to symbiont-mediated chemoautotrophy. They contend with many pathogens, including several lineages of specialized eukaryotic parasites. Their immunobiology is as yet poorly known, in part because most studies focus on a very small segment of gastropod diversity. Gastropod genome sequences are now forthcoming but synthetic overviews of the gastropod immunome are not yet available. Most immunological studies focus on interactions between gastropods and the larval stages of digenetic trematodes (digeneans) such as the medically important schistosomes. Digeneans elicit demonstrable and relevant snail defense responses and provide insights, augmented by the recently available schistosome genome sequences, for how gastropod responses are subverted. Survival of digeneans in snails depends at least in part on their ability to mimic host glycotopes, to overcome the immediate attack of reactive oxygen and nitrogen species produced by host hemocytes, and to induce long-term down-regulation of immune functions. Gastropods can mount distinct responses to different categories of pathogens, and can orchestrate effective elevated secondary responses under certain circumstances. Defense responses of at least one gastropod species, *Biomphalaria glabrata*, involve hemolymph lectins that are diversified by a variety of processes, including somatic diversification. Such observations have played a role in revising our general concept of invertebrate defense to include the possibility of more sophisticated and diversified responses beyond the production of limited repertoires of invariant pattern recognition molecules. The study of gastropod immunobiology is thus of basic interest and has several applied uses as well, including our need to conserve imperiled gastropod diversity.

INTRODUCTION: AN HOMAGE TO GASTROPOD ANTIQUITY AND DIVERSITY

Any attempt to develop an overview of gastropod immunobiology should be framed by a consideration of gastropod antiquity and diversity of life styles. Gastropods were unarguably extant by the Upper Cambrian (488-501 mya) and perhaps even earlier, in the preCambrian, more than 542 million years ago.^{1,2} They have since undergone dramatic radiations, numbering today between 40,000 and 150,000 living species,³ ranging in size from less than 1 mm to nearly a meter in length.⁴ A single New Caledonian coral reef lagoon can harbor over 2,000 gastropod species, and for some families, more than 80% of the diversity may yet to be recorded.⁵ Gastropods are the most speciose class of animals to inhabit marine environments, and have also colonized land and freshwater repeatedly. Some species have life spans measured in decades^{6,7} though many are annuals or have life histories marked by long periods of dormancy.⁸ Most gastropod species are shelled and benthic, but some have abandoned the shell and some have adopted fully pelagic life styles. Some live in extreme habitats such as hydrothermal vents or sulfide seepages. Most are herbivorous though many are carnivorous, some are kleptoparasites, and some are nearly unrecognizable, highly modified endoparasites of other marine invertebrates. Although all gastropods probably depend to some extent on symbionts for their survival, in some this dependence is extraordinary, such that the gastropod host effectively “farms” its symbionts on its gills for eventual consumption,⁹ or the gastropod/symbiont unit approaches the status of a photosynthetic¹⁰ or chemoautotrophic organism.¹¹

INFECTIOUS CHALLENGES TO GASTROPODS

Another important step in comprehending gastropod immunity is to gain an appreciation for the rogue’s gallery of pathogens with which they must contend. Like any group of organisms, gastropods have viruses^{12,13} yet the extent of challenge posed by viruses, and the nature of the gastropod antiviral response, are almost completely unknown. Gastropods in marine habitats are confronted with generalized¹⁴⁻¹⁶ and specialized bacterial pathogens¹⁷ which can cause general die-offs, especially in commercially grown species like abalones, particularly when stressed.¹⁸ For most gastropods, natural die-offs resulting from exposure to viruses or bacteria are very likely to go unnoticed, so the pathogen burden borne by gastropods may be higher than presently appreciated. The increased application of culture-free metagenomics methods to the study of gastropod microbiology¹⁹ will help clarify this issue.

Along with the usual background of viral or bacterial challenge, gastropods face other pathogens that are unequivocally gastropod specialists, the best known being the digenetic trematodes, also known as digeneans or “flukes”. There are ~18,000 nominal species of digeneans, nearly all of which are parasites of molluscs, and most of which are obligate gastropod parasites.²⁰ Digeneans typically exhibit considerable host specificity with respect to their molluscan hosts. They undergo a complex developmental program in their chosen host, one that involves intimate contact with host tissues and extensive proliferation of larval stages that culminates, often weeks after initial infection, in production and release of large numbers of cercariae (Fig. 1). Infection often persists until the death of the host and in long-lived gastropods, infections can last for decades.⁷ Digenian infection imposes a steep fitness cost as the host is partially or completely

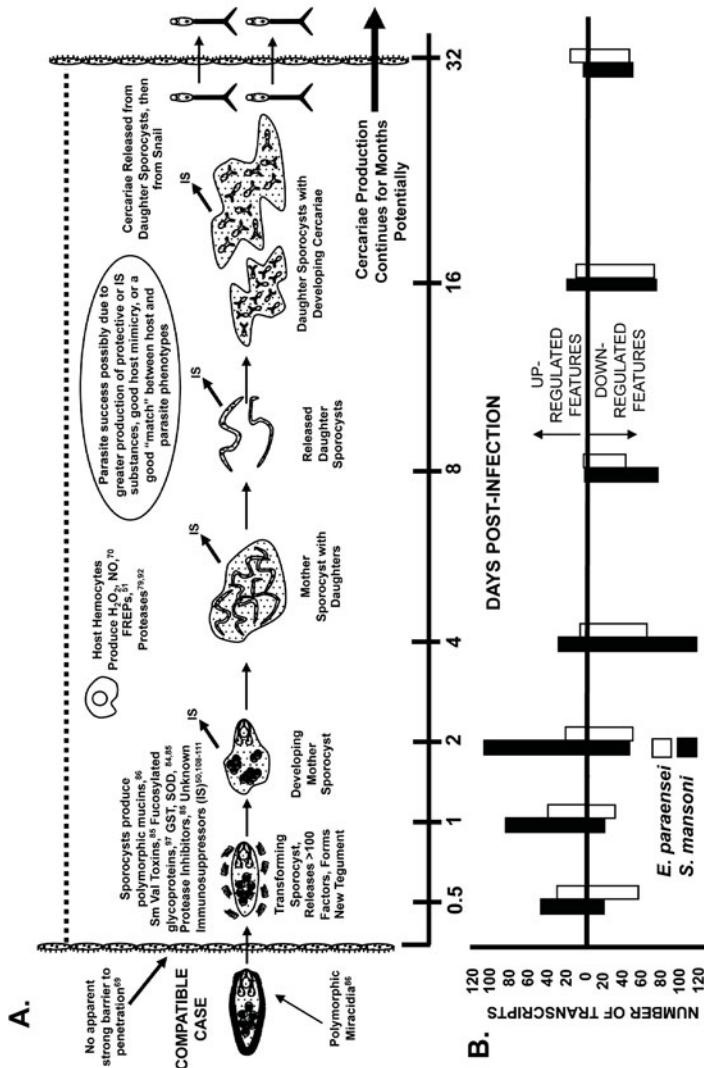


Figure 1. A) A timeline to show the sequence of events involved in the development of a digenetic trematode such as *Schistosoma mansoni* in the snail *Biomphalaria glabrata*, an example of a digenetic life cycle involving sporocyst stages and no rediae. Infection is initiated by penetration of the snail epithelium by a miracidium, followed by the sequence of events noted, culminating in the production of cercariae approximately one month later. Cercariae cross the snail's epithelium to become free-swimming; their production can persist for several months. Indicated are key references that relate to events occurring early in infection, believed to be the critical period when successful infection is established. B) Shown in the same timeframe are the results of a microarray study indicating that following exposure of *B. glabrata* to either *S. mansoni* or *Echinostoma paraensei*, there is an early preponderance of host immune-related features that are up-regulated (especially for *S. mansoni*). Thereafter, throughout the long period required to complete development, a sustained predominance of down-regulated features occurs for both species.⁵⁰ This is indicative of the need for these parasites to induce a long-term state of immunosuppression while at the same time massively proliferating in their host, all without causing its premature demise.

castrated.²¹ Digenean infections are known from a broad spectrum of gastropods ranging from basal limpet species to derived heterobranchs. Some gastropod species play host to dozens of digenean species, and simultaneous infection of individual gastropods with two or more digenean species is not uncommon.

A nonexhaustive list of additional pathogens containing at least some members with a significant degree of specialization on gastropods includes apicomplexans,²² microsporidians,^{23,24} ichthyosporeans,²⁵ nematodes,²⁶ copepods²⁷ and hematophagous mites.²⁸

It must be noted that the vast majority of gastropod species have never been studied in an immunological context: our current information is based on just a few species, biased towards those of medical or commercial significance (Fig. 2). Furthermore, we should resist the temptation to assume that all gastropod defense systems are the same. It seems probable, especially given the age of the gastropod lineage, that the unique challenges posed by different habitats, feeding styles, and levels of commitments to symbiotic associations will have selected for diverse kinds of immune systems. For example, species heavily besieged by digeneans may have adopted very different immune “portfolios” than species living in dense colonies where directly-transmitted microparasites may be more of a concern. These portfolios might feature expansions of different gene families or the emergence of entirely novel defense mechanisms to deal with their respective immune challenges. The study of

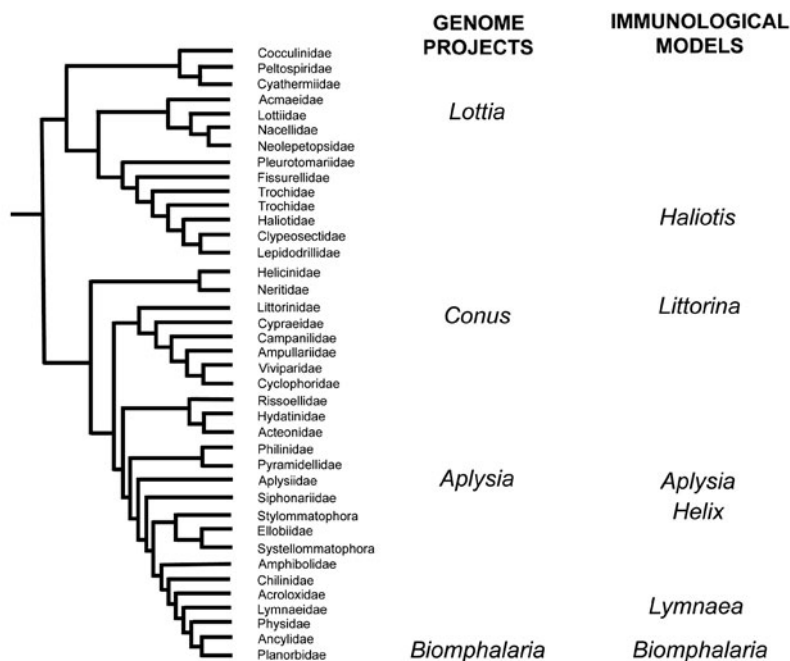


Figure 2. An overview of gastropod phylogeny, with emphasis on taxa most relevant to the study of gastropod immunobiology. This tree should not be viewed as a formal hypothesis of gastropod relationships but as an overview to assist readers in placing the approximate phylogenetic positions among gastropods prominently figuring in studies of gastropod immunology. Relationships outlined in trees presented by Aktipis et al (2008)⁴ and Klussmann-Kolb et al (2008)¹³⁷ were used to delineate the relationships among the groups singled out for discussion.

alternative models of gastropod immunity should be encouraged, and Table 1 highlights some of the recent results forthcoming from the study of other than freshwater pulmonates.

GASTROPOD GENOME PROJECTS—AWAITING THE DELUGE

The completion of genome sequencing projects has revolutionized our understanding of both ecdysozoan^{29,30} and deuterostome³¹ invertebrate immunobiology. Recent years have also seen the initiation of gastropod genome projects, two of which have now been substantially completed with a third well underway. The first gastropod, and molluscan, genome sequence to be obtained is that of the owl limpet *Lottia gigantea* (Patellogastropoda, Lottiidae). The 359.5 Mbp *L. gigantea* genome is also the smallest known molluscan genome, and has been sequenced to 8.87X coverage and both automated and custom annotation provided (<http://genome.jgi-psf.org/Lotg1/Lotg1.home.html>). The initial shotgun sequence (7X coverage) and assembly of the 1.8Gbp genome of the California sea hare *Aplysia californica* (Opisthobranchia, Aplysiidae) has also been completed (<http://www.broadinstitute.org/science/projects/mammals-models/vertebrates-invertebrates/aplysia/aplysia-genome-sequencing-project>), and sequencing of the 931 Mbp sequence of the medically important gastropod *Biomphalaria glabrata* (Pulmonata, Planorbidae) is well underway (<http://biology.unm.edu/biomphalaria-genome/index.html>). A genome project emphasizing the venoms (venomics) of the cone snail *Conus consors* (www.conco.eu) is also underway. Fortuitously, these genome projects provide representation for diverse parts of the gastropod family tree (Fig. 2). As yet there has been no overarching synthetic or comparative interpretations for any of these genomes, particularly with respect to elaboration of the immunome, though a deluge of new information will be forthcoming in the near future, one that will surely also revolutionize our understanding of gastropod immunobiology.

THE DISTINCTIVE ARCHITECTURE OF GASTROPOD IMMUNE SYSTEMS OFFERS MANY OPPORTUNITIES FOR STUDY

The soft, moist body surface of gastropods is protected by a ciliated, mucus-producing epithelium that provides an initial physical trap and barrier to colonization by pathogens, one that is at present under-studied with respect to its role in defense.³² The isolation of achacin from the body mucus of the giant African land snail *Achatina fulica* suggests mucus also provides a chemical barrier. Achacin is an L-amino oxidase that generates H₂O₂ from oxidative deamination of L-lysine and L-arginine and has antibacterial and tumoricidal properties.³³ The regular production and shedding of body mucus is also likely advantageous in cleansing the surface of pathogens like bacteria which are stimulated to grow in biofilms in mucus trails.³⁴ The composition, turnover rates, and dynamics of pathogen shedding of snail mucus are all worthy of more study.

Hemocytes—Multifunctional Guardians of the Snail Body

The open circulatory system of gastropods is populated with circulating defense cells called hemocytes (formerly often called amebocytes) with well appreciated defensive roles in phagocytosis and encapsulation reactions. These cells wander into tissue spaces as well. Gastropods also possess “fixed” defensive cells that are found around internal

Table 1. Important models to increase the phylogenetic breadth of the study of gastropod immunobiology

Function	<i>Lottia</i>	<i>Haliotis</i>	Gastropods Studied <i>Littorina</i>	<i>Aplysia</i>	<i>Helix</i>
Immune cell and system description		hemocyte description ¹⁵	hemocyte description ^{38,56} Signaling pathways ³⁹	hemocyte description ³⁹ Albumen gland source of AMPs ⁴⁰ Aplysianin in eggs ⁴¹	hemocyte description ¹³⁸ Agglutinins from albumen gland ⁷⁴
Genome Mining	G-protein coupled receptors ⁶² FBG-encoding genes ⁹⁵			Ink as source of tumorcidal L-AA ¹⁴² 2 FREPS ¹²⁷	
Immune Gene Discovery		Perforin-like ^{143,144} NF-KB homolog ⁶⁷ Pattern recog ¹⁴⁵ B-thymosin ¹⁴⁶ Cytidine deaminase ¹⁴⁷ Abhisin AMP ¹⁴⁸ TNFa homolog ¹⁴⁹ GST cloned ¹⁵⁰			

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Table 1. Continued

Function	<i>Lottia</i>	<i>Haliotis</i>	Gastropods Studied <i>Littorina</i>	<i>Aplysia</i>	<i>Helix</i>
Immune Gene Discovery		Fas ligand ¹⁵¹ C-type lectin ¹⁵² Mx protein ¹⁵³			
	Immune-Pathogen Interactions		Vibriosis ¹⁵⁴ Withering syndrome ¹⁷ Stress review ⁴⁷ Vibrio and p38 ⁶³ Pedal mucus and bacteria ³⁴	Signalling and trematodes ¹⁵⁵ Hemocytes and digeneans ¹¹¹ Hematopoiesis and digeneans ¹⁰⁶	
			Bacteria for trt of vibriosis ¹⁵⁶ Global warming effects ¹⁸ Herpesvirus ¹⁵⁷ Gene regulation study ¹⁵⁸ Tagged vibrios ¹⁴ Stress markers ¹⁵¹		

spaces and trap and/or phagocytose particulate objects.³⁵ Also present are rhogocytes (also known as pore cells) which are involved in synthesis or processing of respiratory proteins,³⁶ but have also been implicated in ingestion of small foreign particles.³⁷ The relative contribution of fixed cells to defense is not well understood.

In addition to their more familiar roles, hemocytes are also involved in wound healing, nerve repair, shell formation and repair, tissue remodeling possibly including dissolution of gonads after spawning, and metabolite and nutrient movement.³⁸⁻⁴⁰ Gastropod hemocytes also engage in diapedesis, the movement of ingested foreign materials to and across external or gut epithelia.³⁷ The extent to which this occurs and influences production of new hemocytes or potentially increases susceptibility to new pathogen challenges is unknown.

We are far from a comprehensive understanding of where hemocytes are produced, how many distinct lineages of hemocytes are produced, their life spans and how functionally diversified they are: these parameters will almost certainly be shown to vary among different gastropod lineages. A distinct hematopoietic organ, called the amebocyte producing organ or APO, has been identified in some snails like *B. glabrata* and *Lymnaea truncatula*.^{41,42} The APO of *B. glabrata*, if removed from snails of a strain resistant to *Schistosoma mansoni* infection and implanted into snails of a susceptible strain will successfully transfer a degree of resistance.^{43,44} Others have doubted whether this organ alone can provide sufficient numbers of hemocytes for the entire snail, and have postulated that hemocyte production also occurs in peripheral vascular locations.⁴⁵ In other pulmonates, both individual circulating and tissue-dwelling hemocytes can divide,⁴⁶ and circulating blast-like cells in *Littorina littorea* are also capable of division.³⁸ For abalones, a hematopoietic organ has yet to be localized.⁴⁷

The pathways involved in activating hematopoiesis are poorly known and may involve direct stimulation by pathogens or indirect stimulation via mitogenic hemocyte cytokines.⁴⁸ Excretory/secretory products of sporocysts of the digenean *Echinostoma paraensei* stimulated enlargement of the APO of *B. glabrata*,⁴⁹ and extracts of *S. mansoni* stimulated an increase in mitosis in excised APOs prompting the suggestion that the parasite provides a direct mitogenic or nutritive effect on hematopoiesis.⁴⁸ APOs treated with phorbol myristate acetate (PMA), a stimulator of protein kinase C (PKC), showed increased mitotic activity.⁴⁸ Recent studies have identified from *B. glabrata* potential mitogen-related proteins including TGF- β Type 1 receptors and epidermal growth factor (EGF)-related proteins, which are up-regulated early in the course of digenean infection and thus are promising candidates as hematopoiesis-promoting factors.⁵⁰

A monograph could be written on the classification and description of gastropod hemocytes,^{15,39} and suffice it to say here that gastropod blood typically contains hemocytes that do not spread or spread minimally when placed on artificial surfaces (often called round cells, or blast-like cells), whereas the majority of cells typically spread avidly, forming conspicuous filopodia or lamellopodia in the process, and are believed to be the mature effectors in phagocytosis and encapsulation responses. Such spreading cells have been referred to as hyalinocytes or as granulocytes, the difference in terminology relating to the numbers of "granules" within the cells, a parameter likely to vary among gastropod taxa and as a function of hemocyte ontogeny. Round/blast-like cells are frequently considered to be of an earlier stage in hemocyte ontogeny, and in *L. littorea* such cells were shown to have incorporated 5-bromo-2'-deoxyuridine indicative of recent cell division and are believed to differentiate into mature effector cells.³⁸

Related subjects awaiting further study are how long gastropod hemocytes live, and whether mature effector hemocytes are terminally differentiated, although it has been noted hemocytes participating in encapsulation and wound healing responses can still divide.⁴⁶ In comparison to other invertebrates, hemocytes of the small number of gastropods so far studied are relatively few in distinctive morphological types, generally lack large aggregates of conspicuous granules or pigments, lack flagella, and do not have an obvious propensity to lyse when removed from the host or presented with an antigenic stimulus. Gastropod hemocytes are characteristically sticky and readily form aggregates, a tendency noted to be more pronounced in marine than freshwater species.³⁸

Hemocyte Signaling Pathways

Hemocytes are responsible for synthesis and release of several defense-related factors such as reactive oxygen species (ROS),³⁹ lectins such as fibrinogen-related proteins (FREPs)⁵¹ and antimicrobial peptides (AMPs).⁵² As compared to many other invertebrates, the study of gastropod AMPs is in its infancy. Hemocyte membrane-associated lectins have been implicated in detection of intruders. For example, a galectin present on the surface of ~60% of *B. glabrata* hemocytes has been characterized, and in recombinant form binds to the tegument of *S. mansoni* sporocysts in a carbohydrate-inhibitable manner, suggesting it is a hemocyte-bound pattern recognition molecule.⁵³ It was also suggested there may be “counter receptors” on hemocytes, such as integrin-like molecules,⁵⁴ that could be bound by soluble forms of galectin, such that the galectin could also serve in cross-bridging hemocytes to a parasite surface.⁵³

Key to understanding hemocyte effector functions are the intracellular signal transduction pathways likely to be activated by exposure of hemocytes to exotic stimuli. PMA stimulation of hemocytes, acting on PKC, results in H₂O₂ generation in *B. glabrata*,⁵⁵ of superoxide anions in *L. littorina*,⁵⁶ and of NO in *Lymnaea stagnalis*.⁵⁷ PKC activation is likely to result in activation via phosphorylation of mitogen activated protein kinases (MAPKs) like ERK or p38 because inhibitors of MAPKs also prevent hemocyte spreading or H₂O₂ production.^{58,59} More natural stimuli such as laminarin also activate PKC and H₂O₂ production in *L. stagnalis* hemocytes.⁶⁰ A role for phosphatidylinositol 3-kinase in controlling phagocytic activity has been shown in *L. stagnalis* hemocytes,⁶¹ and G-protein coupled membrane receptors have also recently been reported from *L. littorea*.⁶² The notion that components of gastropod signaling pathways such as p38 can be targeted by pathogens, as they often are with pathogens of plants, has recently been documented in abalones contending with *Vibrio harveyi*.⁶³

Toll-like receptors (TLRs) and associated Toll pathway components such as MyD88 are known from cephalopods⁶⁴ and bivalves.^{65,66} Homologs of TLRs are present in the *Lottia* genome, a Rel-like NF- κ B transcription factor is known from abalones,⁶⁷ and both Dorsal-like and Rel-like transcriptional factors occur in *B. glabrata* (Zhang, personal communication) so additional Toll pathway homologs are likely to be present in gastropods, although their functional relevance is as yet unknown. It is somewhat surprising that the numerous studies focused on digenean-snail interactions have not as yet turned up TLRs or transcription factors of canonical Toll-signaling pathways. Acting upstream of Toll-signaling pathways in other invertebrates are pattern recognition molecules like PGRPs (peptidoglycan recognition proteins) and GGBP (gram negative binding protein, or B-1-3 glucan recognition/binding protein or LGBP). Both short and long form PGRP-encoding genes are present in *B. glabrata*⁶⁸ and the latter generates different

transcripts via alternative splicing, some of which may have amidase activity. At least three different GNBPs are also known from *B. glabrata*, and two of these group with other GNBPs that may preserve glucanase activity.⁶⁸ Short form PGRP and GGBP were down-regulated at 6 hours after exposure to 3 types of microbes and neither responded early in the course of digenean infection, whereas the former was modestly up-regulated after two weeks of digenean infection. Long-form PGRP expression was consistently low and not modified by exposure to microbes or digeneans.⁶⁸

A Plethora of Immune-Relevant Molecules Awaiting Functional Validation

One of the conceptual cornerstones of gastropod immunobiology is that hemocytes interact in concert with several different categories of molecular defense factors, many of them circulating in the hemolymph, together more effectively recognizing and disabling pathogens.^{39,56} Among the prominent categories of immune-relevant molecules either produced by, or working in concert with, hemocytes are ROS, lectins, AMPs, PGRPs, GNBPs, proteases, protease inhibitors, and complement-like proteins. Much of the current focus of gastropod immunobiology is on revealing and identifying these factors, many of which are alluded to in the following sections of this document, and have been recapped in recent reviews^{69,70} and in Table 1 and Figure 3. As just a few recent examples of intriguing immune-relevant molecules that have also proven to be responsive to microbial or digenean stimulation on a *B. glabrata* oligo-based microarray,⁷¹ are a complement C1q-like protein with a lectin domain, serpins, several signaling pathway components including an NF- κ B p105 subunit, several FREPs, a FREM which is a unique juxtaposition of fibrinogen and EGF domains, and a macrophage migration inhibition-like factor. All await further study. Also most worthy of further consideration are the many “unknown” sequences (up to 60%) responsive to immune challenge that are often recovered from gene discovery efforts. Since they do not fit established paradigms, further study of immune responsive unknowns may lead to the most novel future discoveries of gastropod defenses.

Some Additional Comments on Gastropod Internal Defense Systems

Gastropod blood does not obviously coagulate upon removal from the animal or after exposure to pathogens. However, tubular helical filaments visible with electron microscopy have been found in hemolymph,^{39,72} and plasma from *B. glabrata* infected with *E. paraensei* often forms precipitates.⁵¹ It has been suggested that matrilin in *B. glabrata* plasma might participate in formation of filaments which could serve to keep hemocytes from contacting digenean sporocysts in susceptible snails.⁷³ Further study is thus warranted to determine if gastropods have as yet unappreciated abilities to alter their hemolymph in ways that might be easily overlooked and are relevant for defense.

The albumen gland, well-known for its role in providing nutrients in the perivitelline fluids packaged around each developing embryo in an egg mass, has received increasing attention for its immunological roles. A unique hexameric “H-type” lectin is produced by the albumen gland of *Helix pomatia* and *Cepaea hortensis* and is believed to protect developing embryos from bacteria.⁷⁴ A recent proteomics study revealed that 16 of 20 identified polypeptides in *B. glabrata* egg mass fluids have a defensive role, and include protease inhibitors, compounds with possible phenoloxidase activity, Cu-Zn superoxide dismutase, C-type lectins, GGBP, aplysianin/achacin type proteins, and LBP/BPI or lipopolysaccharide binding protein/bacterial permeability increasing protein.⁷⁵ Defensive

CONSTITUTIVE DIFFERENCES BETWEEN SUSCEPTIBLE AND RESISTANT SNAILS

- Resistant Snails have more hemocytes than susceptible snails
Bg-Sm: 176
- Resistant hemocytes have different behavior or properties
Bg-Sm: 177
Bt-Sm: 178
- Resistant Snails produce more parasite-toxic reactive oxygen species (ROS)
Bg-Sm: 70,101,175-182
- Resistant Snails have novel recognition/killing capacity in their plasma
Bt-Sm: 183,184
Bg-Sm: 44
- Innate differences in the levels of potentially immune relevant factors
Bg-Sm: 185
Bg-Ec: 73,76,84,186,187
Bt-Sm: 188
- Resistant snails have greater ability to resist parasite offensive strategies, including signaling pathway targets
Bg-Sm: 6,116
Bg-Ec: 110
- Resistant and susceptible snails may differ in the ease with which they are mimicked by parasites
Bg-Sm: 97

RESPONSE CAPACITIES FOLLOWING EXPOSURE TO INFECTION

- Resistant Snails are more stimulated to make hemocytes following exposure
Bg-Sm: 189
- Resistant snails respond more prominently or differently to infection than susceptible snails
Bg-Sm: 92,100,102,103,185,189-191
Bg-Ec: 73,76,77,94,187
Bt-Sm: 188

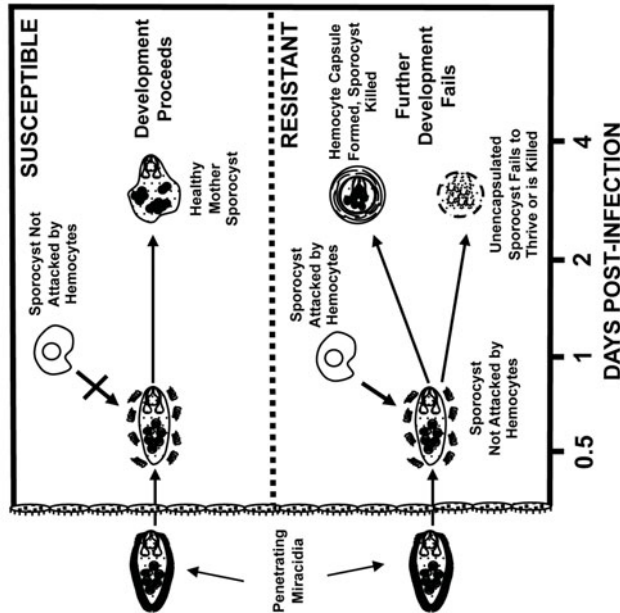


Figure 3. Explanations for the phenomenon of resistance of snails to digenean infection. At least three different experimental model systems have been developed to examine the phenomenon of resistance of snails to trematode infection: Bg-Sm: This is the most prominent and involves *Schistosoma mansoni* and its interactions with *Biomphalaria glabrata* strains either selected for resistance (such as 10-R2 or 13-16-R1 strains) or that are naturally resistant (BS-90 snails, also known as Salvador snails). Bg-Ec: This model involves *B. glabrata* and strains selected for resistance or susceptibility to *Echinostoma caproni*. Bt-Sm: A third model gaining recent prominence involves *S. mansoni* and *Biomphalaria tenagophila* isolates that are either naturally susceptible (Cabo Frio) or resistant (Taim) to infection. Some of the prominent explanations for the underlying reason d'être for resistance derived from these models and recent applicable references are outlined in the attached list. The more recent papers will lead readers to additional older papers on similar topics.

compounds produced by the albumen gland are probably distributed systemically with the plasma to protect the adult snail as well.^{76,77}

One surprising feature of gastropod immunity is the limited extent to which snails so far have been shown to rely on phenoloxidase and the eventual production of melanin to kill or wall off pathogens, at least in the most commonly studied gastropod models.^{39, 78} PPO activity is reported from snails,⁷⁹ and the egg mass fluids they produce,⁸⁰ and it is conceivable that particular groups of gastropods, once investigated, will have a greater reliance on melanization and associated reactions for defense than seen thus far. A molluscan hallmark, the ability to produce the stacked, compartmentalized structure of aragonite crystals to make shell nacre, can also be exploited in defense by encasing macroscopic pathogens such as trematode metacercariae. The molecular basis of nacre formation is being revealed in bivalves⁸¹ and homologs should be sought in gastropods.

SNAILS AND DIGENEANS AS MODELS TO STUDY SPECIFIC, INTIMATE AND LONG-TERM HOST-PARASITE SYSTEMS

The recent study of gastropod immunology has been dominated by the use of freshwater pulmonate gastropods, especially the planorbid snail *B. glabrata*, as study subjects. This is understandable because such snails serve as the intermediate hosts for digeneans of medical or veterinary significance, such as *S. mansoni* in the case of *B. glabrata*. Digeneans are conveniently available, natural pathogens of gastropods that can be used as probes to elicit relevant host immune responses then amenable to study. Snail-digenean systems are excellent models to study the “give and take” involved in intimate, long-term host-parasite associations. The use of digeneans as a means to study gastropod defenses is more fruitful now than ever because genome sequences have been recently forthcoming for *S. mansoni* and the related *S. japonicum*.^{82,83} These have already been of use in illuminating aspects of the intramolluscan biology of digeneans^{84,85} that in turn provide important clues for how the snail response might be orchestrated.^{86,87} Furthermore, a number of important tools and resources have been developed that not only allow further dissection of the outcome of digenean-snail encounters but also enable exploration of responses to other pathogens, or to environmental perturbations. These include the ongoing *B. glabrata* genome project⁸⁸ and associated development of BAC libraries⁸⁹ and EST databases,⁹⁰ the development of microarrays for both digenean⁹¹ and snail,^{71,92} the use of echinostome flukes such as *Echinostoma paraensei* or *E. caproni* to elicit host responses that can be compared and contrasted with those engendered by *S. mansoni*⁶⁹ and the development of strains of *B. glabrata* susceptible or resistant to infection with either *S. mansoni* or *E. caproni* (Fig. 3). Recent reviews can be consulted for more detail,^{21,69,70} particularly with respect to the specific identity of a number of candidate molecules identified for further study.

Use of both EST datasets and a *B. glabrata* microarray have shown that the gastropod response to different insults—wounding, injection of gram positive or negative bacteria, or exposure to infection with *S. mansoni* or *E. paraensei*—is not “one size fits all”^{50, 71} and that snails can discriminate among different kinds of immunological stimuli, a phenomenon also noted by others.^{90,93-95} Wounding alone provoked rather mild responses, the two kinds of bacteria upregulated only partially overlapping subsets of array features, and the response to the two digeneans not only were different, but also easily separable from those elicited by bacteria.⁷¹

With respect to the more specific issue of the determinants of compatibility between digeneans and snails, several recent productive lines of investigation have been developed, many scrutinizing differences between schistosome-susceptible and -resistant snails (Fig. 3). None of the explanations generated should be ruled out and all may apply to one extent or the other. With respect to the role of ROS, hemocytes from snails resistant to *S. mansoni* infection produce higher levels of H₂O₂ than do susceptible snails.⁷⁰ Hydrogen peroxide is particularly lethal to *S. mansoni* sporocysts which have also been shown to be naturally vulnerable to high oxygen tensions.⁹⁶ Furthermore, resistant snails possess a different allelic form of the enzyme Cu/Zn superoxide dismutase involved in hydrogen peroxide production than do susceptible snails,⁷⁰ though it is unclear if this difference or other gene regulatory steps are responsible for the different levels of hydrogen peroxide achieved. On the parasite side, proteomic studies of the larval transformation proteins (LTPs) of both *S. mansoni* and *E. paraensei* sporocysts^{84,85} reveal molecules involved in scavenging of ROS to be prominent, highly suggestive of their importance in protection from hemocyte attack. Also found among the LTPs are intriguing venom-related proteins⁸⁵ and glycoconjugates,⁹⁷ many of which bear fucose residues and make excellent candidates for host-mimicking molecules as host molecules are also known to bear fucose residues.^{98,99} As noted by Wu et al.,⁸⁵ a veritable “cloud” of parasite produced factors are released, at least 99 of which have been characterized.⁸⁵ Among these molecules are highly polymorphic mucins and it has been speculated that these parasite produced molecules might serve as a smokescreen to divert attack by humoral factors such as FREPs,^{86,87} at least some of which have a known binding preference for fucose residues.⁵¹ Thanks to the availability of the *S. mansoni* genome, we can gain a much better perspective on the kinds of molecules released by transforming parasites, or by developing daughter sporocysts, and can begin to formulate hypotheses for how these might confuse or suppress the host defense response.

Although protective and potentially obfuscating parasite molecules are produced, it is clear that *B. glabrata* is nonetheless able to detect the presence of either *S. mansoni* or *E. paraensei* infection within hours of penetration.^{50,92,100-103} Thus unless subverted, a growing host response could increasingly jeopardize the developing parasite. A number of lines of evidence stemming from the original work of Lie and colleagues¹⁰⁴ suggest immunosuppression, or “interference” in the parlance of the snail-digenean literature, is mediated by larval digeneans, and, importantly, that this effect can be implemented within one hour after penetration¹⁰⁵ and likely persists for the duration of the infection.⁶⁹ Hemocytes from digenean-infected snails often show diminished ability to adhere to surfaces, engage in phagocytosis or to encapsulate trematode larvae^{106,107} effects that can be reproduced by exposure of hemocytes to LTP from echinostome sporocysts or rediae.¹⁰⁸⁻¹¹¹ A recent microarray study documenting the responses of *B. glabrata* from 0.5 to 32 days following exposure to either *S. mansoni* or *E. paraensei* revealed that both parasites, after a brief early preponderance of upregulated array features, provoked a persistent excess of down-regulated features relative to up-regulated features after 2 dpe (Fig. 1).⁵⁰ Insofar as this study examined whole body extracts from exposed snails it is not unreasonable to assume that significant down-regulation occurred even earlier among host cells in the immediate vicinity of developing parasites, in accordance with earlier work documenting trematode-mediated interference in *B. glabrata*.^{109,112} Also, it should be emphasized that although interference has been most associated with echinostome development, this recent study, also in agreement with Lie’s classical studies,¹¹² suggests that schistosomes also engage in a strategy of immunosuppression though it seems to take somewhat longer to be manifested.⁵⁰ Indeed, based on results

obtained from other snail-digenean studies where evidence consistent with interference has been obtained,¹¹¹ it seems likely that a strategy of immunosuppression may be a general requirement for parasites like digeneans with a long developmental program and persistent infections.

The specific digenean-produced factors responsible for inducing interference, and their exact mechanisms of action are unknown, but more detailed information regarding the in vitro larval digenean proteome^{84,85} provide good starting points for testing candidates, though the actual in vivo proteome of larval digeneans may be somewhat different. One likely possibility is that digeneans produce factors that alter the activity of hemocyte signaling pathways, likely in more than one way.^{113,114} Exposure of adherent *B. glabrata* hemocytes to LTP of *E. paraensei* sporocysts or rediae caused both induction of calcium waves and rounding, not always both in the same cell, suggestive of at least two distinct effects of LTP.¹¹³ Following exposure to LTP or to whole *S. mansoni* sporocysts, ERK-dependent signaling was impaired in hemocytes from *B. glabrata* susceptible to infection, but not in schistosome-resistant snails. As ERK regulates important hemocyte effector functions, it was suggested that disruption of ERK signaling, possibly via carbohydrates expressed on parasite surfaces or in LTP, facilitated survival of *S. mansoni* in susceptible snails.^{115,116} It was also noted that components in plasma may protect hemocytes from the impact of parasite-mediated suppression of hemocyte signaling pathways,¹¹⁵ providing a potential explanation for why some plasma factors, for instance FREP4, are persistently upregulated in snails that have been successfully infected with digeneans.^{50,117}

In contemplating the study of schistosome-snail interactions, including those studies outlined in Figure 3, it should be kept in mind that the relative amount of genetic diversity present in both digenean and snail laboratory populations is low, and has been skewed by artificial selection. Furthermore, natural populations of schistosome intermediate hosts such as *B. glabrata* from South America, or of *Biomphalaria pfeifferi* from Africa, typically do not exhibit wholesale resistance to infection with most or all *S. mansoni* miracidia. In contrast, in more natural settings, the success or failure of a particular schistosome-snail encounter depends on the genetic constitution of each participant,^{118,119} such that some combinations are compatible and some are not, e.g., parasite and host phenotypes are matched or mismatched. Thus the results of laboratory studies must be interpreted cautiously, and more studies using fresh field isolates encouraged. The underlying basis for phenotype matching remains unknown and building on the intriguing results of Roger et al^{86,87} should be a priority for future study.

ADDRESSING FUNDAMENTAL ISSUES IN IMMUNOLOGY USING GASTROPOD MODELS

Gastropods and their immune systems can also serve as useful models to consider important conceptual issues applicable to all organisms and the problems they confront with pathogens. The standard paradigm emphasizes the view that invertebrates elaborate a basic set of pathogen pattern recognition receptors coupled to standard responses like activation of phagocytosis or production of protective factors such as ROS, lectins or AMPs. An emerging challenge to this paradigm, in which studies of gastropods have figured prominently, has emphasized that the threats posed by the ubiquity, diversity and rapid evolvability of pathogens require a greater degree of sophistication from invertebrate defenses than previously considered.¹²⁰ It also notes that many invertebrates, including

some gastropods live for decades, and must be able to contend with pathogens that could “lock on” and decimate a host with a fixed immune repertoire.^{121,122}

Central to this discussion is the issue of the production of molecules with diverse recognition capability, such that broader spectra of pathogens, including those that may actively alter their own surface molecules or secretions, could be achieved. Lectins play an important role in nonself recognition in gastropods.¹²³ Vasta et al¹²⁴ note that even in the absence of unusual genetic rearrangements, a substantial degree of recognition diversity can be achieved because lectins exist in multi-gene families, can engage in alternative splicing, and the carbohydrate recognition domains of lectins exhibit considerable “plasticity” in their interactions with ligands and so might still bind multiple pathogens with slightly altered surface carbohydrates. A cursory examination of the *Lottia* genome indicates that several C-type lectins are present, and the same is almost certainly true for the *B. glabrata* genome, so these considerations are likely to apply.

Studies of fibrinogen-related proteins, or FREPs, from *B. glabrata*, which were first identified following exposure to digeneans, also exemplify some of the points of Vasta et al,¹²⁴ however, they differ in one important regard, namely the presence of an unforeseen mechanism of genetic rearrangement. FREPs are circulating hemolymph proteins that as presently understood are comprised of either one or two N-terminal immunoglobulin domains connected via a short interceding region to a C-terminal fibrinogen domain.¹¹⁷ They exist in a large family comprised of 14 identifiable sub-families. FREPs have calcium-dependent lectin activity, and bind soluble digenean antigens, or to the surfaces of digenean sporocysts and bacteria.⁹⁵ Recent microarray studies suggest they are highly responsive to digenean infection, some being up-regulated, and others down-regulated. In our experience, the persistent up-regulation of FREP4 is effectively a marker of digenean infection in *B. glabrata*.⁵⁰

In addition to their multiple subfamily representation, FREPs are further diversified by alternative splicing, and because at least some FREPs exist in their native configuration as multimers,⁹⁵ this represents a further opportunity for diversification, as has been noted for fibrinogen-related defense molecules in mosquitoes.¹²⁵ Additionally, study of the FREP3 subfamily surprisingly provided evidence for somatic diversification resulting from a combination of point mutations and putative gene conversion events acting on a relatively small number of source sequences,¹²⁶ the latter process favored by tandem arrangement of FREP3 genes, as has been noted from recent studies of BAC architecture (Adema, personal communication). Interestingly, this diversification process has either not been noted or is far less prominent for members of other FREP subfamilies such as FREP2 and 4, which show levels of variation more in line with expectations of normal allelic variation or representation from a small number of germline loci. This further suggests the FREP3 diversification noted is not artifactual (Adema, personal communication). One noteworthy aspect of FREP3 somatic diversification is that it provides a potential mechanism to generate diversity among different individual hemocytes¹¹⁷ and raises the possibility that not all gastropod hemocytes “are created equal”, offering a potential means to greatly increase the immune recognition repertoire.

The specific combination of IgSF and fibrinogen domains has been found in other planorbid snails such as *Helisoma trivolvis*, and recently two bonafide FREPs have been reported from *Aplysia californica*,¹²⁷ extending their representation to include another major lineages within the heterobranch clade. The *Lottia* genome contains an estimated 70 fibrinogen-encoding genes,⁹⁵ but they are not believed to be associated with IgSF domains. In general, gastropod studies have helped draw attention to the importance of

the fibrinogen domain which is proving to be common among other invertebrate groups as well, frequently being implicated in defense responses.¹¹⁷

Recent studies of *S. mansoni* miracidia and sporocysts indicate they have a remarkable system for generating diversified mucin molecules that are released only in the molluscan host, such that each individual miracidium is potentially unique in the spectrum of mucins produced.^{86,87} Heterogeneity in glycotope expression within and among *S. mansoni* miracidia has also been noted by Peterson et al.⁹⁷ These findings serve to accentuate the point that even metazoan parasites can present a highly diversified face to the gastropod defense system. Although the function of diversified mucins is presently unknown, it has been speculated they could potentially interact with FREPs, possibly serving as a smokescreen to divert host recognition efforts away from the parasite, thus potentially offering an explanation for why schistosome-snail compatibility depends on the specific genetic composition of the schistosome being matched to that of the snail.^{86,87} Although this is an attractive hypothesis, much more work is needed to determine if digenean mucins and host FREPs actually interact with one another, and how these systems might work in the more relevant test bed of naturally occurring snails and schistosomes.

A number of other lines of evidence also indicate that gastropods can produce diversified defense molecules. Bouchut et al⁷³ in a study of genes involved in hemocyte adherence or extracellular filament formation in *B. glabrata* found evidence for point mutations in six of the genes studied, but only if transcripts originated from hemocytes as opposed to whole bodies, and concluded "it raises the question of the existence of a gene wide diversification process taking place preferentially (or exclusively) in hemocytes of *B. glabrata*." Ittiprasert et al¹⁰² noted that *B. glabrata* mucin transcripts were heterogeneous and commented they could match the heterogeneity of *S. mansoni*-produced mucins. In their study of *B. glabrata* egg mass fluids, Hathaway et al⁷⁵ noted that the most abundant protein LBP/BPI existed in variant forms, and at least six similar yet variant C-type lectins were present. Zhang et al⁹⁵ noted the presence of multiple forms of both PGRPs and GNBPs from *B. glabrata*, and in a study of hemocyte-mediate nerve repair in *L. stagnalis*, Hermann et al⁴⁰ speculated that different integrins may exist in different combinations on hemocytes thus creating diverse populations. In a similar vein, recent studies of the bivalve *Mytilus edulis* have revealed that one category of AMPs, myticin C, exhibits extraordinary variation within and among individuals, and suggested that all the variants were generated from more common sequences by an as yet unknown mechanism.¹²⁸ Perhaps the presence of molluscan mobile genetic elements such as *Nimbus*¹²⁹ can help to explain some of the genetic rearrangements these results imply. It is important to remain cautious about the real implications of all the studies documenting diversification in recognition capacity as much more study is needed to document the exact mechanisms involved, and to validate their actual relevance in terms of protection from pathogen challenge.

Gastropods also provide an excellent opportunity to teach us about another fundamental aspect of invertebrate defense: by virtue of prior exposure to immunological stimuli, can defense responses be induced to quantitatively or qualitatively different levels that diminish susceptibility to subsequent pathogen challenge in a manner actually relevant to the animal? This topic is fraught with controversy among invertebrate immunologists and caution is required to assure that outcomes attributable to immunological priming or memory actually have an immunological basis.¹³⁰ One of the most compelling set of studies addressing this general topic was provided by Lie et al¹³¹ who showed that by first exposing *B. glabrata* to irradiated echinostome miracidia, those same snails would

subsequently be resistant upon challenge with normal miracidia up to 10 days later. By contrast, control snails exposed only to normal snails readily became infected. The study was all the more remarkable for showing a degree of specificity: snails with acquired resistance to one echinostome species had partial resistance to other echinostomes, but remained fully susceptible to *S. mansoni*. Snails sensitized twice and then challenged showed an even faster time course of parasite destruction, and a role of humoral factors in immobilizing parasites and making them vulnerable to encapsulation was invoked.¹³² They concluded acquired resistance was “not due to increased numbers of circulating amebocytes, but rather to intrinsic changes in the recognition and probably also killing capacity of amebocytes”.¹⁰⁴ This model system awaits further study and has the potential to help us define the limits of what gastropods, and invertebrates in general, are capable of with respect to priming of their defense systems.

OPPORTUNITIES TO EXPAND THE RELEVANCE OF GASTROPOD IMMUNITY

The primary focus of this chapter has been on basic gastropod immune mechanisms, often as revealed by exposure to digenetic trematodes. However, there are several additional more applied contexts in which an understanding of gastropod immunology is useful, and that provide additional opportunities to justify studies of this topic. Some of these are highlighted and summarized in Table 2.

CONCLUSION

The current state of gastropod immunology is marked by anticipation of the insights soon to be forthcoming from genome projects and by—thanks to the development of a number of new tools and approaches—the possibility of better understanding a backlog of interesting immune candidates that await rigorous testing with respect to their functional significance. The study of gastropod immunology currently is in need of approaches such as RNAi to enable more comprehensive functional testing. RNA knockdown can be achieved in *B. glabrata* by injection of dsRNA of target genes,¹³³ but the degree of knockdown needs to be higher so we can draw clearer conclusions from such experiments, and the consistency with which significant knockdown can be achieved improved. Using a different approach based on injection of multiple short dsRNAs, knockdown of FREP2 has again been achieved and is being further investigated (Hanington, personal communication). RNAi has been reported in *L. stagnalis*¹³⁴ and bivalves,^{81,135} so progress can be anticipated on this front.

Except for recent studies documenting viruses in abalones (Table 1), we are almost completely ignorant with respect to the diversity and impacts of gastropod viruses and the nature of the immune response they engender. Availability of the *Biomphalaria glabrata* embryonic (Bge) cell line as a means to search for viruses from natural populations of planorbid and possibly other snails should be encouraged, both to find viruses that might be potential control agents for medically important snails and to harvest viruses which can then be used to probe basic gastropod defense responses. Another need is to develop more comprehensive overviews of gastropod transcriptional profiles, which should be enabled as genome sequences projects are completed. Representation of the

Table 2. The following topics represent important ways in which the scope and relevance of studies of gastropod immunobiology can be expanded

Maintaining Gastropods as Sources of Food

Learning how abalones respond to viruses^{13,153} or bacteria,⁶³ to improve abalone culture

Control of Snail-Transmitted Human Parasites

Replacing snails in natural habitats susceptible to *S. mansoni* with snails that are resistant, thus achieving biological control of schistosomiasis^{159,160}

To Identify Compounds with Potential Biomedical Relevance

Antimicrobial peptides, like abhisin from the disk abalone *Haliotis discus discus*,¹⁴⁸ may have activity against bacteria, fungi and cancer cells.

To Better Protect Endangered Gastropods, Including those Threatened by Disease

Perhaps an improved understanding of gastropod immunity could have been used to save the land snail *Partula turgida* from extinction at the hands of a microsporidian parasite¹⁶¹

Elimination of Invasive Gastropods

A thorough understanding of the immunobiology of invasive gastropods could lead to approaches to exploit co-evolved or new pathogens in their control.^{162,163}

To Return Balance to Ecosystems

Improve abalone health and abundance to provide normal food sources for endangered sea otters¹⁶⁴

As Models to Understand Commitment of Internal Defense Systems to Symbionts

The bacterial associates of gastropods are poorly known but range from near monocultures of specialized bacteria⁹ to surprisingly diverse floras in the guts of diet generalists like *Biomphalaria* or *Helisoma*.¹⁹ The nature of the accommodations made to symbionts in such diverse cases is unknown.

As Models to Study the Impact of Pollutants or Other Stressors on Immune Function, and to Develop Biomarkers for Stress

Snails with shells fouled by epibionts were also more likely to be infected with digeneans, and though multiple explanations might exist, interactions between stressors are likely.¹⁶⁵

Snails stressed by fasting, exposure to cold or to detergent became more susceptible to trematode infection¹⁶⁶

Toxin exposure may influence susceptibility to pathogens like trematodes.^{103,167} Exposure to herbicides such as fomesafen¹⁶⁸ or organic chemicals¹⁶⁹ may diminish defense capacities Use of selenium-dependent glutathione peroxidase as a possible biomarker for physical stress in abalones¹⁷⁰

continued on next page

Table 2. Continued**To Investigate the Impact of Mounting Immune Responses on Life History Attributes**

Repeated exposure to bacteria necessitating immune responses affected short-lived and long-lived land snails differently, with long-lived species checking growth and mating frequency, thus potentially favoring long-term survival¹⁷¹

B. glabrata snails resistant to *E. caproni* exhibit delayed maturity compared to susceptible snails¹⁷² Rigby and Jokela (2000)¹⁷³ showed that immune response measured by increase in hemocytes trades off with reproduction and survival in snails, and predator avoidance may modify expression and costs of immune defense.

To Examine the Interface between Immunity and Behavior

Zbikowska (2004)¹⁷⁴ found that snails infected with digeneans tended to select lower temperatures than uninfected controls, and postulated this slowed down the rate of parasite production and gave the snail time to respond to the damage produced. No “behavioral fever” was noted.

To Monitor Climate Change and Record its Impacts

A 1°C temperature change increases abalone mortality due to *Vibrio harveyi*¹⁸

Different abalone species respond differently to withering syndrome with temperature change⁷⁵

transcriptomic repertoires on current snail microarrays is modest. More comprehensive arrays can be used to assess responses to different pathogen exposures, and how such responses might be modulated in snails from natural habitats, including those under environmental stress or contending with preexisting infections. Such studies may very well also reveal novel immune responses undertaken by components that today we can only classify as “unknowns”.

Thanks to the continuing availability of new genome sequences for gastropods and other invertebrates, in combination with further studies to assess the importance of various immune pathways to survival, we will begin to approach for the first time higher order questions pertaining to the evolution of immunity. For example, (1) does the presence of prominent lineage-specific pathogens—such as digeneans in snails—drive the development of unique immune capabilities not seen in other host groups?; (2) how similar are the immune systems of diverse gastropods, and do the patterns we discern among their defense systems more strongly reflect a phylogenetic signal, or do they diverge more as a consequence of their immediate ecological challenges?; (3) as previously argued,¹³⁶ are the internal defense systems of gastropods and bivalves fundamentally different, with the former showing narrower recognition diversity than the latter?; and finally, (4) do the immune capacities of major lineages, such as at the phylum level, evolve along similar trajectories, or has selection placed a premium on the development of novel immune solutions that prevent easy pathogen transfer from one group to the next? Exciting times lie ahead, and as noted at the beginning of this chapter, we must take care not to overgeneralize the gastropod case for no doubt, embraced within this ancient and diverse lineage, are fascinating alternative solutions to the problem of achieving effective internal defense.

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CHAPTER 3

BIVALVE IMMUNITY

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Abstract: Bivalves are comprised of animals unclosed in two shell valves, such as mussels, oysters, scallops and clams. There are about 7,500 bivalve species and some of them are of commercial importance. Recently, interest in bivalve immunity has increased due to the importance in worldwide aquaculture and their role in aquatic environmental science and their position in phylogenetic research. This chapter provides a short review of bivalve immunity, including cellular and humoral immunity and the key components and the interactions involved in humoral immunity.

INTRODUCTION

The phylum Mollusca is one of the most large, various and important groups in the animal kingdom. The monophyletic Mollusca is sister to the clade that unites annelids with nemertean, phoronids and brachiopods.¹ Together with platyhelminthes, they constitute Lophotrochozoa and further constitute Protostomia with Ecdysozoa (molting animals).¹ Bivalvia has about 7,500 species and was the second most diverse class of molluscs after Gastropoda. Many of them are sources of seafood and are important for pearl production and therefore of great commercial importance. In addition, as sedentary filter feeders, bivalves may concentrate bacteria, viruses, pesticides, industrial wastes, toxic metals and petroleum derivatives, making them important markers for biomonitoring pollution in aquatic ecosystems and ideal species for investigating the effects of environmental contaminants.²

In the long course of evolution, bivalves have developed an array of effective strategies to protect themselves from the attacks of various pathogens and environmental stresses. Interest in bivalve immunity has increased continuously in recent years due to

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serious diseases and mortality problems threatening the healthy development of bivalve aquaculture. Although pathology information has been accumulating, research on bivalve immune systems and the underlying molecular mechanisms are still at an early stage, with only some investigations in a comparatively small number of species.^{3,4}

As invertebrates, bivalves rely exclusively on an innate, nonlymphoid system of immune reactions.⁵ The internal defense of bivalve is mediated by both cellular and humoral components. The former includes phagocytosis or encapsulation, with subsequent pathogen destruction via enzyme activity and oxygen metabolite release, while the latter includes various reactions mediated by series of molecules.⁶ Here we review our current knowledge of bivalve immunity, mainly focusing on immune recognition, signal transduction and effector synthesis involved in cellular and humoral immunity.

HEMOCYTES AND PHAGOCYTOSIS

Bivalves have an 'open' circulatory system in which the hemolymph, passing out of the open ends of arteries, bathes all the organs before returning to the heart by the way of sinuses and respiratory structures (gills). Hemocytes, the circulating cells of bivalves, are primarily responsible for the defense against pathogens.⁷ Phagocytosis and encapsulation are two major mechanisms for hemocytes to eliminate nonself substances and dead cells.⁸ In addition, antibacterial effectors, opsonins, nonspecific hydrolysis and toxic oxygen intermediates found in bivalve hemolymph together coordinate immune response.

The assortment of bivalve hemocytes has been controversial for a long time. Based on morphology and histochemistry research results from mussels and clams, bivalve hemocytes are generally categorized into hyalinocytes and granulocytes,⁹⁻¹¹ and the latter can be further subdivided into eosinophilic granular hemocytes and basophilic granular hemocytes.^{6,11} There are also other hemocyte types in some species. For instance, the Type 'III' eosinophil, morula-like cells and blast-like cells were found in *Cerastoderma edule*, *Tridacna derasa* and *Scrobicularia plana* hemolymph, respectively.^{6,12} Among all these cell types, granular hemocytes are the most numerous, rich in a variety of hydrolytic enzymes and the major cell, executing the process of phagocytosis.¹⁰

Phagocytosis is a process to recognize and ingest nonself molecules and cell debris. Bivalve hemocytes can engulf a variety of particles including bacteria, algae, yeast, foreign blood cells and latex spheres.⁷ The first step of phagocytosis is the attachment of phagocyte to the targeted particle.¹³ Many studies have revealed that hemocytes of many bivalve species exhibit chemotactic as well as chemokinetic reactions, type of which is dependent upon the nature of the molecules presented.¹⁴ In the mussel *Mytilus edulis*, lipopolysaccharides (LPS) from both *Serratia marcescens* and *Escherichia coli* stimulated the migration of cells. In European flat oyster *Ostrea edulis*, hemocytes migrated from circulatory system to connective tissues after *Bonamia ostreae* infection.¹⁵ Hemocytes of *Mercenaria mercenaria* migrated toward not only peptides or small proteins secreted by both Gram-positive and Gram-negative bacteria. After chemotactism, the phagocytes adhere to pathogens, followed by cytoskeleton modification, internalization and destruction of the engulfed target within phagosomes.¹³ In *Mytilus galloprovincialis*, both hyalinocytes and granulocytes could execute phagocytosis by formation of coated vesicles and uncoated endocytic vesicles,¹⁰ and cells of different morphology presented different levels of phagocytosis towards zymosan, latex beads and bacteria.¹⁶

After the nonself is phagocytosed, the phagosomes and lysosomes fuse together and the engulfed target is destroyed within phagosomes by lysosomal enzymes, reactive oxygen species (ROS), nitric oxide (NO) as well as antimicrobial factors. During phagocytosis, the release of degradative enzymes for the destruction of foreign material is accomplished by a sudden release of ROS within hemocytes, which is referred to as respiratory burst. ROS act as killing agents, either alone or in combination with lysosomal enzymes and are important in the phagocyte-mediated killing of microorganisms.¹⁷ Studies on oyster *Crassostrea ariakensis* revealed that granulocytes were most active in spontaneous ROS production¹². In clam *M. mercenaria*, hemocyte oxidative burst was active in response to the stimulation of zymosan and bacterial extracellular products, accompanied by the increase of ROS production in hemocytes.¹⁸ The internalization of different phagocytic targets and the production of ROS and NO in *M. galloprovincialis* were found to be blocked by inhibition of phosphatidylinositol 3-kinase, protein kinase C and extracellular signal-regulated kinase.¹⁶ Furthermore, many other hemolymph factors also help phagocytosis, including agglutinins (e.g., lectins) and various antimicrobial peptides. Bivalve lectins have different carbohydrate-binding specificities and are involved in nonself-recognition.¹⁹ Antimicrobial peptides are engaged in the destruction of bacteria inside the phagocyte before being released into hemolymph to participate in systemic responses.⁷ These factors will be further discussed in the following parts.

Another mechanism of invertebrate cellular immune response is encapsulation.¹³ In invertebrates, encapsulation is the common immune defense reaction for foreign bodies which are too large to be phagocytosed. In general, a capsule of hemocytes encloses the foreign body (e.g., multicellular parasites) and cytotoxic products (e.g., degradative enzymes and free radicals) are released by the hemocytes in an attempt to destroy the invader. In the encapsulation response of *Crassostrea gigas* against copepods *Mycicola ostreae*, the copepods were observed on the gill surface of *C. gigas* engulfed by a massive agglomerate of hemocyte-like cells encircled by a thin layer of fibroblast-like cells.²⁰ The clam *Dreissena polymorpha* also employed such a defense mechanism when infected with trematode *Bucephalus polymorphus*.²¹ Studies on *C. edule* revealed that positively charged targets stimulated the most vigorous response and that nonspecific electrostatic forces and humoral plasma factors have a synergistic role in hemocyte attachment and the encapsulation response. Phagocytosis and encapsulation are important and complicated processes, the involved molecules and the detailed mechanism should be addressed in future studies.

IMMUNE RECOGNITION

Immune recognition is the first step in activating immune response and occupies a very important position in the immune system to discriminate nonself from self substances. The immune responses begin when specialized, soluble or cell-bound Pattern Recognition Receptors (PRRs) recognize (and bind to) the major targets, called Pathogen-Associated Molecular Patterns (PAMPs).^{22,23} PAMPs are common in microorganisms but rare or absent in host animals, such as LPS or peptidoglycan (PGN) in bacterial cell walls and β -1,3-glucan on fungal cell walls.²⁴ Invertebrates rely only on innate immunity and develop a sophisticated system of PRRs. Seven groups of distinct PRR are identified in bivalves, including peptidoglycan recognition proteins (PGRPs), Gram-negative binding proteins (GNBPs), C-type lectins, galectins, thioester-containing proteins (TEPs), scavenger

receptors (SRs) and Toll-like receptors (TLRs). In the following, we will briefly introduce the bivalve PRRs, focusing on their binding specificity and functions.

Peptidoglycan Recognition Protein

PGRP is a member of PRRs that specifically bind to PGN, a unique cell wall component of all virtual bacteria but not present in eukaryotic cells. The knowledge of invertebrate PGRPs comes mainly from insects. These PGRPs played a central and diverse role in activating immune reactions, such as melanization cascade, phagocytosis and activating the Toll or IMD signal transduction pathways for the production of antimicrobial products to hydrolyze peptidoglycan and protect host against infection. Some bivalve PGRPs have been identified from the Pacific oyster *C. gigas* (*CgPGRP-S1S*, *-S1L*, *-S2*, *-S3* and *CgPGRP-L*),²⁵ bay scallop *Argopecten irradians* (*AiPGRP*)²⁶ and Zhikong scallop *Chlamys farreri* (*CfPGRP-S1*).²⁷ The identified bivalve PGRPs were all short type with a conserved amidase PGRP domain in their C-terminus. Interestingly, there was an additional goose-type (g-type) lysozyme domain in *CgPGRP-L*, while a defensin-like domain was present in both *CgPGRP-S1S* and *CgPGRP-S1L*.²⁵ The recombinant protein rCfPGRP-S1 from scallop could bind not only with PGN but also with chitin and LPS moderately. More importantly, rCfPGRP-S1 exhibited strong activities to agglutinate Gram-positive bacteria *Micrococcus luteus* and *Bacillus subtilis*, while slightly with the Gram-negative bacterium *E. coli* (unpublished data). The short type PGRP is widely present in various bivalve species and seems to be a versatile PRR not limited to the function of recognizing and binding the PAMPs and thus an indispensable component in bivalve innate immunity.

Gram-Negative Binding Protein

The GNBP family includes members that bind Gram-negative bacteria, LPS and β -1,3-glucan.²⁸ Research on bivalve GNBP is still limited and they have only two related reports. The LPS and β -1,3-glucan binding protein (LGBP) is one of the GNBP with various biological functions, including the activation of the prophenoloxidase (pro-PO) system, cytolysis, bacterial aggregation and opsonic reaction. The LGBP gene was cloned from scallop *C. farreri* (*CfLGBP*) with conserved domains of the LPS-binding site and glucan-binding site. The initial up-regulation after *Vibrio anguillarum* challenge indicated that *CfLGBP* was sensitive to bacterial infection.²⁷ The recombinant CfLGBP could bind not only LPS and β -glucan, but also PGN and exhibited obvious agglutination activity towards Gram-negative bacteria *E. coli*, Gram-positive bacteria *B. subtilis* and fungi *Pichia pastoris* in vitro. Additionally, a beta-1,3-glucan binding protein (β GBP) was purified from the plasma of marine mussel *Perna viridis*²⁹ with an inherent serine protease activity. It agglutinated bakers yeast, bacteria and erythrocytes and enhanced proPO activity of the plasma. Although its gene sequence and molecular structure are still unknown, β GBP is thought to be a multifunctional molecule and functions as a recognition molecule for beta-1,3-glucan on the surface of microbial cell walls.

C-Type Lectin

C-type lectins are a superfamily of diverse proteins with one or more carbohydrate-recognition domains (CRDs) of ~130 amino acid residues. They recognize

and bind to terminal sugars on glycoproteins and glycolipids and function in nonself recognition and clearance of invaders.³⁰ In bivalves, lectins are undoubtedly involved in nonself recognition and also have agglutination or opsonic roles in hemocyte phagocytosis. We will quote those lectins in immune recognition of bivalve in the following and adduce their agglutination or opsonic roles later.

The forefront research on C-type lectins mainly focused on characterizing the features or detecting their activities. With the development of molecular biotechnology, much more attention was paid to their gene structure and recognition mechanism. Recent EST analysis in scallops, clams and oysters revealed a high content and variety of lectin gene homologues.²⁷ Some C-type lectins or proteins with similar activities have been characterized from bivalves.³¹ The mRNA expression of these genes could be significantly up-regulated by bacteria or parasite, especially bacteria of *Vibrio* genus, suggesting that they were involved in the immune response against invading microbes.

Bivalve C-type lectins are diverse in their domain structure and organization. Besides most lectins with single CRD, there are also three and four CRDs in Cflec-3 and Cflec-4 from *C. farreri*, respectively, while most insect C-type lectins contain tandem CRDs.³² The architecture and phylogenetic analysis of these proteins together with those from *Drosophila* and *Caenorhabditis elegans* suggested that multidomain C-type lectins in different lineages did not arise from a common multidomain progenitor and these proteins served distinct functions in different animal lineages.

Bivalve C-type lectins have different carbohydrate-binding specificities and are believed to be a kind of antibodies in nonself-recognition. For instance, in clam *Ruditapes philippinarum*, MCL can bind to the surfaces of purified hyphospores and zoospores of *Perkinsus olseni* parasite by recognizing and binding the terminal GalNAc/Gal residues. Meanwhile, the hemo-agglutinating activity of MCL3 could be inhibited by GalNAc, Mannose, lactose, raffinose polysaccharides bovine mucin Type II and *Candida* mannan.³³ These two lectins, as well as MCL-4, contributed to the phagocytic ability to eliminate bacteria or parasite via recognition of terminal carbohydrate residues on the surface of microbes.³⁴ Chiletin from *Ostrea chilensis* could agglutinate sheep red blood cells through binding galactose and mannose.³⁵ The EPN motif in the canonical binding sites of Codakine from clam *Codakia orbicularis* proved to be important for calcium-dependence and mannose/GlcNAc-binding activity.³⁶ In some species, the lectins with similar carbohydrate-binding specificity may distinguish different invading microbes. For example, the recombinant C-type lectins Cflec-1, Cflec-2, Cflec-3 and Cflec-5 from *C. farreri* agglutinated *E. coli*, *Staphylococcus haemolyticus*, *Pseudomonas stutzeri* and *P. pastoris*, respectively, though they all possessed mannose-binding specificity.

The great number of C-type lectins identified from bivalves shared similar structural features and displayed binding and agglutinating activities towards a range of microbes. Since there is no antibody-mediated immunity in invertebrates, abundant lectins with diverse expression profiles and bioactivities might function as nonclonal effectors in the bivalve immune system.

Galectin

Galectins are a family of β -galactoside-binding lectins and they are probably the most conserved and ubiquitous lectin family found in multicellular organisms.^{22,23,28} The bivalve galectin was purified first from oyster *Pinctada fucata martensii* in the 1980s,

but the genes were cloned and identified recently, including *CvGal* from *Crassostrea virginica*³⁷, *CgGal* from *C. gigas*,³¹ *AiGal1* from *A. irradians*,³⁸ *MCGal* from clam *R. philippinarum*³⁹ and *Pf-galectin* from *Pinctada fucata*. Except for *CgGal*, which contains a single CRD, the other four galectins possess multiple CRDs. *AiGal1*, *CvGal* and *Pf-galectin* are quadruple-CRD galectins, which are so unique in bivalves that it has never been reported in other species. From a phylogenetic point of view, all four CRDs of *AiGal1*, *CvGal* and *Pf-galectin* form a single clade, suggesting that the bivalve galectin CRDs share a common ancestor and the four individual CRDs of each galectin are originated by repeated duplication of a single galectin gene.

CvGal and *MCGal* are two major galectins for functional study. *CvGal* could facilitate recognition of a variety of potential microbial pathogens, unicellular algae and preferentially *Perkinsus marinus* trophozoites. Attachment and spreading of hemocytes to foreign surfaces induced localization of *CvGal* to the cell periphery, its secretion and binding to the plasma membrane. *CvGal* subsequently promoted phagocytosis for both potential infectious challenges and phytoplankton components.³⁷ Moreover, *MCGal* had an affinity towards galactose and N-acetylgalactosamine and could bind to the surface of *Perkinsus olseni* and agglutinate *Vibrio tapetis* in vitro.³⁹

Thioester-Containing Protein

TEPs are a family of proteins characterized by the unique intrachain β -cysteinyl- γ -glutamyl thioester bond and a propensity for multiple conformationally sensitive binding interactions.⁴⁰ This protein family consists of complement components C3, C4, C5, protease inhibitor alpha₂-macroglobulin (α_2 M), CD109 and a set of insect TEPs. Among them, the invertebrate C3-like molecules and insect TEPs were thought to be involved in the innate immune defense as PRRs.²⁸

The bivalve TEPs were recently identified from the clam *Ruditapes decussates*⁴¹ and scallop *C. farreri*,⁴² termed as *Rd-C3* and *CfTEP*, respectively. They both contained canonical thioester motif GCGEQ, proteolytic cleavage sites and catalytic histidine residues similar to C3 molecules. However, *CfTEP* possessed additional features distinguished it from C3 molecules, including: (1) the absence of anaphylatoxin-like and C345C domains, (2) a distinctive cysteine signature in the C-terminus which characterized the TEP subfamily apart from complement factors and α_2 M subfamilies and (3) the highly variable central region.⁴² Due to these structural differences, *CfTEP* was phylogenetically related to the insect TEPs, while *Rd-C3* was related to the invertebrate C3-like molecules. The above results supported the view that TEP and complement factors shared a common ancestor but they separated from each other at a rather early lineage.⁴³

The genomic organization of *CfTEP* was similar to human and mouse C3 rather than ciona C3-1 and *Drosophila dTEP2*, indicating a complicated evolutionary history of this gene family. It was of great interest that seven different *CfTEP* transcripts were produced by alternative splicing and they displayed different expression patterns in gonads in response to different bacterial challenges, which suggested an important role of diverse *CfTEP* transcripts in the innate immune defense of scallops.⁴⁴ These results provided new insights into the role of TEPs in bivalve immune responses, as well as the evolutionary origin of this important, widespread and functionally diversified family of proteins.

Scavenger Receptor

Scavenger receptors (SRs) are a main type of endocytic receptors with multifunctions to recognize and engulf various PAMPs.⁴⁵ In contrast with other PRRs, the information about invertebrate SR is extremely limited. There are only two invertebrate SRs identified from *Drosophila* (*dSR-CI*) and *C. farreri* (*CfSR*, GQ260639.1). CfSR is structurally different from all the characterized SRs. It contains six scavenger receptor cysteine-rich (SRCR) domains absent in dSR-CI and UPAR-like and ShK toxin-like domain do not exist in any other member of SRs. CfSR shares a similar attachment site with anchor protein Sgp-2 and it was mainly detected on the outer surface of hemocytes by immunofluorescence approach, indicating that CfSR was anchored on the outer-membrane of cells. The recombinant CfSR displayed a significantly strong activity to bind not only acetylated LDL, dextran sulfate but also with various PAMPs, including LPS, PGN, zymosan particle and mannan.

CfSR is one of the most primitive SR found so far in invertebrates. It displays unique structure and broader ligand binding ability. The existence of SR protein in bivalve will contribute not only to the origin and evolution of the diverse molecules, but also to the understanding of the complex mechanism of immune recognition.

Toll-Like Receptor

TLRs are an ancient family of proteins with the hallmark structure of extracellular leucine-rich repeats (LRRs), intracellular Toll/Interleukin-1 receptor (TIR) domains, which play key roles in detecting various nonself substances and then initiating and activating immune system. The reports of TLRs in mollusc are still rare although they were widely distributed in nearly all animal phyla. Only one TLR gene (*CfToll-1*) and three TLR EST fragments had been identified in *C. farreri*, *C. virginica*⁴⁶, *A. irradians*⁴⁷ and *M. mercenaria*.⁴⁸ *CfToll-1* shared the same domain architecture with *D. melanogaster* Tolls (*DmTolls*) and *Tachypleus tridentatus* Toll (*tToll*). The mRNA expression of *CfToll-1* was upregulated by LPS in a dose-dependent manner.⁴⁹ The finding suggested that *CfToll-1* might be involved in immune response against bacterial invasion. Considering their importance in innate immunity, the study of the number, assortment and the roles of bivalve TLR in immune recognition and signal transduction (discussed in other section) should be addressed in the future.

IMMUNITY SIGNALING PATHWAYS

Innate immune system is under the control of a complex network of evolutionary conserved signaling pathways, which are activated depending upon different invasions or stimuli. Several signaling pathways, such as TLR, the Janus kinase/signal transducer and activator of transcription (JAK-STAT), mitogen-activated protein kinase (MAPK) and NF- κ B pathways have been extensively studied in recent years for their important roles in regulating the immune system in both vertebrates and invertebrates. In bivalves, effective immune defense systems have been developed during evolution which can protect them from infection successfully. When bivalves are challenged by a pathogen, different signaling pathways triggered by PRRs induce the systemic immune response and produce responding effectors. Although the studies are preliminary, some genes

involved in conserved signaling pathways have been identified in bivalves. In this section, we will briefly introduce what is currently known about these signal transduction pathways, which may be a crucial step for our understanding about bivalve immune system.

The Canonical NF- κ B Signaling Pathway

NF- κ B signaling pathway is an evolutionarily conserved process to activate NF- κ B,⁵⁰ a nuclear factor which has a central role in coordinating the expression of a wide variety of genes that control immune responses.⁵¹ In resting cells, NF- κ B proteins bind to the inhibitor of κ B (I κ B). The NF- κ B signaling pathway is activated once I κ B is degraded. This degradation of I κ B is catalyzed by serine kinase I kinase (IKK) which leads to the translocation of released NF- κ B dimers to nucleus. The activated NF- κ B proteins then bind to DNA and activate gene transcription.⁵⁰

In recent years, the key molecules involved in bivalve NF- κ B signaling pathway, such as NF- κ B, I κ B, IKK genes, have been identified successively, offering clues for the existence of NF- κ B signaling pathway in bivalves. Now three I κ B genes have been cloned from scallops *C. farreri*²⁷ and *A. irradians*,⁴⁷ and oyster *C. gigas*,⁵² and they showed a high level of identity with insect I κ B-like proteins and vertebrate I κ B isoforms. The *Cg-Rel*, a gene encoding the first NF- κ B homologue in bivalve, shares the structural organization with Rel/NF- κ B family members. The C-terminal transactivation domain and κ B binding sites were indispensable for activation of the expression of genes controlled by NF- κ B pathway. The NF- κ B signaling pathways were well studied in pearl oyster *P. fucata*. Three genes, *Pf-Rel*,⁵³ *Pf-IKK*⁵⁴ and *pol κ B*⁵⁵, were cloned and they were constitutively and ubiquitously expressed in tissues of pearl oyster. LPS could transiently stimulate I κ B degradation, but couldn't influence the expression level of *Pf-IKK*. Transfection experimentation in NIH3T3 cells with *Pf-IKK* demonstrated that *Pf-IKK* triggered the gene expression by activating NF- κ B in an I κ B-dependent manner just as mammalian counterparts do.⁵⁴ The findings above favored the hypothesis that NF- κ B signaling pathway was an ancient scheme of immune gene regulation pathway, which was conserved in bivalves. Although the complex members and their detailed information of NF- κ B signaling pathway are not well understood in bivalve, the accumulating evidence indicates that it bears considerable similarity with mammalian NF- κ B signaling pathway.

MAPK Pathway

MAPK cascades are one of the most important signaling pathways controlling a variety of physiological processes including cell proliferation, growth, differentiation, cell death, innate immunity and development.⁵⁶ Three subfamilies of MAPKs have been well-characterized in multicellular organism. They are extracellular signal-regulated kinases (ERKs), c-Jun amino-terminal kinases (JNKs) and p38 MAPKs. The pathway represents a characteristic phosphorylation system in which a series of three protein kinases phosphorylate and activate one another.⁵⁷ MAPK pathways were proven to exist in bivalve by immunoblot or ELISA techniques.⁵⁸ The pathways are not only sensitive to various environmental stressful stimuli, but also can be activated by growth factor insulin, cytokines, hormone substances and bacterial challenge. Several ESTs homologous to the MAPK pathway components were screened from cDNA library of oyster (*Cg-MAPKK1*,

Cg-MAPKK2, *Cg-c-jun* and *Cg-phosphatase*, *Cg-focal*, *Cg-FAK*) and Manila clam (c-jun).⁵⁹ Further investigations in molecular structure and their functions are needed for the precise defining of all molecules involved in MAPK pathway of bivalves.

JAK-STAT Pathway

JAK-STAT pathway is one of the important signaling pathways, downstream cytokine receptors.⁶⁰ Although JAK-STAT pathway seems to be present throughout evolution, there is no such report regarding the existence of JAK or STAT molecule in bivalves. Nevertheless, STAT activation has been observed recently in *M. galloprovincialis* under the stimulation of cytokine and bacteria. The microbicidal activity against *E. coli* in mussel hemocytes increased significantly after the pretreatment with human recombinant IFN γ , while no effect was observed with IFN α . IFN γ induced a rapid and time dependent increase of phosphorylated STAT1-like protein, evaluated by polyclonal antibodies specific for the tyrosine phosphorylated sites of STAT1.⁵⁸ It was suspected that JAK-STAT pathway possibly existed in bivalves. More and more cytokines and their receptors found in bivalves (details in other section) provided the clue for further investigation of bivalve JAK-STAT pathway. The roles of JAK-STAT pathway and their relationship with the newly identified cytokines from these animals may be an interesting research field which is helpful to our comprehensive understanding of bivalve immunity.

The Toll-Like Receptors Signal Pathway

The characterization of TLR and its signaling pathway is one of the greatest propellers to acquaint with the immune system in the last decades. The receptors, adaptors and transducers in this pathway, such as TLR, myeloid differentiation factor88 (Myd88), tumor necrosis factor receptor-associated factor 6 (TRAF6) and IL-1 receptor-associated kinase (IRAK) displayed striking similarity from *Drosophila* to mammals, while the function of this pathway is different between human and fly. TLR pathway in mammals is a skillful system that detects invasion of various pathogens and plays a key role in bridging innate and adaptive immunity. Toll pathway in *Drosophila* is indispensable for both development and antifungal/anti Gram-positive bacteria immunity. It is worth noting that the absence of Myd88 orthologue and their very low similarity of TRAF and IRAK homologues strongly imply the evolutionary 'loss' of Toll signaling pathway in nematodes.⁶¹ Moreover, *tol-1* in *C. elegans* is not involved in immune response but only in development. More information about TLR pathway in other invertebrates is required for better understanding of its function and evolution.

Fortunately, the key components involved in TLR pathway, including TLR,⁴⁹ Myd88,^{62,63} TRAF6,⁶⁴ TRAF3,⁶⁵ and IRAK⁴⁶ have also been identified in bivalves. And the expressions of *CfToll-1*,⁴⁶ *CfMyd88*,⁶² *CfTRAF6*⁶⁴ in scallops were up-regulated by stimulation of LPS or PGN. Furthermore, when the *CfToll-1* was "knock down" by RNAi technique, the scallop was more susceptible to pathogen and the expression of downstream genes was also down-regulated. Undoubtedly, the canonical TLR signaling pathway existed in bivalve. Considering scallop is a relatively primitive animal, the presence of TLR signaling pathway is epochmaking to understanding its origin, evolution and crucial roles in innate immunity.

Complement Pathway

The complement system is a major component of vertebrate innate immunity and also an essential bridge between innate and adaptive immunity. It can be activated mainly through four separate pathways, referred to as the classical, alternative, lectin⁶⁶ and coagulation⁶⁷ pathways. Recent investigation on the evolution of complement system has demonstrated the origins of complement system could be traced to near to the beginnings of multi-cellular animal life and the components of complement system should have some evolutionary traces in inferior species. However, there are very few reports on complement system in invertebrates and even less in bivalves. To our knowledge, only two complementary factors, C3 (*Rd-C3*) and B factor-like (*Rd-Bf-like*) molecules have been identified as representatives of the alternative pathway from carpet-shell clam (*Ruditapes decussatus*). The Rd-C3 shares distinctive structural characteristics with complement proteins (C3/C4/C5). Rd-Bf-like is composed of two complement control protein modules (CCP domains) and shares about 30% similarity with other known Bf proteins.⁶⁸ Additionally, three complement-like factors, including *CfTEP*⁴² (details in other section), *CfC1qDC*⁶⁹ and *AiFREP*⁷⁰ were identified from scallops. TEP reported in scallop suggested that it might be an ancient equivalent component of the key factor, C3, and there was likely to be a primitive, simple complement-like system in bivalves. CfC1qDC is a novel C1q-domain-containing protein with LPS binding activity and is suspected to be a candidate of classical pathway in scallop. AiFREP, a member of protein with fibrinogen-like (FBG) domain at the C-terminus, exhibited a very high similarity with FBG domain of mammalian ficolin (at 61% approximately with *Rattus norvegicus*) and other invertebrate FREP proteins. *AiFREP*, the EST fragment of MBL and ficolin from oyster and kinds of lectin in bivalves indicated the high existent probability of lectin pathway. The results place us in a puzzle about the bivalve complement system and its activation.

IMMUNE EFFECTOR

Innate immunity encompasses a complex array of defense reactions, in which immune effectors are fundamental molecules and utilized as executor for the incapacitation and elimination of invaders. The immunity of bivalves relies upon the production of immune effectors that are active against a large range of pathogens or sensitive for the environmental stress. The importance of those molecules is underlined by increasing outbreak of a variety of diseases and summer mortalities in bivalve aquaculture. The following details recent knowledge of immune effectors, such as antimicrobial peptides (AMPs), cytokines, complement components, antioxidant enzymes, acute phase proteins and draws their repertoire in bivalve immune response.

Antimicrobial Peptides

AMPs have been characterized as one of the key immune effectors in innate immunity and are widespread in plants and animals. All the known AMPs are classified into four structure groups according to their amino acid sequences, secondary structures and functional similarities: (1) the linear basic peptides forming amphipathic α -helices

conformation and deprived of cysteine residues; (2) peptides containing cysteine residues with one to six intra-molecular disulphide bonds; (3) peptides rich in regular amino acids like proline with a variable structure; (4) the peptides produced by the hydrolysis of large inactive or proteins with little activity. Approximately 20 AMPs have been identified from bivalves, with 10 from *M. edulis* and *M. galloprovincialis*, which were organized into four distinct groups according to their antimicrobial action based on their hydrophobic and cationic properties and the amphipathic structure: MGD (defensin), mytilin, myticin and mytimycin.^{71,72}

MGDs (MGD1 and MGD2) characterized from *M. galloprovincialis* with eight cysteines showed high similarities to the arthropod defensin family, which contains 6 cysteines. They were believed to be the original members of the arthropod defensin family because of the presence of two extra cysteines.⁷³ MGD1 and MGD2 share significant homology with the high conserved cysteines, hydrophilic and hydrophobic residues,⁷² and they are essentially active against Gram-positive bacteria, including some pathogens for marine invertebrates. There are five kinds of mytilin isoforms in the bivalve mussels. Mytilin A and B were isolated from *M. edulis* plasma and mytilin B, C, D and G₁ were isolated from *M. galloprovincialis* hemocytes. Mytilin A, B, C, D exhibited significant activities against both Gram-positive and Gram-negative strains, while mytilin G₁ was only active on Gram-positive bacteria. Mytilin B and D were also active against the filamentous fungus *Fusarium oxysporum*. Mytilin C was tested against the protozoan parasite, *P. marinus*.

Myticins were identified as novel cysteine-rich peptides. At present, three classes of myticin, A, B and C have been reported. Only one isoform has been described for myticin A and B and a total of 74 different isoforms have been reported for myticin C so far.⁷⁴ Myticin A and B were purified from the hemocytes and the former was also from the plasma of *M. galloprovincialis*. The mature peptides of myticin A and B comprise 40 residues with four intra-molecular disulfide bridges and a cysteine array in the primary structure different to that of the previously characterized cysteine-rich AMPs. The two myticins had marked activity against the gram-positive strains, *M. luteus*, *Bacillus megaterium* and *Aerococcus viridans*. Myticin B was also active against the filamentous fungus, *F. oxysporum* and moderately against the Gram-negative, *E. coli* D31. Myticin C had a high variability on the nucleotide sequence. It was a ubiquitous peptide expressed early in the development and associated with mussel survival. Different myticin C isoforms were observed along their life when mussel touched with new potential pathogens or nonself molecules in general. Therefore, myticin C could deal with a huge range of potential pathogens present in the marine ecosystem. However, the mechanisms to generate diverse isoforms and its implication in the fight against pathogens should be further studied.⁷⁴

There is only one mytimycin identified from *M. edulis*⁷⁵ with molecular weight of 6.2 kD. The search in the peptide sequence data bases did not yield any homology with known peptides. There are twelve cysteines engaged in the formation of six intra-molecular disulfide bridges. Its antimicrobial activity was strictly antifungal.

In addition to the AMPs found in *M. galloprovincialis*, defensins have been identified from other bivalves, including *Cg-Def*, *Cg-Defh1* and *Cg-Defh2* from *C. gigas*,⁷⁶ and a big defensin (*AiBD*) from *A. irradians*.⁷⁷ *Cg-Defh1* and *Cg-Defh2* identified from hemocyte were almost same and shared around 80% identity with mantle *Cg-Def*.⁷⁶ Recombinant *Cg-Def* was active in vitro against Gram-positive bacteria but displayed

no or limited activities against Gram-negative bacteria and fungi. *AiBD* was the first bivalve big defensin gene cloned from bay scallop *A. irradians*. It consisted of 531 nucleotides with a canonical polyadenylation signal sequence AATAAA and a poly(A) tail, encoding a polypeptide of 122 amino acids. Recombinant AiBD showed activities against both Gram-positive and Gram-negative bacteria and some fungi.

As the most important effector in bivalve immunity, more AMPs will be found and well characterized, which will provide a better understanding of the immune defense mechanisms of bivalve and new insights into health management and disease control in aquaculture.

Lysozymes

Lysozyme is a ubiquitous enzyme existing in numerous phylogenetically diverse organisms such as bacteria, bacteriophages, fungi, plants and animals, which catalyzes the hydrolysis of β -1,4-glycosidic linkage between *N*-acetylmuramic acid and *N*-acetylglucosamine of PGN and causes bacterial cell lysis.⁷⁸ It has been widely accepted that lysozyme functions as a crucial effector molecule in innate immunity. In general, the lysozymes are classified into six types according to the organisms where they are first identified, chicken-type (c-type), goose-type (g-type), invertebrate-type (i-type), phage, bacterial and plant lysozyme.

Several lysozymes and their activities have been characterized in bivalves. All previous reported bivalve lysozymes are i-type.⁷⁹ But recently, two g-type lysozymes (*CFLysG*, *AILysG*) were identified in scallops and no such thing has been reported before. It demonstrated that g-type lysozyme was not vertebrate-specific and its origin should precede the divergence of invertebrate and vertebrate.⁸⁰ Chlamysin from *Chlamys icelandic*,⁸¹ lysozymes from *Tapes japonica*⁸² and *C. virginica*⁸³ have been verified as i-type lysozymes with remarkable antibacterial activity against both Gram-positive and Gram-negative bacteria. The g-type lysozyme CFLysG possessed all conserved features critical for the fundamental structure and function of g-type lysozymes, such as three catalytic residues (Glu 82, Asp 97, Asp 108) and showed more potent inhibitive activities against Gram-positive bacteria. The self-defense activity of lysozyme have been detected widely in the body fluid and various tissues of bivalves, including the genus of *Mytilus*, *Bathymodiolus*, *Calyptogena* and *Chlamys*. A recent study has demonstrated that lysozyme could augment the activity of AMPs through a synergistic mechanism.⁸⁴ The lysozyme can also serve as a digestive enzyme in the digestive organs,⁸⁵ which has been further verified by lysozyme activities in hepatopancreas of several bivalves.

Lectins

The function of lectins includes not only self/nonsel self recognition but also engaging associated effector mechanisms, such as complement-mediated opsonization and killing of potential pathogens. Diverse lectins have been reported in oysters,^{31,35} scallops and clams,^{86,87} and they are involved in the immune response against pathogens. The recombinant C-type lectins Cflec-1, Cflec-2, Cflec-3 and Cflec-5 from *C. farreri* agglutinated *E. coli*, *S. haemolyticus*, *P. stutzeri* and *P. pastoris*, respectively. The lectin from *C. virginica* agglutinates a wide variety of bacteria.⁸⁸ The sialic acid-binding lectin of horse mussel, *Modiolus modiolus*, has strong antibacterial activity against *Vibrio* strains.⁸⁹ A calcium independent lectin isolated from the foot muscle of marine

bivalve *Macoma birmanica*, named MBA, could interact with both Gram-positive and Gram-negative bacteria.⁹⁰ There is increasing evidence to support the suggestion that the presence of isoforms and diverse roles of lectins provide bivalves with a functional diversity to the innate immune response.

Cytokines

Cytokines comprise a large number of regulatory molecules, interleukins (IL), IFN, TNF and chemokines and many of them function vitally in the vertebrate immune system.⁹¹ In view of the significant functions, the studies of cytokine in bivalves have been pursued since the early 1990s. In the first publication, oyster cells were found to respond to IL-1 and TNF in a manner similar to that of human granulocytes.⁹² Subsequently, the information about the effect of cytokines on immunity such as cell motility, chemotaxis, phagocytosis and cytotoxicity has been accumulating in bivalves.⁹³ However, the previous knowledge about bivalve cytokine is from immunological experiments with human antibodies, which lacks molecular evidence.⁹⁴ Recently, cytokine homologs and their receptors have been validated at molecular level.

The first identified cytokine is oyster IL-17 homologue (*CgIL-17*). Bacteria challenge induced a large and rapid elevation of *CgIL-17* transcript in oyster.⁹¹ This is the solely IL identified in bivalve. Although other cytokines are not available in bivalves, two TNF receptor genes were cloned from Zhikong scallop, which was homologous respectively with p75 neurotrophin receptor (*p75NTR*) and osteoprotegerin.⁹⁵ Another receptor for the transforming growth factor beta (*TGFβ*) has been identified from oyster *C. gigas*.⁹⁶ The phylogenetic and structural analysis as well as the expression pattern during early development suggested that Cg-TGFβRI belonged to the TGFβs.s./activin type I receptor. The existence of different receptors suggested the corresponding cytokines should be present in bivalves.

It is exciting that a bivalve TNF (*RpTNF*) gene was cloned from clam *R. philippinarum*. The typical structure indicated that RpTNF was a true Type II (i.e., intracellular N terminus and extracellular C terminus) transmembrane protein. The recombinant RpTNF induced the death of tumour cells just like its mammalian homologs (unpublished data). Considering the broad immune response of cytokines, a multiplex cytokine-receptor system must exist and mediate bivalve immunity in a specific “ligand-receptor” manner, which may take important roles in regulating the bivalve immunity.

Complement Components

The complement system is one of the major effector arms of immune response in vertebrates and a necessary complement for antibodies to play a role in cytolytic. Its effector functions include opsonization leading to enhanced phagocytosis and lysis of microbes. In the evolutionary progress, it appears earlier than the acquired immunity. The previous research on sea urchins, tunicates and horseshoe crabs also revealed a simple opsonic complement defense system in invertebrates.⁹⁷ However, there are few reports on its role in bivalves.

The recent identification of C3 and B factor-like (details in other section) from carpet-shell clam provides insight into their conserved characters critical for function. Three other molecules with the hallmark domain structure of complement factors

have also been cloned from scallops (details in other section). A C1qDC protein has been identified from *C. farreri* to have LPS binding activity.⁶⁹ The *CfTEP* transcripts were mainly detected in the tissues of hepatopancreas and gonad and remarkably up-regulated by microbial challenge.⁴² Its expression was complicatedly mediated by alternative splicing mechanism. Recombinant AiFREP agglutinated chicken and human A, B, O-type erythrocytes. The agglutinating activities were calcium-dependent and could be inhibited by acetyl group-containing carbohydrates. It also agglutinated Gram-negative bacteria *E. coli* JM109, *V. anguillarum* and Gram-positive bacteria *M. luteus* in the presence of calcium ions. The attractive studies collectively favored that those complement factors play significant role in bivalve immune responses and complement system should be an intriguing driver for understanding the characteristic of innate immune system in bivalves.

The Antioxidant Enzymes

ROS are free radicals that contain the oxygen atom, constantly generated when the organism is attacked by invaders or contaminant exposures.⁹⁸ Low concentrations of ROS, such as superoxide anion (O_2^-), hydrogen peroxide (H_2O_2) and hydroxyl radical ($OH\cdot$) may be beneficial or even indispensable in processes including intracellular signaling and defense against micro-organisms. Excessive production of ROS may, however, lead to oxidative stress, loss of cell function and ultimately apoptosis or necrosis.⁹⁹ As with chemical antioxidants, cells are protected against oxidative stress by an interacting network of antioxidant enzymes. In bivalves, a large number of antioxidant enzymes have been identified over the last decades and their response against microbe challenge and environmental stress have been studied. The knowledge of superoxide dismutase (SOD), catalase (CAT), glutathione peroxidase (GPx) and glutathione-S-transferase (GST) is summarized briefly in this section.

Superoxide Dismutase

SODs are a class of antioxidant enzymes that catalyze the dismutation of superoxide into oxygen and hydrogen peroxide. According to their metal content, SODs are classified into four distinct groups: iron SOD (FeSOD), manganese SOD (MnSOD), copper/zinc SOD (Cu/ZnSOD) and nickel SOD (NiSOD). However, there were only Cu/ZnSOD and MnSOD identified presently in bivalves.¹⁰⁰ Most of SODs were constitutive proteins that could play a crucial role in bivalve defense. Many studies have indicated the importance of SODs to the immune response as well as the role in protecting cells against various challenges.¹⁰⁰ For example, the SOD genes have been identified from scallop *C. farreri* and *Nodipecten subnodosus* and a rapid elevation of SOD activities were observed after microbe infection.^{101,102} The SOD activity in gills and mantle of *M. galloprovincialis* increased obviously at the toxic chemical pollutants, especially heavy metals.¹⁰³ While in *Chamelea gallina*, the SOD activity decreased in haemocytes with increasing temperature or exposure to benzo[a]pyrene.^{104,105} A higher-level of mRNA expression of MnSOD was detected in gill and mantle of *A. irradians* after being challenged with *V. anguillarum* and the expression level in gill was even higher, indicating that MnSOD was necessary in the immune responses against *V. anguillarum* infection. The responses of bivalve SOD to environmental stress were quite variable depending upon isoforms

and tissues. When exposed to heavy metals, SOD is predominantly detected in the cytosolic fractions of gill and digestive gland, with the highest amount in the gill. The mitochondrial Mn-SOD activity in the gill and digestive gland is lower than cytosolic SOD.¹⁰⁶ The whole set of results make them potential target biomarkers aiming at the given environment factors in monitoring strategy.

Catalase

CAT is one of the central enzymes involved in scavenging the high level of ROS. It catalyzes the decomposition of hydrogen peroxide to gaseous oxygen and water molecules.¹⁰⁷ This enzyme is ubiquitous and present in archaea, prokaryotes and eukaryotes. In mammals, the functional catalase is a tetramer of four identical subunits with a molecular weight of approximate 240 kD.¹⁰⁸ Recently, more and more proteins with catalase activities and the genes have been reported in bivalves, such as *M. edulis*,¹⁰⁹ *D. polymorpha*,¹¹⁰ *C. gigas*¹¹¹ and *C. farreri*.¹¹² The expression of CAT from *C. farreri* increased gradually after-*Vibrio* infection. *M. galloprovincialis* CAT activity was increased 2-3 times at the polluted coastal areas, with high activity in winter and spring.¹⁰³ This rise of *M. galloprovincialis* catalase activity was significant with temperature, salinity and light duration. The importance of these enzymes in regulating oxidative stress is recognized. CAT is considered as an important and sensitive biomarker of environmental stress, used to reveal the biological effect on the redox status of bivalve organisms.

Glutathione Peroxidase

GPx is the general name of an enzyme family with peroxidase activity whose main biological role is to protect the organism from oxidative damage. It reduces lipid hydroperoxides to their corresponding alcohols and reduces free hydrogen peroxide to water. Two isoforms of GPx had been identified, selenium-dependent GPx (Se-GPx) and selenium-independent GPx. Se-GPx catalyses the reduction of both organic and inorganic peroxides like hydrogen peroxide (H₂O₂) while selenium-independent GPx reduces only organic peroxide. The detection of GPx activity and identification of GPx molecules collectively suggest that GPx may respond to bacterial infection and hydrogen peroxide exposure and be involved in protection against oxidative stress and immune defense in bivalves.^{2,113-115} To date, the activity of GPx has been detected in bivalves *Corbicula fluminea*,¹¹³ *Pinna nobilis*,¹¹⁴ *M. galloprovincialis*¹¹⁵ and *Ruditapes decussatus*.² There have also been several reports on the GPx genes cloned from bivalves, such as *D. polymorpha*, *Unio tumidus* and *C. gigas*.

Glutathione S-Transferase

GSTs are comprised of classes of dimeric enzymatic proteins that catalyse the conjugation of glutathione to a wide variety of hydrophobic compounds through the formation of a thioether bond with their electrophilic centre. Most mammalian GSTs are cytosolic enzymes with molecular masses of 23 to 28 kD as homodimers or heterodimers. The cytosolic GSTs have been classified into at least 12 different classes based on their N-terminal amino acid sequence, substrate specificity, antibody cross-reactivity

and sensitivity to inhibitors. To date, many novel classes of GST sequences have been identified and classified from nonmammalian organisms. A few GSTs have been studied in bivalves, including *Atactodea striata*, *R. decussates*, *M. edulis*, *C. fluminea*, etc.

GSTs are major Phase II detoxication enzymes found mainly in cytosol and function as a substrate of antioxidant enzymes to eliminate the reactive oxygen induced by xenobiotic compounds,¹¹⁶ providing protection against electrophiles and products of oxidative stress. Most GST studies in bivalves focused on purification and/or biochemical measurement of total GST or different GST isoforms. GST has been used as a biomarker in environmental assess in bivalves for a number of years. The level of GST in bivalves was responsive, at some extent, to the potential environmental contamination exposure. Mussel population affected by nontreated wastewaters exhibited significantly higher GST activities as a result of an increase in conjugating activities.¹¹⁷ A rapid increase of GST activities was observed in primary cultured digestive gland acini of *Pecten maximus* treated with Tributyltin, ethylmethane sulfonate and the water-soluble fraction of crude oil, in a time and dose dependent manner.¹¹⁸

Acute Phase Proteins

In addition to the discription above, there are other effectors, such as acute phase proteins, to protect bivalve against the toxic effects of contaminants or bacteria challenge. Although nonspecific, acute phase response serves as a core of the innate immune response involving physical and molecular barriers and responses that serve to prevent infection, clear potential pathogens, initiate inflammatory processes and contribute to resolution and the healing process. Acute phase proteins, an integral part of the acute phase response, have been identified in many bivalve species. Among these, heat shock protein and metallothioneins (MT) are two families of acute phase proteins recently studied in bivalves.

Heat shock proteins (HSPs) are ubiquitous and highly conserved stress proteins, not only playing important roles in response to potentially deleterious stress conditions, but also preventing cell toxicity and cell death to protect cells and tissues against damage. More recently, it has also been suggested that HSPs could function as potent activators of the innate immune system.¹¹⁹ The principal HSPs range in molecular mass from 15 to 110 kD. According to apparent molecular mass, they are classified into several families. Members of HSP22, HSP60, HSP70 and HSP90 have been identified from invertebrate and most of them were found in bivalves. Recent studies in different species of bivalves have confirmed the relevant physiological roles of HSP expression in thermal tolerance and multiple stress response.¹²⁰ The different forms of HSPs might be highly affected by temperature and salinity, as well as by a great variety of chemical stressors that might be often found in seawater or in the sediment.^{120,121} Meanwhile, the response of some HSPs against bacteria challenge has also been reported in bivalves. After *Vibro* stimulation, a clearly time-dependent expression pattern of HSP22, HSP70 and HSP90 was observed in scallops. HSPs might serve as powerful biomarkers of marine pollution and be helpful for health management of bivalve aquaculture.

Metallothionein (MT) is a superfamily of cysteine-rich proteins with low molecular weight, no aromatic amino acids and high metal binding affinity and widely found in a large variety of organisms.¹²² In the past years, multiple functions have been attributed

to MT proteins, such as homeostasis of essential metals, detoxification of toxic metals, protecting against ionizing radiation and oxidative stress, the scavenging of free radicals and response to estrogenic compounds. Bivalves are known to accumulate high concentrations of heavy metals in their tissue and are widely used as bioindicators for pollution in marine and freshwater environments. There has been considerable research dedicated to the diversity of metal-inducibility and expression in different MT isoforms from bivalves *C. virginica*, *M. galloprovincialis* and *D. polymorpha*.^{123,124} The mRNA expression of MT from scallop was increased drastically to hundred fold post *Vibrio* challenge respectively.¹²⁵ The sensitivity of MT from scallop to bacteria challenge offers us a hint of its regulation in scallop immune defense. The progress opens research perspectives for the use of this marker to assess the effect of various pollutants in the aquatic environment and a better knowledge of its functional multiplicity in the bivalve immune system.

CONCLUSION

Bivalves are always challenged by their environment with high bacterial and viral loads, pollutants and they have evolved a multiplicity of efficient defense strategies to defend against microbial attack. In recent years considerable progress has been made in our understanding of the bivalve immunity. As described in this chapter, bivalve immune system is effective, multifaceted and incorporates cellular and humoral components. It is well accepted that the immune system in bivalves lacks the components of adaptive immunity present in the vertebrates, but there is also a complex mechanism involving an array of different molecules and various multi-step cascade processes. Many aspects of bivalve immunity are not well understood. Even some of the molecules and immune response mechanisms are found to be structurally and functionally similar to that in vertebrate animals. The research progress about the bivalve immunity has been hampered by absence of genome, tools for genetic manipulation and mutants and stable long-term cell lines for in vitro studies. Thus, there is no doubt that the collaborative efforts among immunologists, cell biologists, physiologists and geneticists are necessary for the study of bivalve immunity. It is still an essential task for us to fully characterize the molecules, responses, cascade pathways involved in bivalve immunity. The analysis of the precise underlying molecular mechanism of bivalve immunity will assist in understanding the nature and evolution of immune system and the connections between immune defense in invertebrates and vertebrates.

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CHAPTER 4

EARTHWORM IMMUNITY

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Abstract: Earthworms belonging to oligochaete annelids became a model for comparative immunologists in the early sixties with the publication of results from transplantation experiments that proved the existence of self/nonself recognition in earthworms. This initiated extensive studies on the earthworm immune mechanisms that evolved to prevent the invasion of pathogens. In the last four decades important cellular and humoral pathways were described and numerous biologically active compounds were characterized and often cloned.

INTRODUCTION

Considering the fact that the majority of the immunologists all over the world are focused on the mammalian or particularly human immunology, it may sound surprising that invertebrates—and among others earthworms—have been an important experimental model since the very beginning of immunology. For example, phagocytosis, an important and evolutionarily conserved defense mechanism of innate immunity, was discovered in the late 1800s by a Nobel Prize winner Elie Mechnikoff while studying the origin of the digestive organs in the floating larvae of starfish.¹

From the total number of extant animal species, certainly surpassing 2 millions, 95% are included in the invertebrate taxa. Invertebrates have evolved for hundreds of millions of years, often surviving in very hostile environments. Their successful survival strategies are likely based on short life span combined with numerous offspring. More importantly, all invertebrate species have developed a variety of defense mechanisms efficiently recognizing and responding to nonself substances.^{2,3}

In contrast to adaptive immunity, which is a highly sophisticated system based on antigen-specific T and B cells and antibodies and which is observed in vertebrates only,

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many innate immunity mechanisms are conserved from invertebrates to vertebrates. Cellular mechanisms of invertebrate innate immunity include wound repair, clotting and coagulation responses, phagocytosis of invading microorganisms and encapsulation reactions. Apart from these cellular mechanisms, invertebrates possess a broad range of antimicrobial factors such as lysozyme-like proteins, proteases, cytolytic proteins, antimicrobial peptides and enzyme activation-base cascades; humoral defense also includes lectin-like and pattern recognition molecules that are designed to recognize a few highly conserved structures present in many different microorganisms. The majority of the above immune responses of invertebrates are nonadaptive with no or very limited ability either to “remember” or to respond more vigorously and effectively to repeated exposures to the same pathogens.⁴

BASIC INFORMATION ON EARTHWORM ANATOMY

The earthworms are protostomian animals possessing true coelom of mesenchymal origin. The coelomic cavity is filled with coelomic fluid containing free wandering cells, named coelomocytes, originating in the mesenchymal lining of the cavity. The coelomic cavity is metameric and the segments are separated by transversal septa. Transport of the coelomic fluid and coelomocytes between the segments is regulated by channels. Each segment of the coelomic cavity is opened to the outer environment by a pair of nephridia and by a dorsal pore (Fig. 1).

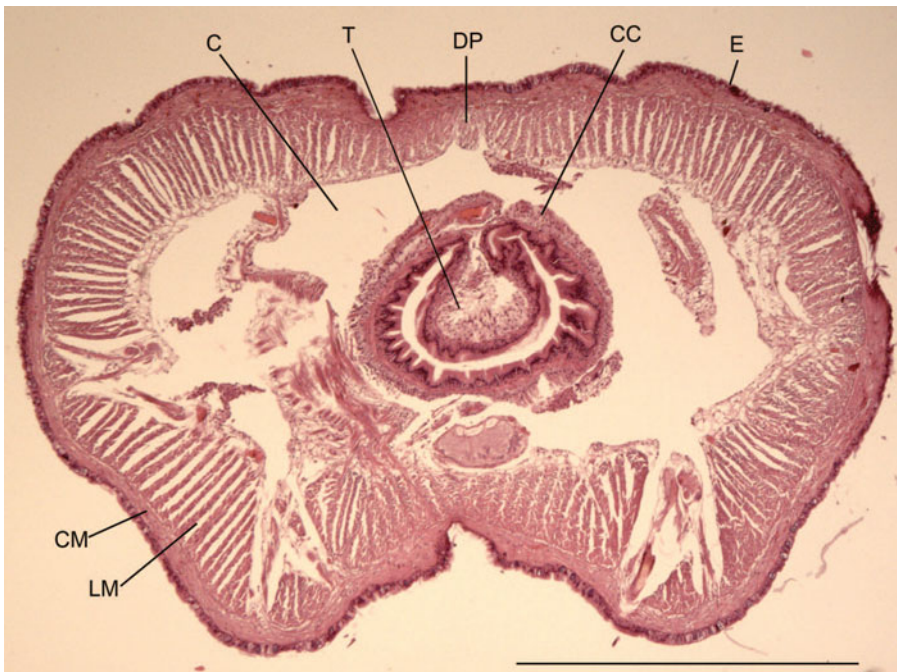


Figure 1. Transverse section of *Eisenia fetida* stained with hematoxylin-eosin. C—coelom, T—typhlosole, DP—dorsal pore, CC—chloragogen cells, E—epidermis, CM—circular muscles, LM—longitudinal muscles. Bar 1 mm. Reproduced by courtesy of Dr. Pavel Rossmann, Prague.

In terms of systematic biology, the earthworm family—Lumbricidae—is the largest member of the class Oligochaeta, phylum Annelida. The earthworms are found in leaf litter, manure, under stones and logs as well as some arid areas, but most species prefer wetter, more heavily vegetated regions. Earthworms range in size from two centimeters to over one meter. There are over 3000 described earthworm species known worldwide and they have adapted to a wide range of soil habitats as well as freshwater lakes and streams (for a review see ref. 5). Immunological research is performed mainly in two genera—*Lumbricus* and *Eisenia*.

CELLULAR DEFENSE MECHANISMS

The first nonspecific barrier of earthworms is the skin covering their entire body. The skin consists of the epidermis and a thin cuticle, which contains mucopolysaccharides acting as an antimicrobial barrier.^{6,7} The epidermis is formed by a single layer epithelium of supporting cells, basal cells and secretory cells. The basal cells play an important role in wound healing and graft rejection, often exerting phagocytic activity.^{8,9} Thus, these basal cells are sometimes considered not to be of epidermal origin, but rather homologous to coelomocytes.¹⁰⁻¹²

Each segment of the coelomic cavity communicates with outer environment by a dorsal pore; the skin therefore cannot sufficiently prevent the microorganisms from entering the coelomic cavity. Consequently, the coelomic cavity is not aseptic and always contains bacteria, protozoans and fungi from the outer environment. Nevertheless, there are efficient mechanisms that keep the growth of microorganisms under control.¹³ It was reported that coelomic fluid contains 6×10^5 /ml naturally occurring bacteria while the number of potentially phagocytic cells is more than ten times higher. These abundant phagocytes combined with the presence of various humoral factors can easily prevent the microorganisms from outgrowth.

The invading microorganisms can be eliminated by a number of ways. First, they can be excreted by nephridia¹⁴ or engulfed by the cells of nephrostome or middle tube.¹⁵ Second, as mentioned above, the microorganisms can be phagocytosed by certain coelomocytes and phagocytic cells that, when they become exhausted, are expelled through dorsal pores. The pores are equipped with muscular sphincters controlling intracoelomic pressure and the exchange of material between the outer and inner environments.¹⁴ Third, large foreign bodies, e.g., agglutinated bacteria or parasites are eliminated by encapsulation.^{16,17} This process begins, similarly to phagocytosis, by the recognition of foreign material which, however, cannot be engulfed due to its size. Within the first day, the foreign body is surrounded by free coelomocytes and after several days a dense capsule (often called a brown body because of its melanin content as a consequence of the prophenoloxidase cascade) composed of flattened cells is formed. When the capsule is about 1-2 mm in diameter, its external cells lose their adhesiveness so that the capsule can migrate towards the posterior segments of the coelomic cavity where it is eliminated by autonomy followed by wound.¹⁸⁻²¹ It was documented in *Eisenia fetida* earthworms that most brown bodies contain tissue wastes, agglutinated bacteria, grearines or nematodes.¹⁷

As mentioned above, coelomic fluid contains different types of coelomocytes. Their nomenclature is based mainly on morphological and cytochemical criteria (for a review see refs. 22,23) though more recent studies attempt to determine superficial and

functional markers for cell classification.²⁴ In general, there are three main coelomocyte types—eleocytes, free chloragogen cells with nutritive and accessory functions and either hyaline or granular amoebocytes, both representing effector immunocytes involved in a broad range of defense functions including phagocytosis.

Although both types of amoebocytes have phagocytic properties, their activity differs. In contrast to granular amoebocytes, the cytoplasm of hyaline amoebocytes is occasionally full of engulfed material.²² It should be mentioned that amoebocytes engulf all kinds of material including inert particles, microbial cell wall components as well as foreign cells. However, the phagocytosis of eukaryotic cells depends on the source of the cells. Unlike allogeneic cells, xenogeneic cells (both from different earthworm species and from noninvertebrate species) are rapidly phagocytosed.^{14,25}

Phagocytosis by coelomocytes, similarly to that of vertebrates, can be modulated by humoral components, opsonins, which coat the engulfed particle and thus promote its phagocytosis. It was proven that preincubation of both yeast and synthetic copolymer particles with the coelomic fluid significantly increased their phagocytosis.^{26,27} It is noteworthy that also mammalian opsonins, IgG immunoglobulin and C3b complement fragment, were described to enhance coelomocyte phagocytic activity, in contrast to IgM and C3d fragment, which did not affect phagocytosis.²⁸

The earthworms are regarded as an important model organism of comparative immunology since 1960s when transplantation experiments were performed (for a review see ref. 29) and cell-mediated short-term memory was observed.³⁰ All these experiments proved the existence of self and nonself recognition in earthworms and initiated extensive studies of earthworm immune mechanisms. The ability to recognize and respond to allografts as well as xenografts and, on the other hand, the ability to accept or not to destroy autografts was observed in many annelid species.²⁹ This process begins like the reaction to injury. The first major change, which occurs after the healing of wounds is, regardless of the graft origin, the accumulation of coelomocytes near the graft sites and their infiltration into the matrix. The response to the xenografts results in complete walling off of the graft and its destruction by encapsulation reaction.³¹ The number of invading coelomocytes during the autograft transplantation is markedly lower^{32,33} but the reaction seems to be more rapid. The maximum number of coelomocytes surrounding the graft was detected within 24 hours, returning to the normal level by 72 hours. In contrast, the peak response to xenografts is on day 3 or 4 and normal levels are not reached before day 7. The destruction of xenografts is completed approximately by day 17 after the transplantation. If a second graft is transplanted at this time, an accelerated rejection within 6 or 7 days occurs. Moreover, the number of the invading coelomocytes is 20-30 % higher. The increased number of coelomocytes during the retransplantation is probably caused by an increased proliferating activity of mesenchymal lining of the coelomic cavity and the septa. These data suggest the existence of short-term and very limited memory, which is based solely on cells as the transfer of either the coelomic fluid or other substances does not induce any accelerated reaction.^{30,33}

Allo- and xenorecognition mediated by earthworm coelomocytes was evidenced in experiments showing cell-mediated cytotoxicity to allogeneic coelomocytes³⁴ and, more interestingly, capability of earthworm non/phagocytic coelomocytes to kill efficiently tumor cell targets such as K562.^{35,36}

HUMORAL DEFENSE MECHANISMS

The coelomic fluid of annelids exerts numerous biological activities that are involved in effective defense mechanisms against invaders. It was documented that it contains various antimicrobial factors like lysozyme^{37,38} and antimicrobial peptides.³⁹⁻⁴¹ Moreover, coelomic fluid was found to cause the lysis of vertebrate erythrocytes and, subsequently, several hemolytic factors were isolated and described. The majority of proteins with hemolytic properties have hemagglutination activity as well and, more interestingly, a spectrum of antibacterial and bacteriostatic activities against pathogenic soil bacteria.⁴²⁻⁴⁴ Furthermore, it was observed that coelomic fluid lyses eukaryotic cells other than erythrocytes, namely fibroblasts and insect hemocytes⁴⁵ and various tumor cell lines.^{46,47} Hereinafter, each activity of the coelomic fluid is thoroughly described.

Lysozyme is a bacteriolytic enzyme which catalyzes the hydrolysis of 1,4- β -D-links between N-acetylmuramic acid and N-acetyl-D-glucosamine residues in the peptidoglycan of bacterial cell walls and thus efficiently protects against infections caused particularly by Gram-positive bacteria. Lysozyme activity was observed in coelomocyte extracts as well as in the coelomic fluid.³⁷ Later, the active protein was isolated and partially sequenced.⁴⁸ Based on the N-terminal sequence, a novel class of lysozymes including those of molluscs, echinoderms, nematodes and earthworms was proposed.⁴⁸ Recently, cDNA coding for lysozyme-like molecule of *E. andrei* earthworms was characterized and cloned.³⁸ Earthworm lysozyme exhibited both lysozyme and isopeptidase activity and shared homology with other invertebrate lysozymes, with the highest similarity (72 % identity) to destabilase I from medicinal leech. Moreover, lysozyme expression can be up-regulated after a challenge with Gram-positive as well as Gram-negative bacteria.

Antimicrobial peptides are an abundant and diverse group of molecules that are produced by many cell types in invertebrates, vertebrates and plants. To date, only a limited number of bioactive peptides have been described in annelids. An antimicrobial peptide named Lumbricin I was identified in *Lumbricus rubellus*.³⁹ Lumbricin I is a proline-rich antimicrobial peptide which is constitutively expressed in adult animals and is not induced by bacterial infection. A Lumbricin I analog named PP-1 was found in the Asian earthworm *Pheretima tschiliensis*, is synthesized in the body wall only and its localization in the mucus of the epidermis suggests its role in the mucosal defense.⁴⁰ Furthermore, an antimicrobial short peptide OEP3121 of only 5 amino acids was found in *Eisenia fetida* earthworms.⁴¹

The coelomic fluid of *Eisenia* earthworms was described to exhibit strong *hemolytic activity* that is tightly connected with *bacteriostatic and antibacterial properties* against pathogenic soil bacteria. The first hemolytic proteins were described by Du Pasquier and Duprat⁴⁹ and later on they were named *EFAF* (*Eisenia fetida andrei* factors) and characterized as two glycoproteins secreted by chloragocytes and eleocytes.^{42,50,51} The 45-kDa protein is encoded by a single nonpolymorphic gene and has a pI of 6.0, while the 40-kDa protein is encoded by a gene having four alleles, each representing one of isoforms with pI of 6.3, 6.2, 5.95 and 5.9. Each individual earthworm possesses the 45-kDa protein and 1 or 2 isoforms of the 40-kDa protein.^{42,52} In addition to EFAFs hemolytic activity, these proteins were found to agglutinate red blood cells⁵³ and to participate in the cytotoxic activity of the coelomic fluid.⁴⁵ Moreover, they exhibit antibacterial activity against both Gram-positive and Gram-negative bacteria,^{43,54,55} particularly against strains

that are pathogenic for earthworms.^{44,52,56} In addition to their bacteriolytic activity, they may also mediate opsonization⁵⁷ and participate in the clotting of the coelomic fluid.⁵⁸

It was documented, that upon binding to sphingomyelin, a major lipid constituent of plasma membranes of most mammalian cells, these proteins polymerize and form 10-nm channels through the lipid bilayer.^{50,59}

Later, EFAs were characterized at the molecular level and were named **fetidins**. A gene for the 40-kDa protein was cloned^{60,61} and it was found that its putative amino acid sequence comprises an N-glycosylation site and a peroxidase motif. This is in accordance with the finding that both fetidins have peroxidase activity.

Independently, a 41-kDa hemolytic protein, which is produced by coelomocytes and causes contraction of rat vascular smooth muscles, was characterized and named **lysenin**.⁶² Simultaneously, two 42-kDa lysenin-related proteins with weak contractive activity were identified.⁶³ More recently, a new member of this lysenin-like multi-gene family has been cloned and provisionally called lysenin-related protein 3.⁶⁴ Lysenin has a high amino acid sequence homology with fetidin (89% identity, 95% positivity), with lysenin-related protein 1 (76% identity, 89% positivity) and lysenin-related protein 3 (81% identity, 90% positivity). Amino acid sequence of lysenin-related protein 2 corresponds to that of fetidin. All these data suggest a close relationship between these lytic molecules. Individual sequence analyses have revealed that fetidin and lysenin are encoded by two distinct highly homologous genes but their expression level differs in individual earthworms.⁶⁵

The hemolytic activity of lysenin is dependent on the presence of sphingolipids in the membrane.⁶⁶ Moreover, the presence of cholesterol in the membrane facilitates hemolysis. Upon binding to the sphingomyelin, lysenin forms oligomers and subsequently pores 3 nm in diameter in the target membranes.^{66,67} Oligomerization does not occur on bacterial membranes since they are devoid of sphingomyelin. Therefore the mechanism of antibacterial activity must be different from its cytolytic activity.⁶⁴

As sphingomyelin is crucial for the cytolytic activity of lysenin, it has been proposed to use lysenin as a valuable probe for sphingomyelin detection in sphingomyelin storage diseases, particularly in the cells of Niemann-Pick A patients,⁶⁶ although the multiplicity of hemolysins in the natural source and the cytolytic activity appeared to be a major obstacle.

Independently on fetidin and lysenin, eiseniapore and hemolysins H₁, H₂, H₃, CL₃₉ and CL₄₁ were described.⁶⁸⁻⁷¹ Nevertheless, their more detailed analyses revealed close relationships with fetidin, lysenin or lysenin-related proteins.

Coelomic Cytolytic Factor as a Pattern Recognition Molecule

The coelomic fluid does not cause either the lysis of the coelomocytes of other earthworm species or of the hemocytes of mollusks, nematodes and protozoans. However, it was documented that coelomic fluid of *E. fetida* lyses a broad spectrum of various cell types including chicken fibroblasts, guinea-pig polymorphonuclear leukocytes and insect hemocytes.⁴⁵ A proteinase-independent cytolytic effect of the coelomic fluid was observed in experiments with TNF-sensitive tumor L929 cell line. Subsequent isolation of lytic proteins led to the identification of a 42-kDa protein, which was named coelomic cytolytic factor—CCF.⁴⁶

CCF acts in earthworm defense as a **pattern-recognition molecule**. Upon binding microbial pathogen-associated molecular patterns, namely O-antigen of LPS

of Gram-negative bacteria, muramyl dipeptide and muramic acid of peptidoglycan from the cell walls of Gram-positive bacteria and β -1,3-glucans and *N,N'*-diacetylchitobiose of yeast, CCF triggers the activation of the **prophenoloxidase cascade**, which results in the formation of cytotoxic and antimicrobial compounds and thus represents an important invertebrate defense mechanism.⁷²⁻⁷⁵ The broad specificity of CCF for pathogen-associated molecular patterns results from the presence of two spatially distinct pattern recognition lectin-like domains. One domain, which shows homology with the polysaccharide and glucanase motifs of β -1,3-glucanases and invertebrate defense molecules, is located in the central part of CCF molecule and interacts with LPS and β -1,3-glucans. The C-terminal tryptophan-rich domain mediates interactions of CCF with *N,N'*-diacetylchitobiose, muramyl dipeptide and muramic acid (Fig. 2).⁷⁴

The binding activity of C-terminal domain of CCF is rather unique for *Eisenia fetida* earthworms. Comparative analysis of CCF-like pattern-recognition proteins in seven other lumbricid species (*Aporrectodea caliginosa*, *A. icterica*, *A. longa*, *A. rosea*, *Dendrobaena veneta*, *Lumbricus rubellus* and *L. terrestris*) revealed high homology in polysaccharide-binding and glucanase motifs while C-terminal part was more heterogeneous.⁷⁶ This is in a good agreement with the absence of cytolytic activity and binding capacity for *N,N'*-diacetylchitobiose and peptidoglycan components. *E. fetida* is an epigeic earthworm living in decaying organic matter, in compost and mold, where

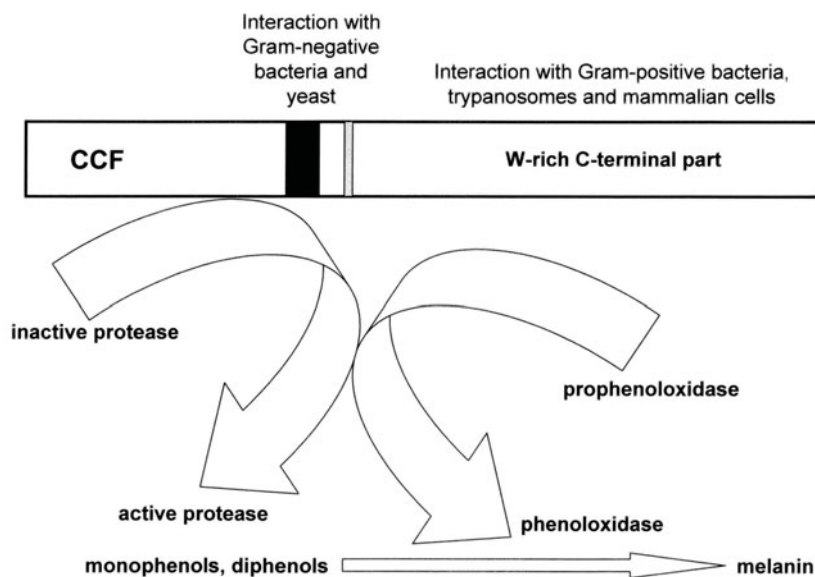


Figure 2. The role of CCF as a pattern-recognition molecule. CCF encompasses two distinct domains: the central part with polysaccharide recognition (black box) and glucanase (grey box) motifs interacts with LPS of Gram-negative bacteria and β -1.3-glucan of yeast while C-terminal tryptophan-rich domain interacts with peptidoglycan constituents of Gram-positive bacteria and N-acetylglucosamine-linked polysaccharides on trypanosomes or mammalian cells. Upon binding of pathogen-associated molecular patterns CCF activates proteolytic enzymes that cleave inactive prophenoloxidase to active phenoloxidase. Phenoloxidase then catalyzes hydroxylation and oxidation of monophenols and diphenols that leads to melanin production. Melanin exhibits antimicrobial and cytotoxic activities and potentiates other defense functions (phagocytosis, encapsulation).

the diversity as well as quantity of microorganisms is substantially higher as compared to other soil layers. It is therefore obvious that *E. fetida* appears to be best equipped to resist microbial load as reflected by the broader CCF pattern-recognition repertoire.

As indicated above, CCF displays amino acid sequence homology with bacterial and animal β -1,3-glucanases but it does not exhibit their enzymatic activity.⁷⁷⁻⁷⁹ Moreover, CCF shows homology with the α subunit of the β -1,3-glucan sensitive factor G from the horseshoe crab *Tachypleus tridentatus*,⁸⁰ with the Gram-negative bacteria-binding proteins of various insects⁸¹⁻⁸⁴ and β -1,3-glucan recognition protein of arthropods.^{85,86} All these invertebrate homologs have been suggested to play a role in invertebrate innate immunity by acting as pattern recognition molecules.

Further, it was shown that CCF agglutinates both Gram-positive and Gram-negative bacteria⁷² and contributes to the opsonizing properties of the coelomic fluid, thereby providing an efficient mechanism for phagocytosis in earthworm defense reactions.⁴⁶ CCF is also involved in the cell-mediated cytotoxic reactions and potentiates the lytic activity of coelomic fluid against red blood cells from various species.³⁴

More interestingly, CCF shares functional analogies with mammalian tumor necrosis factor. The ability of the coelomic fluid to lyse TNF-sensitive tumor cell line L929 is caused by CCF. This activity is not inhibited by anti-TNF neutralizing monoclonal antibodies, suggesting that the structure of TNF and CCF as well as the mechanism of TNF and CCF mediated lysis differ. In addition to this TNF-like lytic activity, CCF exhibits other similarities with this cytokine. CCF is secreted by phagocytic coelomocytes upon LPS stimulation (Fig. 3), while TNF is produced by LPS-activated macrophages.^{34,87} TNF and CCF have opsonizing properties^{46,88} bind β -1,3-glucans and *N, N'*-diacetylchitobiose via lectin-like interactions.^{72,89} In addition, monoclonal antibodies elicited against the lectin-like TIP domain of TNF cross-react with CCF and, conversely, monoclonal antibody against CCF reacts with TNF without impairing the interaction of TNF with its specific receptor.^{90,91} However, the activity of CCF is not inhibited by anti-TNF antibody suggesting different mechanisms of TNF- and CCF-mediated lysis.

The lectin-like domain of TNF was shown to be involved in the killing of African and American trypanosomes.⁹⁰⁻⁹² Hence, in view of the similar lectin-like activity of CCF and TNF, the possible trypanolytic activity of CCF was investigated.⁷³ The coelomic fluid of *E. fetida* as well as purified CCF were described to have a potent trypanolytic activity that can be inhibited not only by anti-CCF monoclonal antibodies but also by *N, N'*-diacetylchitobiose and anti-TNF antibodies. The possible target for both CCF and TNF on the trypanosome surface is the N-linked *N, N'*-diacetylchitobiose core of the variant-specific glycoprotein (VSG) that acts as a protective coat. This idea is strongly supported by the fact that CCF and TNF are able to lyse only bloodstream forms of parasites expressing VSG but not insect-stage procyclic forms expressing procyclin as a surface protein.

It was documented that TNF increases the membrane conductance in mammalian cells, interacting with ion-channels or ion-channel-coupled molecules through a lectin-like domain.^{93,94} Similarly, when endothelial cells or macrophages were activated with CCF, an increase in membrane conductance occurred.⁹⁵ As observed with TNF, the ion-gating effect of CCF appeared when cells from TNF-receptor I and TNF-receptor II knockout mice were used. Moreover, this effect is blocked by *N, N'*-diacetylchitobiose and amiloride—an epithelial sodium channel inhibitor—suggesting that the effect is mediated by the lectin-like domain of CCF. In macrophages, CCF-induced depolarization results in the release of TNF, IL-6 and nitric oxide via NF- κ B signaling. This pathway based

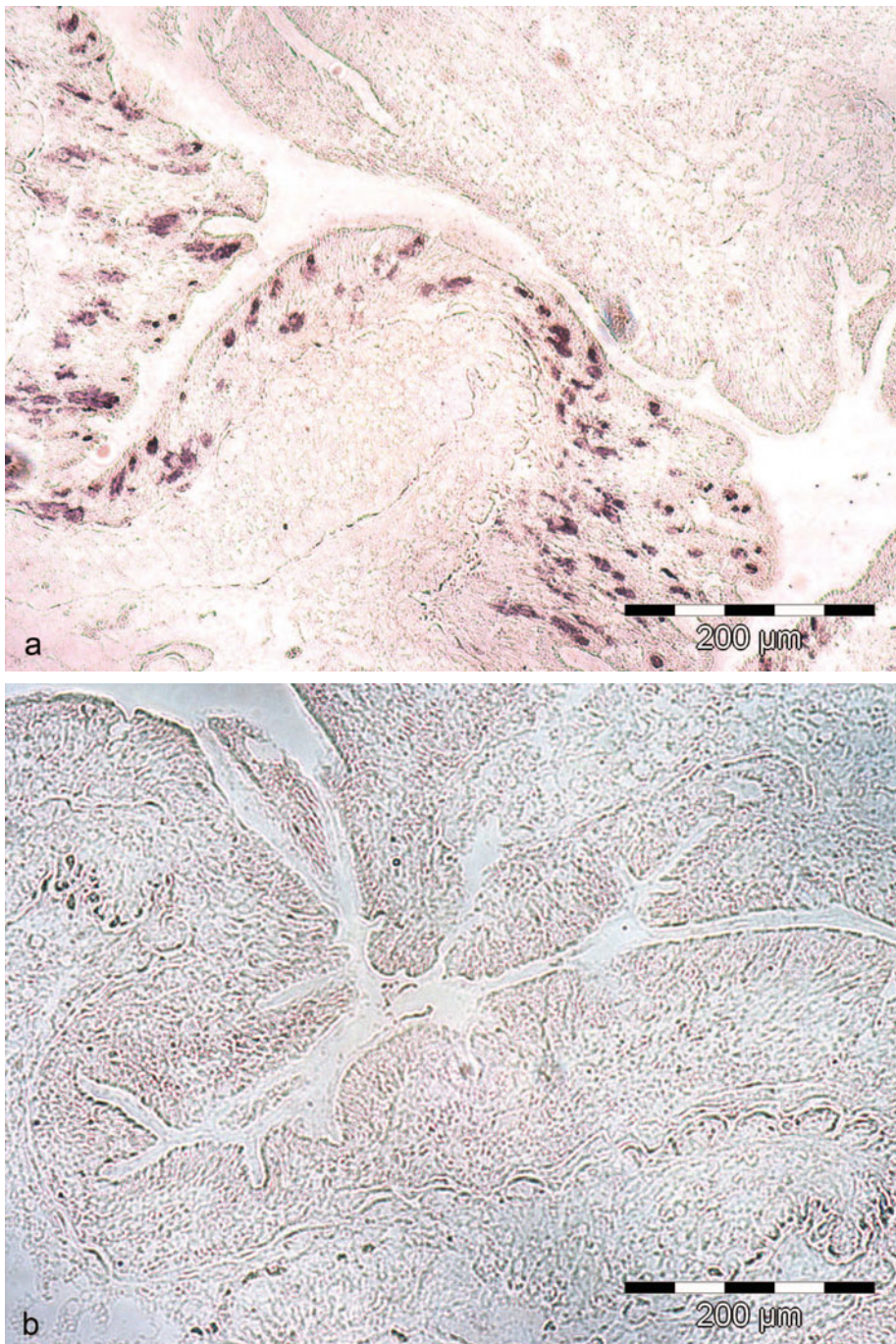


Figure 3. Up-regulated tissue expression of CCF in chloragogen tissue after LPS challenge (a) as compared to nonstimulated controls (b) followed by in situ hybridization. Reproduced by courtesy of Dr. Ellen Kauschke, Giessen.

on an interaction of lectin domain with saccharide moiety of ion channel may represent an evolutionary ancient mechanism of cell activation.⁹⁶

Surprisingly, despite the functional analogies of CCF and TNF and cross-reactivity of anti-CCF and anti-TNF antibodies, these molecules do not show any gene or amino acid sequence homology, indicating a lack of common evolutionary origin.⁷³

CONCLUSION

Earthworms rely on innate defense mechanisms that are sufficient for survival in often hostile environment. The choice of earthworms for comparative immunology studies was pertinent since they represent an inexpensive, appropriate and noncontroversial model for experimentation. Described defense mechanisms and molecules help to better understand more sophisticated immunity in vertebrates. Moreover, earthworms were found useful in monitoring environmental pollution. The Organization for Economic Cooperation and Development (OECD Guidelines for testing of Chemicals 1984) and the American Environmental Protection Agency accepted official protocols involving screening of earthworm immunological parameters as markers/indicators of impaired environmental conditions.^{97,98} Furthermore, earthworms might be considered as a source of biologically active compounds with potential industrial or medical use. Actually, earthworm powder has been used as a traditional medicine in some South Asia countries for years to treat various diseases. Currently, the therapeutic effect of earthworm active factors is being evaluated by a modern scientific approach. Some therapeutics containing fibrinolytic enzymes from *Lumbricus rubellus* and *Eisenia fetida* earthworms⁹⁹⁻¹⁰² are already commercially available to support coagulation and fibrinolysis balance in the body and thus prevent or treat cardiac and cerebrovascular diseases (Boluoke[®] (lumbrokinase), Canada RNA Biochemical Inc.).

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CHAPTER 5

LEECH IMMUNITY: From Brain to Peripheral Responses

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Abstract: In the present chapter, we will emphasize the immune response in two compartments (Central nervous system and peripheral system) in two blood sucking leeches i.e., the medicinal leech and the bird leech *Theromyzon tessulatum*. In the medicinal leech, the neuroimmune response has been described in the context of septic trauma at the cellular and humoral levels through microglia, Toll-like, cannabinoids and chemoattractant factors activation and modulation. In the bird leech, the antimicrobial responses have been dissected at the cellular and molecular levels. Altogether, this chapter presents a complete integrate immune response from the brain and the systemic compartments with high similarity to the vertebrates one. These points that the neuroimmune and immune responses evolved sooner than can be expected.

INTRODUCTION

Leeches are derived from their cousin's earthworm in the class of annelids called hirudinea. Leeches evolved during the Cambrian Explosion, a time of rapid biological development 540 million years ago.¹⁻⁶ There are now approximately 700 species of leeches distributed throughout freshwater, marine and terrestrial ecosystems worldwide. Phylogenetic studies assess that the common leech ancestor was probably a bloodsucking leech with a proboscis rather than an unspecialized ectocommensal.⁷ During the course of leech evolution, a reduction of the proboscis could have taken place in predatory arhynchobdellid ancestors to enable swallowing of larger prey. A second gain of sanguivory by the jawed Hirudiniforms could have been facilitated

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by pre-adaptations to ectoparasitic blood feeding.⁷ Recently, leech EST data from the jaw leech *hirudo medicinalis* have been obtained⁸ and it can be seen high sequence homologies between the medicinal leech and mammals⁸ suggesting a co-evolution between the parasite and the host. Thus, taking into account such evolutive data, we will present in this manuscript, data obtained from two haematophageous leeches, the jawed leech *Hirudo medicinalis* and the gut leech *Theromyzon tessulatum* focused on the immune responses angle. Due to their anatomic differences, *Hirudo medicinalis* appears as a good model for studying the immune response of the nervous system although the easy access to the body fluids of *Theromyzon tessulatum* makes this animal interesting for understanding the systemic response. Taken together, we will try to show that immune response from these blood sucking animals evolve closely to their vertebrate host, pointing the fact that their use currently in hospital never trigger patient immune responses. Host immune tolerance is due to co-evolution and molecular mimicry.⁹

THE MEDICINAL LEECH AS A MODEL FOR STUDYING THE IMMUNE RESPONSE OF THE CNS

The central nervous system (CNS) of the leech has a fixed number of bilateral neuromeres, 32. The 4 anterior-most neuromeres fuse to form the sub-esophageal ganglion and the 7 posterior-most fuse to form the tail ganglion; single bilateral neuromeres comprise the individual ganglia found in each of the corresponding body segments. A supra-esophageal ring of nonsegmental origin, together with the sub-esophageal ganglion, comprises the head ganglion. The central ganglia are connected to each other by a bilateral pair of nerves (the lateral “connectives”) and a single small medial nerve (Favre’s) and to the periphery by two bilateral pairs of nerves (the “roots”) that branch in a stereotypic pattern that allows the identification of branches up to fourth-order and even to fifth-order in some cases (Fig. 1). In hirudinid leeches, each segmental ganglionic primordium gives rise to about 400 neurons.¹⁰ Most of these are bilateral pairs (~180-190 pairs), but perhaps 5-8% are unpaired, with at least some becoming unpaired through cell death.^{10,11} Thus, understanding how a leech segmental ganglion functions requires, in principle, detailed knowledge of the function and connectivity of only ~200-220 individual neurons. Moreover, since each segmental ganglion is a variation on a theme (with the exception of the “sex” ganglia of body segments 5 and 6, which have additional complements of neurosecretory cells), the leech has one of the most accessible nervous systems from a systems analysis point of view.

An important property of leeches is their capacity to regenerate neurites and synaptic connections in the adult CNS. Neurites that have been damaged or severed can sprout, establish de novo growth cones and extend and reconnect specifically with normal targets.¹² Early stages of leech CNS regeneration following a mechanical lesion are characterized by two events that appear to be crucial for successful repair: one is the increased activity of epithelial nitric oxide synthase (NOS) in the area of the lesion and the generation and diffusion of nitric oxide (NO) and the second is the induced migration of microglia towards and their accumulation at, the injury site.^{13,14} Microglial cells are considered as the brain immune cells.

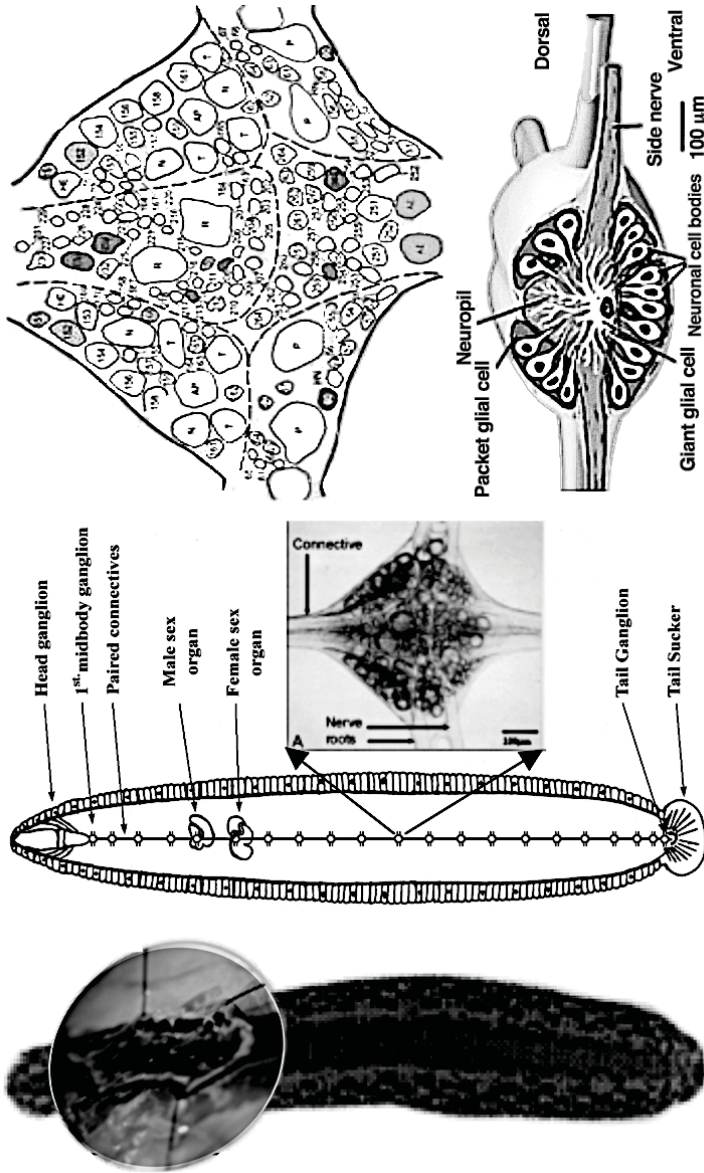


Figure 1. The central nervous system (CNS) of the leech has a fixed number of 32 bilateral neuromeres. The four anterior-most neuromeres fuse to form the sub-esophageal ganglion and the seven posterior-most fuse to form the tail ganglion; single bilateral neuromeres comprise individual ganglia found in each of the corresponding mid-body segments. The central ganglia (insert, inset photograph) are connected to each other by a bilateral pair of nerves (lateral “connectives”) and a single small medial nerve (Faivre’s) and to the periphery by two bilateral pairs of nerves (“roots”) that branch in the body wall and carry motor activity centrifugally and sensory signals centrally. **B** (Top) Diagram of a leech ganglion with some well-known neurons labeled. One of our goals is to create protein profiles for all of these neurons in order to understand the molecular mechanisms that give rise to their different functions. (Bottom) Drawing showing the typical organization of a leech ganglion: a ring of cell bodies and a central neuropil containing processes and synapses.

NO and Cannabinoids in Leech Microglia Chemotaxis Involvement

To assay directly for a role of NO on microglial accumulation at the injury site, Chen et al¹⁵ modulated NO levels in several ways. As demonstrated by NOS immunoreactivity, a large increase in NOS occurs at the crush site within 5 min of injury and this high level persists for at least 24 hrs. Microglial accumulation at the lesion, however, is not detectable at 5 min but is quite strong after a few hours and peaks at ~24 hrs. Inhibition of NO synthesis by the prior application of the NOS inhibitor L-NAME effectively blocks microglial accumulation, while the presence of its inactive enantiomer D-NAME has little or no effect. Interestingly, increasing NO levels with the NO donor spermine NOate (SPNO) also inhibits accumulation of microglia at the crush, but not in the presence of the NO scavenger cPTIO (2-(4-Carboxyphenyl)-4,4,5,5-tetramethylimidazoline-1-oxyl-3-oxide). Examination of microglial kinetics in living nerve cords shows that the effect of SPNO application occurs by the reduction of average microglial migratory speeds, even to no movement. Thus, NO is clearly implicated as a modulator of microglial movement and indeed appears to function as a stop signal at high levels, leading to the higher density of these cells at the injury site. Moreover, we recently found that the NO time-course profile in the injured leech brain is partially under the control of endocannabinoids, namely, anandamide (AEA) and 2-arachidonylglycerol (2AG), which affect the NO time-course through their modulation of cannabinoid-like receptors.¹⁶ AEA blocked microglial cell accumulation before their arrival to the lesion site in a concentration dependent-manner and this effect of AEA on microglial cell recruitment was also demonstrated in vitro. Moreover, stimulation with a concentration of AEA determined elsewhere by (Matrix assisted laser desorption ionization time of flight/time of flight (MALDI TOF/TOF) mass spectrometry¹⁷ enhanced the release of NO at the lesion site of the harmed connective. This NO release may be related to the AEA-activated CB1-like receptor carried by neurons of the leech present in the injury site. This same report suggested that this pathway leads to a microglia stop signal. In a complementary mode of action, the second most described endocannabinoid 2AG has been shown to enhance and drive the chemotaxis of microglial cells in a dose-dependent manner. Moreover, a better result in chemotaxis was obtained using a concentration of 2AG detected 30 min after injury of the leech nerve cords.¹⁷ This result is reinforced by Cabral et al who suggested that 2AG could act through the autocrine/paracrine system to chemoattract microglial cells after brain insult.¹⁸ In leech brain, AEA level declines¹⁷ and in contrast the 2AG climbs during the first 4 h following a lesion of the leech CNS.¹⁶

Knowing that these two endocannabinoids are able to act as chemoattractant factors on leech microglia in a dose-dependent manner in combination with NO release, the role of their receptors in the control of NO release and chemotaxis towards the lesion site has been investigated.¹⁶ Blocking the CB1-like receptor in the leech with a specific antagonist of the mammalian CB1 receptor (AM-251) failed to block the accumulation of microglial cells at a distance from the lesion site. This latter result underscores the role of the CB1-like receptor in microglial chemotaxis. On the other hand, blocking the CB2-like receptor in the leech with the specific antagonist AM-630 completely abolished the recruitment of microglial cells at the lesion site and costimulation of a crush with 2AG and AM-630 failed to reverse the accumulation of microglia as compared to the 2AG treatment alone.¹⁶ This latter result suggests that the CB2-like receptor in the leech is triggered by 2AG in order to promote chemotaxis and the direct recruitment of microglial cells to the site of injury. In parallel, the treatment of injured connectives with a physiological concentration of 2AG released upon CNS injury in the leech provoked a

specific response in NO production at the lesion site just after the stimulation.¹⁶ Blockade of the CB2-like receptor by AM-630 revealed an inhibition of NO secretion and the time-course of NO production was significantly delayed in comparison with the control condition. In this way, the whole of these data demonstrated that at least part of the NO produced by the injured leech was related to the 2AG-activated CB2-like receptor, as was described in experiments using Guinea pig mast cells.¹⁹

Even if 2AG and AEA are both able to produce NO during a lesion of the leech brain, only 2AG seems to play a major role in the chemotaxis of microglia and its capacity to do so is closely related to a putative functional CB2-like receptor expressed by activated microglia. Previous pharmacological studies have named the CB2 receptor as playing a crucial role in the early inflammatory process, thereby implicating microglia in this process as well in mammals. The CB2 receptor is expressed very early in the different activation steps of microglia, thus describing a “window” of functional relevance for the expression of the CB2 receptor in microglial cells.²⁰ The delayed time of the initiation of microglial cell activation is linked to changes of their morphology from resting to responsive and allows them to acquire CB2 receptors in correlation with chemotaxis and phagocytosis. The next steps of activation, named “primed” and “responsive,” were discarded from the initial step of CB2 receptor expression. As a diffusible molecule, NO has been demonstrated to take part in the migration of microglia and their accumulation at lesions of the leech CNS in a dose-dependent manner. The mechanisms by which the resting microglia becomes responsive are not yet fully understood. However, we hypothesize that the NO immediately released by the damaged neurons might stimulate the resting microglial cells present at the lesion site and allow them to be responsive for CB2-like receptor expression.

Thus, the cannabinoid system becomes activated at the lesion site and produces the two major endocannabinoids (AEA and 2AG) from membrane precursors in an opposite concentration time-course of,¹⁷ consistent with our previous work where the 2AG concentration was found to progressively increase in opposition with the AEA concentration after the lesion. It has been hypothesized that when the diffusible lipid 2AG increases, the microglia become responsive and start to accumulate at the lesion site via changes in cell morphology related to the activation of the CB2-like receptor expressed in the lamellipodia of responsive microglia.²¹ This hypothesis is reinforced by our *ex vivo* results regarding the stimulation of crushed connectives with 2AG (30 μ M) 1 h after the lesion.¹⁶ In comparison with the connectives simultaneously crushed and treated with 30 μ M 2AG, the same stimulation 1h after the crush is stimulated NO production more rapidly. This observation suggests that microglial cells are more responsive to 2AG 1h after the lesion than at the instant of the lesion, at which time microglia might be resting. During the responsive step, the leech microglia might express functional CB2 receptors at the cell surface and when these receptors are activated by the endocannabinoid 2AG, NO might be produced and participate in cell recruitment as a chemical gradient from the lesion site towards the periphery. On the other hand, the AEA released at the lesion site decreased in concentration after the lesion and it can activate NO release by targeting the CB1-like receptor and inhibiting the accumulation of microglia at the lesion site. This pathway has been described as leading to a microglia stop signal.²² The dual activity of these two endocannabinoids working in apparent opposition might be a means of controlling microglial cell recruitment to the lesion site. However, both molecules control the long-term release of NO, as shown in our time-course measurements of NO after stimulation of injured connectives with cannabinoids.¹⁶ This can be explained by the immunosuppressive response of microglial cells described both in vertebrate²³ and invertebrate models.²²

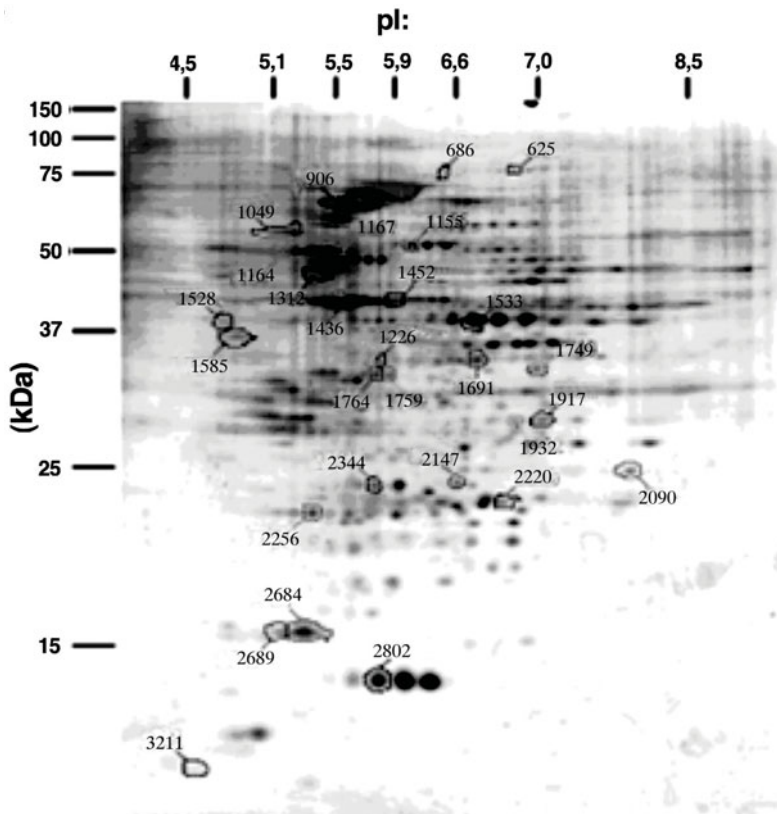


Figure 2. 2D gel between control and immune challenged leech nervous systems. Differentially represented spots are highlighted by arrows and depends on the variation times (variations at 1 h, 6 h, 12 h and 24 h) with permission from Vergote D et al. *Proteomics* 2006;6(17): 4817-25.²⁶

Bacterial Infection and Leech Brain Regeneration

Differential display proteomic analyses using 2D gel electrophoresis (Fig. 2), coupled to mass spectrometry, yielded evidence that the leech CNS responds to bacterial infection by modulating the expression of at least sixteen proteins. These proteins appear between 1 and 24h after bacterial challenge and have been assigned to the immune response because they are not induced by exposure to control sterile medium. These immune-response induced proteins include cytoskeletal and metabolic proteins, foldases, calcium sensors, kinases and neurohemerythrin, reflecting specific cytoskeletal rearrangements linked to cell migration, vesicular trafficking and/or phagocytosis, as well as the modulation of synaptic activity. Interestingly, several of these up-regulated proteins, such as gliarin²⁴ and neurohemerythrin²⁵ are expressed specifically in glial and microglial cells, suggesting a key role for these cells in the immune response of the leech nervous system, similar to what has been observed in vertebrates.²⁶ Gliarin up-regulation, in particular, could thus serve as a new marker of proliferation and maturation of leech glial cells, its up-regulation reflecting glial activation in response to the immune challenge. The accompanying

cytoskeletal rearrangements might result from morphological changes associated with phagocytosis or with the migration of cells. Indeed, it has been shown that leech microglial cells are able to migrate within the ganglionic chain to the site of a lesion and that they can play a phagocytic role.²⁷ By contrast, the expected properties of neurohemerythrin have led to the proposal of several putative functions for this protein in the responses of leech nervous tissue: (i) a role as an oxygen supplier for metabolism, (ii) a role as a trap for reactive oxygen species and NO, protecting cells from cell death and (iii) a role as an antibacterial factor depriving bacteria of iron.²⁸

The data from these initial studies of the effects of bacterial toxins show that the leech CNS is able to respond in an intrinsic manner to a septic stimulus, mounting a “neuroimmune” response. Careful study of potential roles of the identified proteins will be essential to fully understand the mechanisms involved, but some candidates can be proposed on the basis of the proteins identified in this study (ref. 26): (i) cytoskeletal rearrangements potentially responsible for morphological changes, cell migration, vesicular trafficking and/or phagocytosis, (ii) modulation of synaptic activity, (iii) calcium signalling and (iv) unfolded protein response controlling the functionality of proteins affected by the stress generated by the sepsis. The involvement in innate immunity of some proteins or protein families we identified here as previously been described by transcriptomic studies, but at the peripheral level and not within the nervous system.^{29,30} Moreover, the protein families involved in the immune response of the medicinal leech nervous system appear to also be involved in nerve regeneration, as shown by Blackshaw *et al*, also in the medicinal leech³¹ (discussed above) and by Perlson *et al* in *Lymnaea stagnalis*³² (Table 1).

These observations suggest certain parallelism between CNS defence mechanisms and CNS regeneration leading the possibility of the involvement of neuroinflammation in such phenomena in leeches. Modulation of neuroinflammation after a crush seems to be necessary but highly regulated in the CNS during regeneration through cannabinoids as we previously demonstrated.^{22,33-36} Interestingly, evidences that microbial infection triggers the leech brain regeneration (Fig. 3) has led us to perform some biochemical studies focused on antimicrobial peptides present in the leech CNS and expressed in course of infection or trauma. Two novel antimicrobial peptides, *Hm*-lumbricin and neuromacin have been fully characterized.³⁷ Neuromacin and *Hm*-lumbricin exert bactericidal activities against Gram positive bacteria without any haemolytic properties.³⁷ We have observed that in addition to exert antimicrobial activities, *Hm*-lumbricin and neuromacin have regenerative effects on the leech CNS.³⁷ The capacity of both peptides to promote the regeneration of the leech nerve cord was tested *ex vivo* by adding the neuromacin and/or *Hm*-lumbricin antibody(ies) to axotomized nerve cords in presence of killed bacteria.³⁷ Due to the presence of bacteria, the reconnection process should have started two days post-axotomy. It appeared that the presence of antibodies in the culture medium blocked the regeneration process since no reconnection was observed even seven days post-axotomy.³⁷ These observations were corroborated by the data obtained by adding native neuromacin to axotomized nerve cords under aseptic conditions. Nerve repair was evident sooner in the presence of neuromacin, reconnection starting in less than one day instead of four without an exogenous contribution in neuromacin. The participation of endogenous neuromacin and *Hm*-lumbricin in the neural repair is sustained by the accumulation of both peptides at the lesion site upon bacterial challenge of injured nerve cords.³⁷ Further investigations based on single cell RT-PCR analysis and on immunohistochemical analysis of a model, developed by our group, of leech CNS almost completely devoid of microglial cells allowed

Table 1. Similarities between proteins involved in innate immunity within invertebrates and those involved in central nervous system regeneration within Lophotrochozoaria, all identified by proteomic or transcriptomic approaches

Immunity		Leech CNS	Leech Periphery	Insect Periphery
Cytosk.	Microfilaments IF	Tropomyosin gliarin	Tropomyosin-2 Actin-2	Myosin II reg. light-chain Actin 5C
Calcium	Calcium sensor Others	NCS-2/Neurocalcin		Calmodulin, sarc. CaBP1
Metabolism	AA/nt me- tabolism Energy Others	AA dehydrogenase ATP synthase β subunit Acetyl transferase	Aldehyde dehydr.	Aldehyde dehydr. ATP synth. β subunit
Hsp and chaperones	Cyclophilin/PPI PDI	Cyclophilin PDI	PPI	FK506-BP-PPI, Cyclophilin PDI (ERp60), CaBP1-PDI
Metal- oxidation	Resp. molecule Others	Neurohemerythrin		Hemocyanin Thioredox., Transf., ferritin
Regeneration		Leech CNS	Mollusc CNS	
Cytosk.	Microfilaments IF Microtubules	Protein 4.1 Synapsin α - and β -tubulin	Tropomyosin Intermediate filament Tubulin	
Calcium	Calcium sensor Others	Calmodulin-like	Calmodulin, Calbindin Calpain	
Metabolism	AA/nt metabo- lism Energy	ATPase inhibitor	Glutamine Synthase ATP Synthase	
Hsp and chaperones	Cyclophilin/PPI PDI Others	Hsp90	Cyclophilin PDI Hsp60, 14-3-3	
Metal- oxidation	Resp. molecule Others	Myohemerythrin COX I	Peroxiredoxin, Ferritin	

AA, aminoacyl; ATP synth., ATP synthase; calcin., calcineurin; CNS, central nervous system; COX I, Cyclooxygenase I; Cycloph., cyclophilin; dehydr., dehydrogenase; IF, intermediate filaments; nt, neurotransmitters; phosph., phosphatase; PK, protein kinase; PPI, peptidyl-prolyl cis-trans isomerase; sarc., sarcoplasmic; reg, regulatory; resp. molecule, respiratory molecule; Transf., transferring. Used with permission from Vergote D et al. Proteomics 2006;6(17): 4817-25.

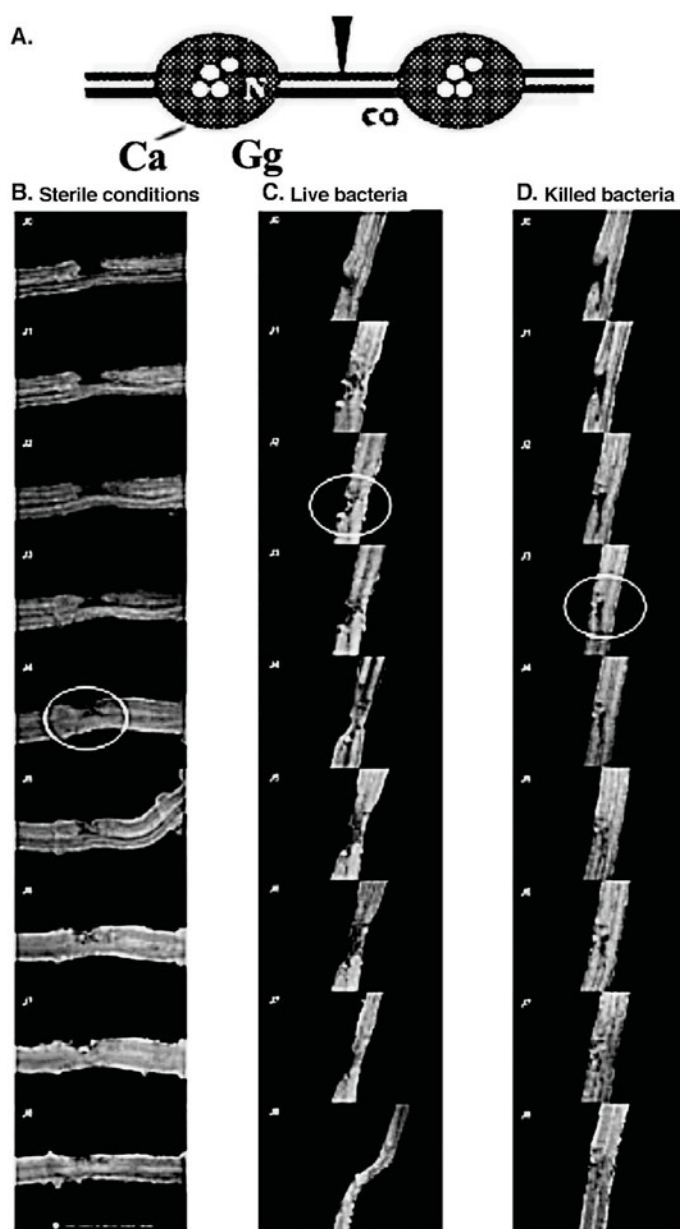


Figure 3. Effects on nerve regeneration of exposure of excised leech CNS to live or heat-killed bacteria (A) Diagram of the leech CNS in culture preparation. Neuron cell bodies (N) within ganglia (Gg) project axons into connectives (co) towards adjacent ganglia. V indicates the location of the cut of one of the two connectives linking two segmental ganglia. Microglial cells, evenly distributed in the nerve cord, are represented by dots. The nervous system is protected by a fibrous capsule (Ca). B-D) Sequential micrographs, taken 24 hr apart, from one (J1) to eight days (J8) post-axotomy, documenting the regeneration of the severed connective nerve. B) Preparation in sterile culture medium, (C) incubated with live bacteria and (D) incubated with killed bacteria. Used with permission from Schiskorski et al. *J. Immunol* 2008;181(2): 1083-95. ©2008. The American Association of Immunologists, Inc.

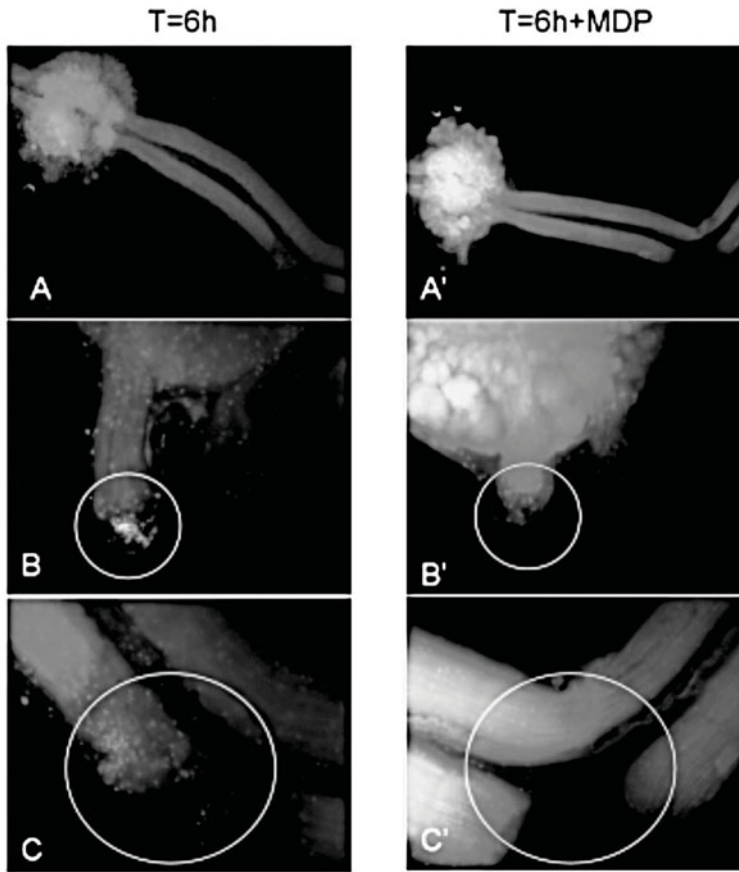


Figure 4. Effects on the recruitment of microglial cells of exposure of injured leech CNS to Muramyl DiPeptide (MDP). A) without addition of MDP to the medium culture, an accumulation of microglial cells is observed by nuclear staining at the injured site of both lesioned lateral (B) and interganglionic connective (C). (A', B', C') This cell accumulation is not visible anymore post treatment with MDP.

us to conclude that the presence of both *Hm*-lumbricin and neuromacin at the axotomized site implicates peptide production by neurons and by the microglial cells recruited at the lesion site.³⁷ However silencing studies have not allowed connecting such antimicrobial peptides with *Hm*TLR1 receptor. Moreover, although bacteria cocktail enhance leech brain regeneration, Muramyl DiPeptide (MDP) blocks microglia migration (Fig. 4) which is line with the data obtained on macrophages migration, also inhibited by MDP.^{38,39}

Taken together, these data confirm the immune role of the leech microglia and the presence of specific sensing receptor in these cells. The next step is the characterization of these receptors and effectors contained in leech brain. For this purpose, we started by *in silico* analyses through medicinal leech adult EST obtained by a consortium between Professor E. Macagno, Professor Terry Gaasterland and Professor Michel Salzet. 91,233 transcripts were obtained from Genoscope (France) and JGI (USA) before annotated and 31,232 sequences were obtained.⁸

Leech Brain Immune Receptors and Effectors

Medicinal Leech Toll-Like Receptors (HmTLRs)

Five medicinal leech Toll-like receptors (TLRs) have been detected, one has already been fully characterized and data from the other ones are due soon. The first *HmTLR* characterized is presented in Figure 5.⁴⁰ TLRs in general, share similarities in their extracellular Leucine Rich Repeat (LRR) and their intracellular Toll/IL-1 Receptor (TIR) domains. The TIR domain plays a central role in TLR signaling. All TLRs contain a cytoplasmic TIR domain, which, upon activation, acts as a scaffold to recruit adaptor proteins. It is well established that the differential recruitment of adaptors to TLRs provides a significant amount of specificity

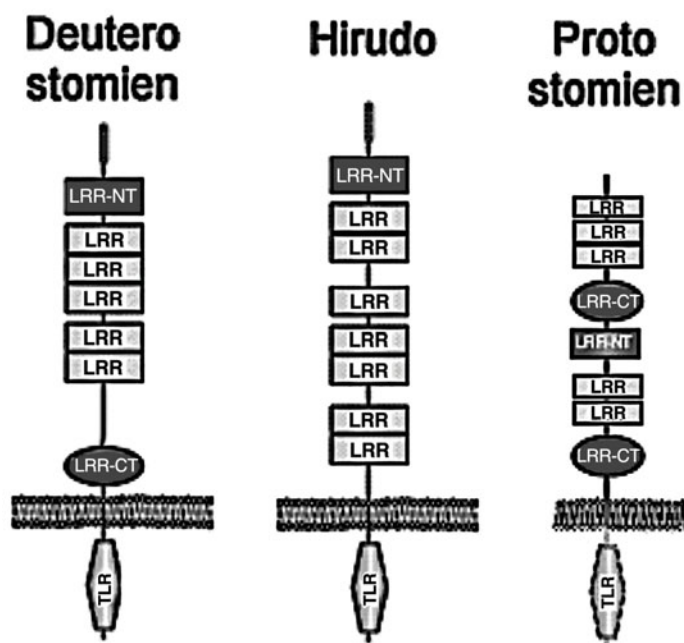


Figure 5. Structure comparison of *HmTLR1* and Protostomian and Deuterostomian TLRs.⁴⁰ SMART™ sequence analysis of the Nterminal part of *HmTLR1* revealed the presence of one LRRNT followed by six LRRs. *HmTLR1* presents the originality to exhibit an array of LRRs capped by one LRRNT only. By contrast to most TLRs described in invertebrates and vertebrates species, no LRRCT domains were identified from the analysis of the ectodomain of *HmTLR1*. Blastp analysis was realized on the entire amino acid sequence of *HmTLR1*. Data reveal great homologies mainly with TLR13s characterized in vertebrate species such as the mouse *Mus musculus* and the opossum *Monodelphis domestica* and in a lesser extent with TLRs of animals living in freshwater such as the zebra fish *Danio rerio*, the goldfish *Carassius auratus*, the salmon *Salmo salar* or again the rainbow trout *Oncorhynchus mykiss*. No homology with molecules identified in other lophotrochozoa was noticed. A second step, Blastp analysis of the LRR and the TIR domains of *HmTLR1* were performed separately, in order to get information on the function and the signalling pathway associated with this receptor respectively. The LRR domain of *HmTLR1* significantly (e-values < e-12) matches with the sequences of LRRs implicated in (i) pathogen recognition such as the LRR domain of the TLR3 and those of some Variable Lymphocyte Receptors (VLRs) and (ii) with the LRRs of vasorin, decorin and netrin known to participate in tissue remodelling and/or axonal guidance in vertebrates. Thus, both regenerative and immune functions could be attributable to this receptor. Concerning the TIR domain of *HmTLR1*, Blastp analyses evidence a great percentage of homology with the TLR13.

to the TLR-signalling pathways. Among these adapter proteins MyD88 and TRIF are now considered as the signalling ones and hence the TLR pathways can be categorized as MyD88-dependent and TRIF-dependent. The LRR domain is an extracellular domain implicated in the detection of pathogens. Based on the organization of the extracellular LRR array, two types of TLRs have been described. Vertebrate TLRs have an array of LRRs capped by cysteine-rich domains located at the N- and C-terminal LRR domains (LRRNT and LRRCT, respectively). By contrast, most of invertebrate TLRs also contain LRRNT and LRRCT domains, but instead of capping the LRR array, these are located within the array in a tandem orientation. Interestingly, leech *HmTLR1* presents the originality to exhibit an array of LRRs capped by one LRRNT only sharing sequence similarity with mouse TLR3. Based on EST medicinal leech sequence and *Helobdella* genome, four other *HmTLR* has been detected and their complete characterization is now in progress.

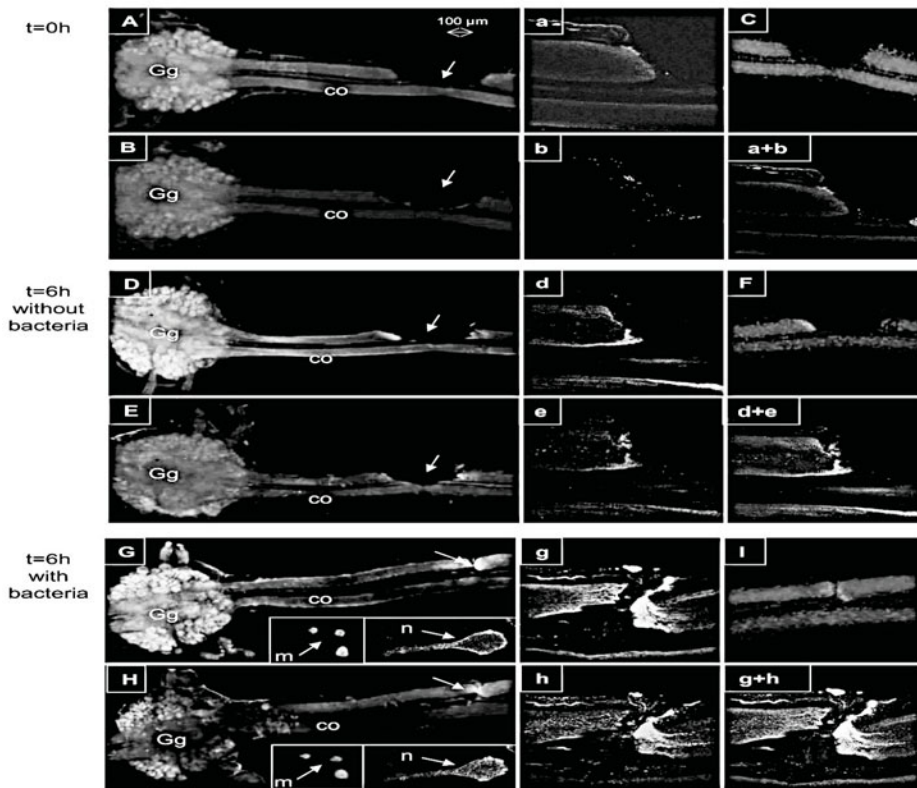


Figure 6. Co-appearance of *Hmp43/EMAPII* (Green) and *HmTLR1* (Red) in the injured CNS incubated or not for 6 h with killed bacteria. Double staining of injured nerve cords was performed at $t = 0$ and $t = 6$ h post-axotomy using the fluorescent nuclear dye Hoechst 33258 (C, F and I) with both the anti-p43/EMAPII (A, D and G) and the anti-*HmTLR1* (B, E and H) polyclonal Ab. Immunodetection was performed using green-labelled anti-EMAPII and red labeled anti-*HmTLR1* secondary Ab. The results demonstrate an accumulation of *Hm-EMAPII* (G, g) and *HmTLR1* (H, h) at the lesion site 6 h after axotomy in the presence of bacteria from reference 40. Used with permission from Schiskorski et al. *J. Immunol* 2009;183(11): 7119-28. ©2009. The American Association of Immunologists, Inc. A color version of this figure is available at www.landesbioscience.com/curie.

We recently demonstrate that *HmTLR1*⁴⁰ is localized in both neurons and microglia and expressed upon septic trauma (Fig. 6). *HmTLR1* is co-expressed with a cytokine related to the Endothelial Monocyte Activating Polypeptide (*HmEMAPII*) sharing chemoattractive activity (Fig. 6). The complete intracellular trafficking upon pathogen challenges has to be undertaken to confirm the functional similarity between *HmTLR1* and mammalian TLR3.

***HmTLR1* and the Leech Cytokine (EMAPII)**

Functional studies using silencing studies have not allowed connecting such antimicrobial peptides with *HmTLR1* receptor. By contrast, a cytokine sharing microglia chemoattractant activity recently characterized by our group in the medicinal leech e.g., *HmEMAPII*,⁴⁰ shown based on RNAi silencing qPCR, western blot experiments (Fig. 7) and biological tests an association to *HmTLR1*.⁴⁰ *HmEMAPII* is processed from *Hmp43* like mammals EMAPII. We hypothesized that *HmEMAPII* could exert a chemoattractant effect on microglial cells as mammalian EMAPII does on monocytes. The chemoattractive effect of *HmEMAPII* is blocked when an antihuman EMAPII antibody underscoring for the first time the ability of EMAPII to exert chemotactic effect toward microglial cells through CXCR3.⁴⁰ Leech CXCR3-related receptor cloning is on the way. These data points out that *HmTLR1* is linked to leech EMAPII and gives for the first time an immune function to a TLR in a noncyclozoan model *i.e.*, in an invertebrate model different from *Caenorhabditis elegans* and *Drosophila melanogaster*.

All together, these data reflect that the medicinal leech express *HmTLR* related to mammalian TLR. The first receptor characterized showed an intracellular localization and *HmTLR1* is linked to the cytokine related to EMAPII which exerts chemottractive effect after brain trauma or upon pathogen challenge. All these data confirm for the first time the presence of a complete TLR-signalling-Cytokine pathway implicated in immune response in medicinal leech nervous system. Such complex is also present in mammals reflecting again a co-evolution between the medicinal leech and its host mammals. It has also to be noted that such mechanism conservation is in line with a common origin of nervous system centralization between annelids and vertebrates as shown in the polychaete, *Platynereis dumerilii*.⁴¹⁻⁴³ Thus, the data presented above indicate that major players in innate immune response like danger sensing receptors coupling to cytokines or antimicrobial peptides and microglia are present in leech and strongly resemble that in vertebrates.

THEROMYZON TESSULATUM AS A MODEL FOR STUDYING THE PERIPHERAL IMMUNE RESPONSE

T. tessulatum is an ectoparasite of aquatic birds. Its life cycle was arbitrarily subdivided in stages (these are not larval stages) defined by taking, as indicators, the three blood meals. The third stage which corresponds to the gametogenesis phase is characterized by an important water uptake making the collection of the body fluid easy (Fig. 8). For this reason, *T. tessulatum* constitutes a convenient model for studying the antimicrobial response which takes place at the systemic level in coelomic fluid.⁴⁴ As a comparison, the medicinal leech has a parenchymatous body, coelomic cavities are reduced and the botryoidally tissue is immersed in a connective tissue. This makes the collection of the body fluid impossible. The antimicrobial response of *T. tessulatum* was investigated at the molecular level by

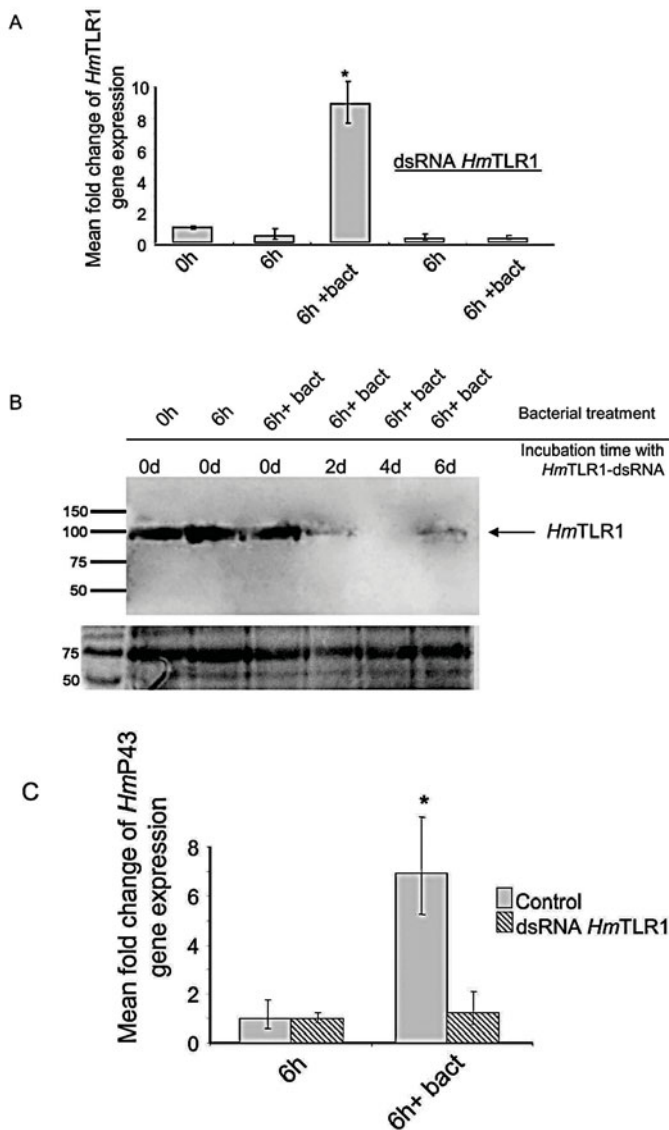


Figure 7. Impact of the *HmTLR1* gene silencing on induction of the *HmP43/EMAPII* gene in the leech CNS incubated with bacteria. **A)** The efficiency of the knock-down was quantified by measuring the level of *HmTLR1* expression in nerve cords incubated with or without dsRNA. Data are expressed as relative levels comparatively to the basal level of expression measured in nerve cords processed immediately after sampling (0 h). *HmTLR1* expression was quantified after 6 h of culture without bacteria (6 h) or with a mix of heat-killed Gram+ and Gram- bacteria (6 h + bact), revealing an induction of *HmTLR1* gene under septic conditions. The bacterial induction of *HmTLR1* is significantly reduced when the CNS is incubated for 4 days with *HmTLR1*-dsRNA (**B**) Western blot analyses of *HmTLR1* protein level in the same conditions as for Figure 1. Best protein extinction is observed at 4 days (4d) of incubation with specific dsRNA confirming the efficiency of the knock down and the specificity of the anti *HmTLR1* antibody. d, day (**C**) *HmTLR1* gene silencing abolished the bacterial gene induction of *HmEMAPII* observed in the control (without dsRNA *HmTLR1*), indicating a role of the *HmTLR1* in the gene regulation of this cytokine in the leech CNS under septic conditions.⁴⁰

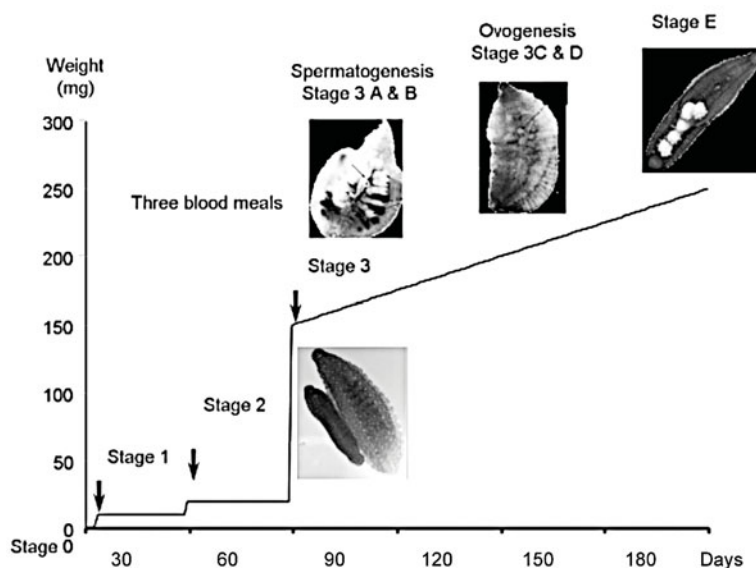


Figure 8. Life cycle of *Theromyzon tessulatum* from reference 61.

focusing on the antimicrobial peptides (AMPs) released in the coelomic fluid and at the cellular level by determining the immune functions of the coelomocytes.

AMPs of *T. tessulatum*

Three antimicrobial peptides (AMPs) were isolated and fully characterized from the body fluid of *T. tessulatum*. These are theromacin, a cysteine rich AMP exhibiting bactericidal activities, theromyzin an anionic peptide with bacteriostatic properties⁴⁴ and peptide B an anionic peptide matured from a neuropeptide precursor, proenkephalin A (PEA).⁴⁵ They all present an activity directed against Gram positive bacteria. Recently, a cDNA encoding a peptide presenting high percentage homologies with lumbricin-1, an AMP firstly characterized from the earthworm *Lumbricus rubellus* was cloned in *T. tessulatum* (Fig. 9).

Theromacin belongs to the cysteine rich AMP family. In invertebrates, most of them share the disulfide array of the insect/arthropod defensin.⁴⁶ In addition to having ten cysteine residues instead of six, theromacin does not harbor this consensus sequence. Theromacin by contrast with neuromacin (see before) has never been evidenced in other lophotrochozoan models or in ecdysozoan and thus seems to be restricted to leeches.

Lumbricin is a linear peptide without any posttranslational modifications. As the majority of AMPs described in the literature, theromacin and lumbricin possess a global positive charge presumably allowing their interaction with the negatively charged bacterial membrane (Fig. 10).

Theromyzin and peptide B, in contrast to theromacin and lumbricin, are anionic molecules. The mode of action of anionic AMPs is still unknown even if several hypotheses have been advanced. In vertebrates, AMPs with anionic properties were evidenced in the human and the sheep lung⁴⁷ AMPs are anionic because of homopolymeric regions of aspartate and require zinc as a cofactor for bactericidal activity.⁴⁸ Histatins,

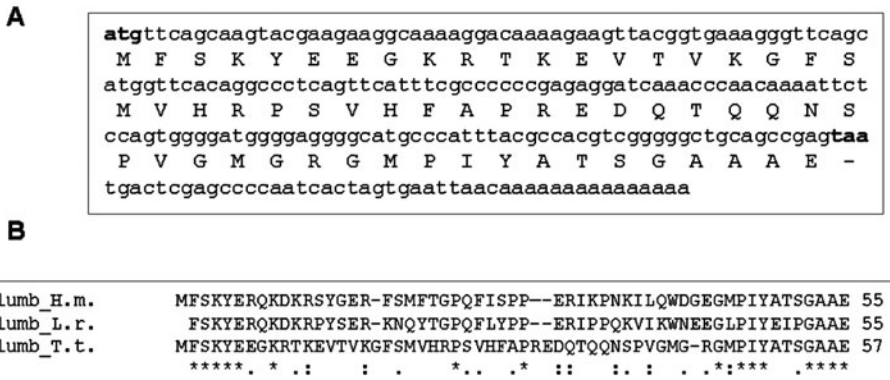


Figure 9. A) Nucleotide sequence of *Theromyzon tessulatum* lumbricin cDNA. The deduced amino acid sequence of the open reading frame is presented under the nucleotide sequence. B) Alignment of the *Tt* lumbricin (Lum T.t.) with the lumbricins characterized from the earthworm *Lumbricus rubellus* (Lum L.r.) and from the medicinal leech *Hirudo medicinalis* (Lum H.m.).

a family of histidin rich AMPs found in human saliva, also need the presence of zinc ions for bactericidal activities. Circular dichroism studies showed that the antimicrobial activities of histatin-5 require a conformational change that results from the interaction of the peptide with both zinc ions and negatively charged membranes.⁴⁹ The abundance of histidine residues at the N-terminal part of theromyzin could argue in favor of some common structures between the leech antibacterial peptide and histatins.

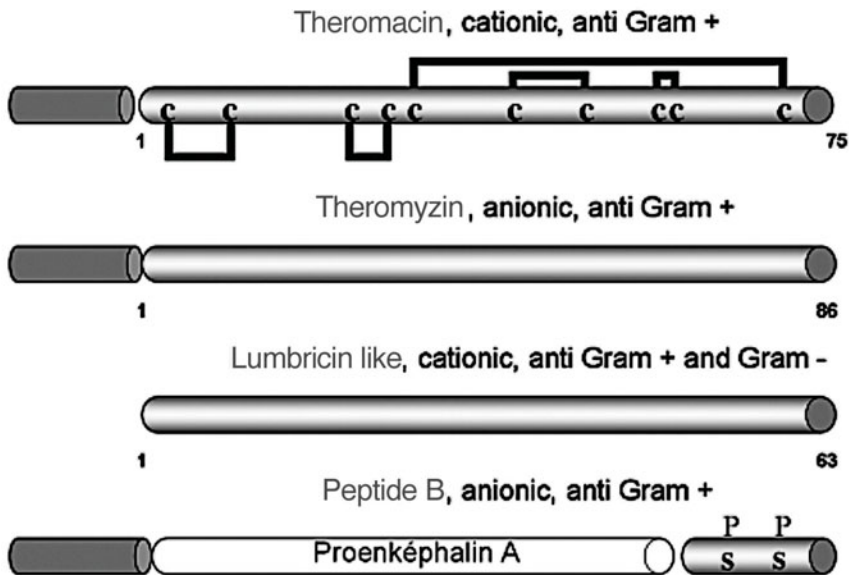


Figure 10. AMPs characterized in *T. tessulatum* from the coelomic liquid.⁵⁷

Consequently, it clearly appears that leeches present a relatively large variety of AMPs. It is interesting to remark that at present no defensins have been isolated from leeches and from annelids in general. Defensins which are considered as the most widespread family of invertebrate AMPs have not been found neither in the genomes of the leech *Helobdella robusta* and the polychaeta *Capitella* nor in the EST libraries of the earthworm *Lumbricus terrestris*, *Eiseinia fetida* and *Hirudo medicinalis*. Reciprocally most AMPs described in annelids have not been found in the genomes of ecdysozoan invertebrate such as *C. elegans* and *Drosophila melanogaster*.

Expression Site and Regulation of the AMPs Synthesis in *T. tessulatum*

Theromacin, *theromyzin* and *Tt lumbricin* genes are preferentially expressed in large fat cells (LFC) evenly distributed in the leech and in contact with the coelomic liquid from which the AMPs were purified. Their transcriptional level is enhanced after bacteria challenge evidencing a regulation of the leech AMPs similar to that of the insect antimicrobial peptides genes. Indeed, in the fruit fly, genes encoding antibiotic peptides are rapidly induced following a septic injury.⁵⁰ The similarity between the antibacterial response of the leech and those of holometabol insects is also supported by the functional resemblance between the leech LFC and the insect fat body which possess the common capacity to produce egg-yolk proteins.⁵¹

Moreover, based on our first investigations, no difference in gene expression was observed after Gram positive or Gram negative injection suggesting that the antibacterial response of *Theromyzon* is aspecific. This nonspecificity has also been assumed in *Drosophila* until the work of Lemaitre *et al* demonstrated that the humoral antimicrobial response of the fruit fly discriminates between various classes of microorganisms and mounts a response that is adapted to the infection.⁵² That suggested that in a more natural mode of infection the leech could also adapt its antimicrobial response, what was recently confirmed in our annelid models by using bacteria living in the environment of the leech.³⁷

The peptide sequences deduced from the *theromacin* and *theromyzin* genes contain putative signal peptides, indicating that mature peptides correspond to secreted molecules. As for the lumbricin like characterized in the medicinal leech, the *Tt lumbricin* precursor lacks the typical signal peptide. We assume that *Tt lumbricin* as demonstrated in *Hirudo* could be secreted through a nonconventional mechanism already observed but still unexplained for several molecules in mammals also.

Peptide B is not produced by the LFC although it was also isolated from the body fluid of the leech. Its precursor, PEA, was immunodetected into circulating coelomocytes suggesting that peptide B could be released from these cells. In contrast to the other leech AMPs, the production of peptide B seems to be more regulated at the translational level by the enzymes implicated in the PEA processing than at the transcriptional level. Of equal importance is the finding that enkephalin such as methionin enkephalins (ME) and peptide B are simultaneously released from PEA. Invertebrate and vertebrate immunocytes contain delta 2 opioid receptors that appear to mediate activation of these cells. In this regard, ME can be envisioned to activate immunocytes and provide a chemotactic signal to further stimulate cell recruitment.

However, since this process may take many minutes to accomplish, the bactericidal peptide B may cover this activation latency period. In this scenario, peptide B is broken down with time, it could release during this time, the heptapeptide MERF. Since we demonstrated that this peptide was able to interact with delta 2 opioid receptors, MERF

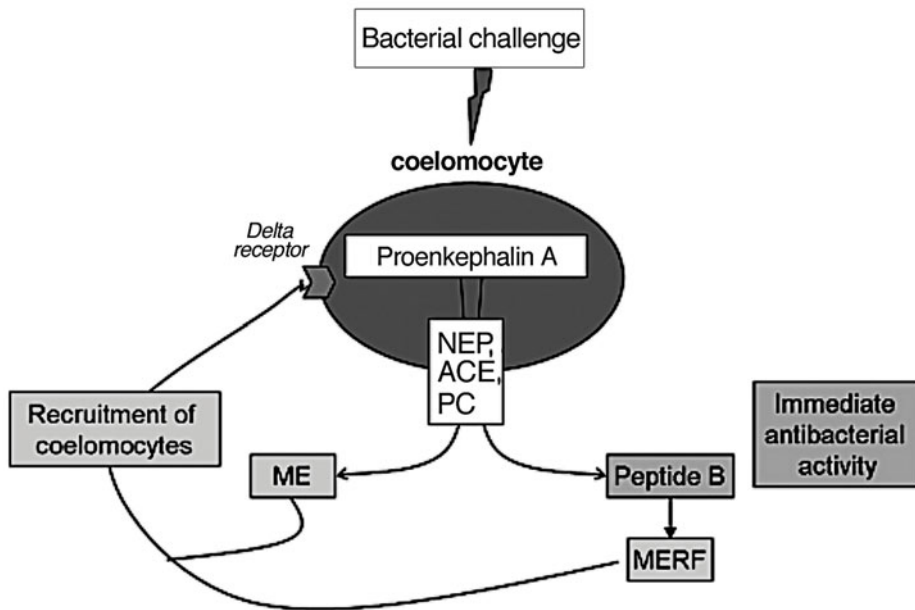


Figure 11. Illustration for the involvement of PEA derived peptides in the immune response of *T. tessulatum*. Upon an initial stimulus, i.e., bacterial challenge, smaller bioactive peptides Met-enkephalin (ME) and peptide B processed by different enzymes (PC, NEP, ACE) are released into the coelomic fluid of the leech. Peptide B plays its antibacterial role and is cleaved a second time into Met-enkephalin arg-phe (MERF) which, as ME does, can recruit coelomocytes.

could keep the rate of immunocyte activity.⁴⁵ This hypothesis is supported by previous studies demonstrating an immune activating role for MERF in human and invertebrate immunocytes. MERF as well as ME was shown to induce rounded invertebrate immunocytes to become mobile and amoeboid as well as to initiate chemotaxis. (Fig. 11).

Whatever the mode of regulation is, the accumulation of AMPs into the coelomic fluid after septic injury suggests that these peptides play their antimicrobial activities through a systemic action. Moreover, the presence of theromacin and theromyzin in the intestinal epithelial cells and at the epidermis level also evokes participation in epithelial defense and/or in the control of the symbionts. The localization of antibiotic molecules in the gastrointestinal tract has also been reported in insects and in vertebrates where they provide a rapid local immune response against exogenous pathogens brought in during feeding.⁵³ Theromacin, lumbricin and theromyzin were detected in the mucous covering the animal. That reminds the local defensive response reported in frogs in which antibacterial peptides secreted in the mucous prevent bacteria colonization and/or subsequent infection.⁵⁴

As for lumbricin-1 in *L. rubellus*, physiological events occurring during gametogenesis phase appeared to be inducers of the AMPs gene expression in *T. tessulatum*. These data suggest that several hormonal factors implicated in sexual maturation may participate in the induction of genes encoding AMPs in annelids as described in *Drosophila* by Meister et al.⁵⁵ Interestingly, leech AMPs were detected

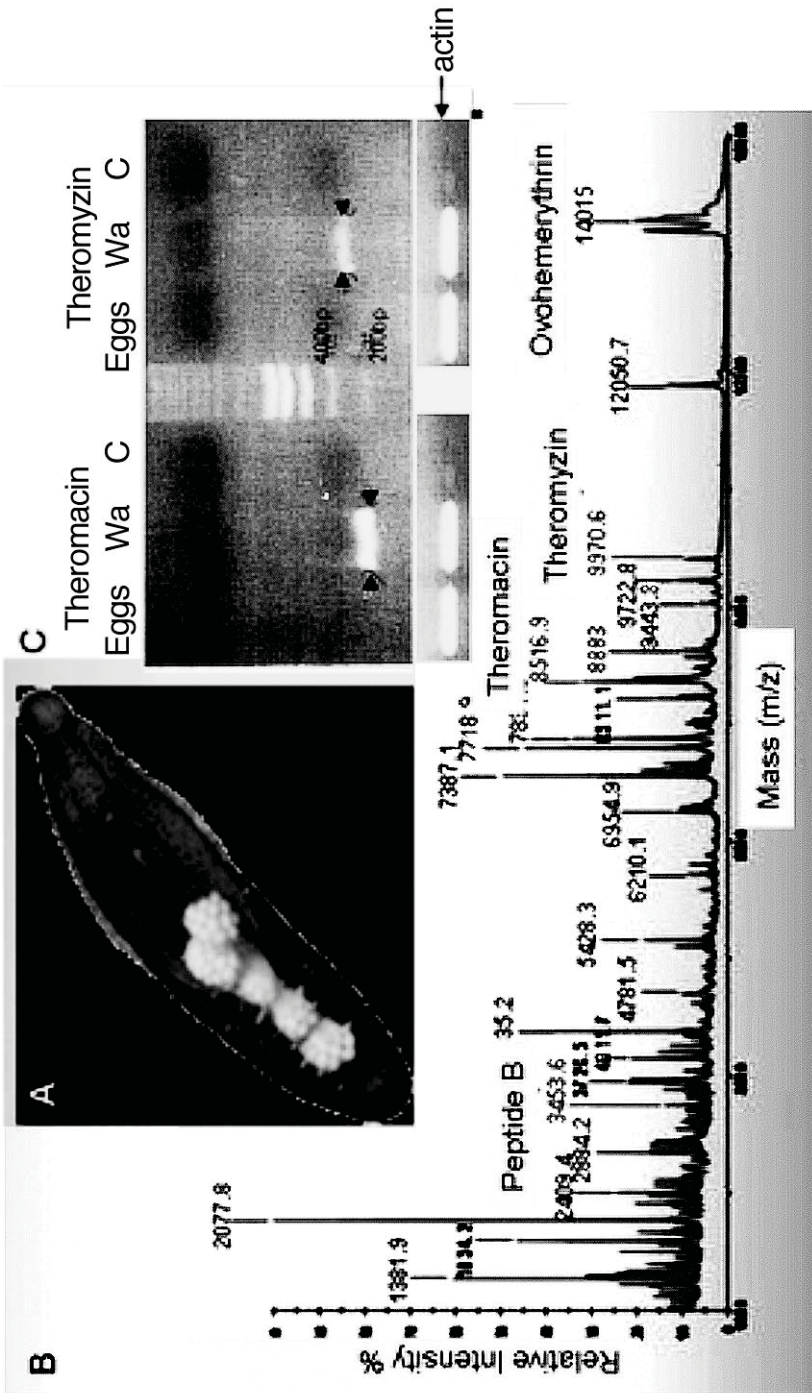


Figure 12. A) Ventral view of a Stage 3 *Theromyzon tessulatum* showing eggs (yellow balls) regrouped within five cocoons. B) Detection by mass spectrometry of molecular masses corresponding to theromyzin and theromycin in eggs extracts. C) Transcripts encoding theromycin and theromyzin are amplified by RT-PCR in whole animals (Wa) but not in eggs. C corresponds to the water control. A color version of this figure is available at www.landesbioscience.com/curie.

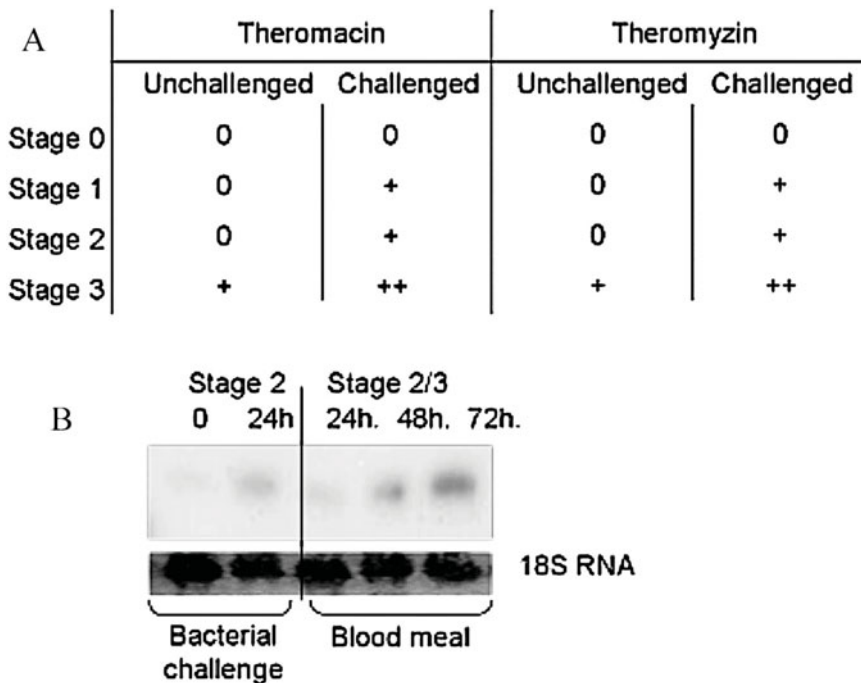


Figure 13. A) Table presenting the detection of transcripts coding for theromacin and theromyzin in leeches pricked or not with a mix of bacteria, at different stages of their life cycle. B) Northern blot showing that the theromacin gene expression is inducible upon bacteria challenge in Stage 2 leeches and after the last blood meal which marks the transition between the Stage 2 and 3.

in the eggs by mass spectrometry analysis whereas the transcripts were not amplified by RT-PCR (Fig. 12).

These results suggest a vertical transmission of the PAMs which could exert a protective role against bacteria during eggs development. AMPs may accumulate into the eggs by being captured from the body fluid of the mother through a pinocytosis mechanism before laying as described for egg yolks proteins and/or from the mucous covering the eggs after laying. A mass corresponding to ovohemerythrin, an egg yolk protein abundantly presents in the leech body fluid was detected by mass spectrometry analysis of eggs extracts supporting the first hypothesis without excluding the second one since AMPs were also detected in the mucous covering the leech. The study of the gene expression in course of the post-embryonic development demonstrated that *T. tessulatum* starts to synthesize its own AMPs from the Stage 1 of its life cycle and that a bacterial challenge is necessary for observing this synthesis. Indeed, a basal level of transcripts encoding theromacin and theromyzin was detected in unchallenged Stage 3 leeches only. We presume that AMPs are expressed at a basal level in adults for providing an immune protection to the eggs by a vertical transmission of the antibiotic molecules. By contrast, a bacterially inducible response is observable from the Stage 1 to the Stage 3 suggesting that leeches acquire the ability to establish an immune response after their first blood meal (Fig. 13).

Consequently, *T. tessulatum* is an original invertebrate model which has developed two modes of fighting infections by AMPs: (i) storage of antibacterial peptide derived

from PEA and release of the peptide into the coelomic fluid after immune challenge (ii) induction after septic injury of gene coding for more classical AMPs, mainly in LFC and rapid release into the body fluid of the antibiotic peptides. Data collected from our group (unpublished) suggest that the same AMPs participate to the systemic antimicrobial response of the medicinal leech, *Hirudo medicinalis*. Interestingly, the PEA processing appeared to be very well conserved in course of evolution since we have demonstrated that the same mechanism is observable during surgery in human patients undergoing cardiopulmonary bypass.^{56,45}

The Cellular Immune Response

Annelids are primitive coelomates known to possess specially developed cellular immunity against microorganisms including phagocytosis, encapsulation and spontaneous cytotoxicity against allogenic or xenogenic cells.^{57,58} In leeches, our group has characterized three distinct populations of coelomocytes in *T. tessulatum* (Fig. 14).

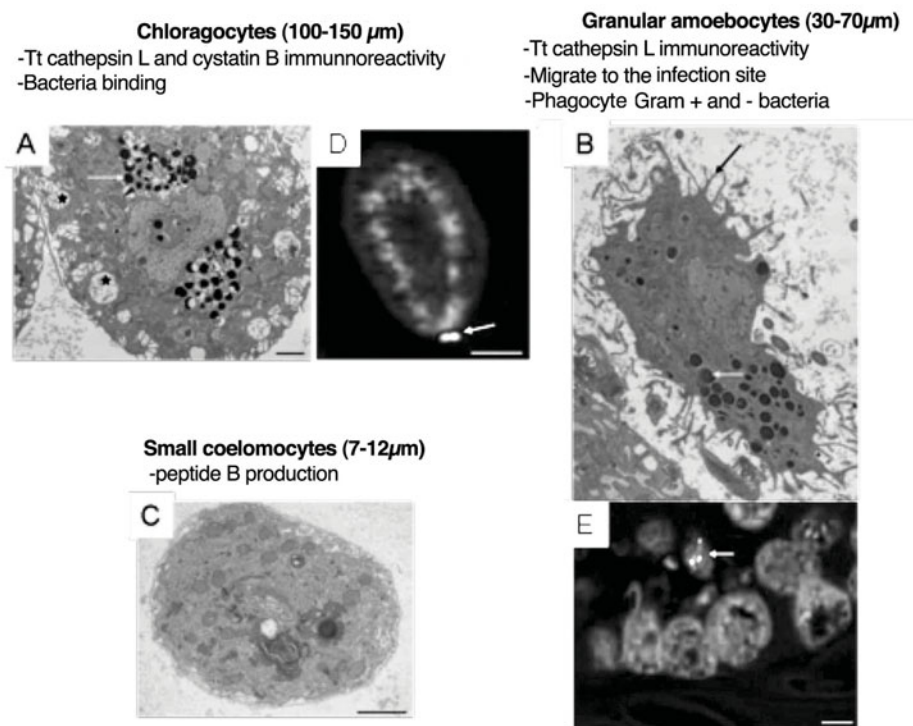


Figure 14. Coelomocytes of *T. tessulatum*. A) Large coelomocytes present large electron-dense granules (white arrow) and electron-lucent vesicles (black stars). B) Intermediate size coelomocytes show long cytoplasmic pseudopods (black arrow) and large electron-dense granules (white arrow). C) Small circulating cells are rich in endoplasmic reticulum and small granules. D-E) Animals were injected with FITC-labelled bacteria (killed *M. luteus* or *E. coli*). After 24 hours of incubation, bacteria are still observed near chloragocytes (D, white arrow) but phagocytosed bacteria are observed in granular amoebocytes only (E, white arrow).

These are constituted by chloragocytes, granular amoebocytes and small coelomocytes.³⁰ Leech chloragocytes are the only cells expressing both the *Tt* cathepsin L and the *Tt* cystatin B. Granular amoebocytes present an immunoreactivity to the anti *Tt* cathepsin L antibody (Ab) although the small coelomocytes are not recognized by neither the anti *Tt* cathepsin L Ab nor the anti *Tt* cystatin B Ab. The immune functions were investigated by essentially focusing on the phagocytic activity and the migrating property of these circulating cells. As resumed in the Figure 14, the granular amoebocytes are able to migrate to the injection site of microorganisms and to phagocyte without any apparent distinction both killed Gram positive and Gram negative bacteria. The molecular mechanism of recruitment may imply enkephalin peptides derived from the PEA maturation process, such as ME or MERF. By contrast, leech chloragocytes are not able to phagocyte bacteria. However, while no phagocytosis was detected, confocal microscopy analysis evidenced chloragocytes-bacteria interaction suggesting the presence of recognition molecule expressed at the surface of these cells (Fig. 14D). Leech chloragocytes may also be implicated in encapsulation reactions as described in oligochaeta annelids. The third type represented by the small coelomocytes presents the morphology of invertebrate hyaline cells i.e., a cytoplasm deprived of granules. However the leech cells have been considered not to be hyaline cells due to their incapacity to phagocytose killed bacteria. Interestingly, our group has very recently observed that the use of live bacteria was a prerogative for inducing the phagocytosis process in a population of leech blood cells. Thus, the small coelomocytes should be incubated with live bacteria to determinate whether they could be assimilated to hyaline cells or not.

This heterogeneous population of coelomocytes reminds the population described in other annelids. Numerous studies performed in oligochætes as *Lumbricus sp.* and *Eisenia sp.*⁵⁹ used optical and electron microscopy. Although monoclonal antibodies were carried out against various invertebrate taxa, the majority of the studies related to insect hæmocytes. De Eguileor *et al* identified three coelomic cell populations, i.e., macrophage-like, NK-like and granular cells, using human monoclonal antibodies in the hirudinea *Glossiphonia complanata*.⁵⁸ Engelmann *et al* produced monoclonal antibodies against coelomic cells in *Eisenia fetida* earthworm.⁶⁰ While anti-EFCC1 antibody (*Eisenia fetida* coelomocyte differentiation cluster) is able to recognize antigenic motifs on various tissues, three other antibodies named anti-EFCC2, anti-EFCC3 and anti-EFCC4 allowed to respectively discriminate chloragocytes, hyaline amoebocytes and granular amoebocytes.⁶⁰ Because oligochætes and leeches are closed relatives, it should be interesting to test these anti-EFCC antibodies for discriminating our coelomic cells and possibly identify the presence of phagocytic hyaline cells in *T. tessulatum*.

Unlike other invertebrates, one of the particularities of annelids is to possess a closed circulatory system separated from the coelomic cavity. Our group is investigating the immune function of the circulating blood cells of the medicinal leech. Most of the reported data being focused on coelomic cells, the obtained results will constitute the first description of the function of the blood cells in an annelid. Moreover since the leech central nervous system permanently baths into the blood, these data may open up new avenues for discovering the impact of the immune response on the neural repair of the medicinal leech.

CONCLUSION AND PERSPECTIVES

The observations we have reviewed in this chapter attest to the broad range of inquiry, from structure to function, and to the breadth of the techniques currently employed to study the defence, repair and maintenance of the leech nervous and immune systems. Clearly, the increasing application of biochemical and molecular genetic tools is beginning to yield insights into the nature of the molecular mechanisms responsible for these phenomena. But, while progress is being made, it is also clear that there is a strong need to accelerate the implementation and application of genomic, transcriptomic and proteomic tools to the leech. The leech model has many important advantages, perhaps the most important being the ability to bridge from the immune response to neural repair, in the context of having, or the possibility of having, detailed knowledge about all the neurons in the CNS and thus an unparalleled level of completeness. Most other invertebrate systems, as well as all vertebrate systems, under current study, can only afford partial, because many or most of their neurons are inaccessible and only population properties are attainable. However, other systems, including *C. elegans*, *Drosophila*, zebrafish, and some rodents, greatly benefit from the possibility of doing standard genetics, their shorter reproductive cycles (particularly in comparison to *Hirudo*, not as much with respect to small leeches like *Helobdella*), and the availability of the complete sequence of their genomes. Nonetheless, the ability to conduct systems level functional studies, and the possibility of relating the physiological programs to genetic programs that encode the underlying circuitry and its properties, the capacity of the system to defend itself, regenerate and repair, and the availability of reverse genetic tools to study gene expression and regulation, all strongly justify continuing and enhancing our efforts to understand neuroimmune responses in the leech.

ACKNOWLEDGMENTS

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INNATE IMMUNITY IN *C. ELEGANS*

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Abstract: The nematode *Caenorhabditis elegans* is proving to be a powerful invertebrate model to study host-pathogen interactions. In common with other invertebrates, *C. elegans* relies solely on its innate immune system to defend itself against pathogens. Studies of the nematode response to infection with various fungal and bacterial pathogens have revealed that the innate immune system of *C. elegans* employs evolutionary conserved signalling pathways. They regulate the expression of various effector molecules, some of which are also conserved. Here, we summarize the current knowledge of the pathways and effector molecules involved in the nematode immune response, with a particular focus on the antifungal immune response of the *C. elegans* epidermis.

INTRODUCTION

C. elegans is a free-living soil nematode that feeds on bacteria and is therefore constantly exposed to potential pathogens.¹ Like other invertebrates, *C. elegans* lacks an adaptive immune system. In contrast to many invertebrate species, however, *C. elegans* does not appear to have specialized immune cells. For example, while *Drosophila* has macrophage-like hemocytes, which engulf invading microbes, the only cells in the nematode body cavity, the 6 coelomocytes, do not seem to be capable of phagocytosis but function as scavenger cells with a high endocytic capacity.²

C. elegans possesses three major mechanisms of defences against microbial attacks:¹ Avoidance behaviour: It has been demonstrated that worms are able to distinguish between different bacteria. Whereas most bacteria attract *C. elegans*, some repel the nematode and cause an avoidance behaviour. Such an aversive response can be specifically directed against the pathogenic strains of a bacterial species (reviewed in ref. 3). Olfactory neurons, G protein coupled receptors and the only Toll-like receptor (TLR) in *C. elegans*, TOL-1,

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are involved in triggering the avoidance behaviour to pathogenic *Serratia marcescens*.^{4,5} Worms can “remember” odours⁶ and can even learn to avoid bacteria that are recognized as noxious.⁷ This discrimination relies in part on pairs of asymmetric chemosensory neurons.⁸ Their correct development requires a signalling cassette that includes an intracellular TIR-domain adapter protein (TIR-1) acting upstream of a p38 MAPK cascade.⁹ This cassette, which will be described in more detail below, appears also to play a direct behavioural role as it has been found to be involved in the neuroendocrine regulation of serotonin-dependent aversion to *Pseudomonas aeruginosa*.^{10,2} The second axis of protection against pathogen invasion is a strong cuticle, made of collagen and chitin and constituting the exoskeleton of the worm. It acts as a physical barrier that is relatively resistant to puncturing. As a complement, the pharyngeal grinder destroys pathogens that are taken up during feeding. It prevents live pathogens from reaching the intestine and establishing an infection. Indeed, mutants with defective grinder function are more susceptible to infection.^{11,12,3} The third line of defence involves inducible mechanisms. These will be the main focus of this chapter. *C. elegans* possesses a complex inducible defence system involving multiple signalling cascades that regulate the production of antimicrobial peptides (AMP) and proteins in a pathogen- and tissue-specific way.

ROUTES OF INFECTION

Most of the known pathogens of *C. elegans* use two main routes of infection, through the pharynx or the epidermis (Fig. 1). Many Gram-positive and Gram-negative bacteria as well as yeast, infect worms upon oral up-take during feeding and establish an intestinal infection. They must survive the passage through the grinder to reach the intestine, proliferate and establish an infection. In some cases, it has been shown that the pathogen destroys the grinder,¹³ in others it appears that the infectious particles, such as the spores of *Bacillus thuringiensis* are resistant to the mechanical action of the grinder.¹⁴ Almost all characterised intestinal pathogens of *C. elegans* remain extracellular, apart from *Salmonella typhimurium* and the microsporidium *Nematocida parisii*, that have been shown to establish intracellular infection in the intestinal cells.^{15,16}

Some pathogenic bacteria and fungi can adhere to the cuticle and infect the *C. elegans* epidermis. For example, *Microbacterium nematophilum* adheres to the anal region of the nematode and induces hindgut swelling¹⁷ and *Leucobacter chromiireducens* is capable of causing lethal uterine infections¹⁸ (Fig. 1). Different fungi that are pathogenic for nematodes, including *Drechmeria coniospora* and species of *Haptocillium*, produce spores that adhere and then penetrate the cuticle and grow into the epidermis (Figs. 1 and 2).^{19,20} Although some pathogens, such as certain strains of *P. aeruginosa*, produce fast-acting toxins,²¹ against which *C. elegans* appears defenceless, in many cases, infection provokes an immune response.

PATHOGEN RECOGNITION

The first step of an inducible defence is the recognition of the pathogen. Conserved structures on pathogens that are not present in the host and thus recognized as foreign, so called microbe-associated molecular patterns (MAMPs), bind to pattern recognition

receptors (PRRs) in many organisms.²² PRRs include peptidoglycan recognition proteins (PGRP), Gram negative binding proteins (GNBP), nucleotide-binding oligomerization domain (NOD) and NACHT domain proteins.²³ Genes encoding proteins of these families are absent from the *C. elegans* genome.

One prominent class of PRRs, in vertebrates the TLRs, can sense outer membrane components of the bacteria, RNA or DNA.²² As mentioned above, the single worm TLR, TOL-1, is involved in behavioural avoidance of some pathogenic bacteria,^{4,5} but does not seem to play a role in the resistance to several pathogens,⁵ nor in the regulation of certain immune effectors.²⁴ One study showed that *tol-1* mutants are more susceptible to *S. typhimurium* infection,²⁵ but it is unclear whether this is due to an involvement of *tol-1* in a protective immune response or rather due to a defect in cell adherence in the pharynx of the *tol-1* mutant leading to a defect in a physical barrier thus favouring pathogen invasion.

TLRs, as well as a number of other PRR families, in both plants and animals, share a common domain, the leucine rich repeat (LRR) domain. In a recent study, the role in host defences of each of the 14 predicted transmembrane proteins with LRR domains encoded in the *C. elegans* genome, was assayed. Loss-of-function mutants in one gene, *fshr-1*, which encodes a glycopeptide hormone receptor homologue, were found to be more susceptible to infection by Gram positive and Gram negative bacteria. It has yet to be determined if FSHR-1, which is expressed in the intestine, acts as a pathogen receptor or rather functions as a positive modulator of the nematode immune response.²⁶

C-type lectins are carbohydrate-binding proteins that can exhibit very narrow ligand specificity. In mammals, a number of C-type lectins have established roles in innate immunity. For example, Dectin-1 is highly expressed on macrophages and recognizes beta-glucan, a component of the fungal cell wall and thereby acts as a PRR.²³ *C. elegans* possesses 278 genes encoding C-type lectins, but it is currently unclear as to whether any of them function as PRRs or rather as effector molecules (see below).

While there is no clear Dectin-1 orthologue in *C. elegans*, there are a number of potential scavenger receptors (SR), another class of protein known to be involved in pathogen recognition in other species.²⁷ Indeed, there are six proteins homologous to CD36 and Croquemort, members of the SR-B family and one well-characterised SCARF orthologue CED-1. Because of its expression in the intestine throughout development, one of these, C03F11.3, was suggested a number of years ago to be potentially involved in the recognition of microbial molecules.²⁸ A study published last year supports such an idea, as CED-1/SCARF and C03F11.3/CD36 appear to function in host resistance to *Candida albicans* and *Cryptococcus neoformans* in *C. elegans*.²⁹ Whether in the nematode these proteins in fact recognize yeast cell wall beta-glucans and act as PRRs has not been formally demonstrated. Alternatively, given CED-1's known function in recognizing dying cells during programmed cell death, it might instead recognize damaged host material and then induce the unfolded protein response (UPR, see below) in an attempt to contain this damage. So we still do not know whether the worm responds through the detection of specific MAMPs or more generally to the cellular damage and stress caused by the pathogen (so-called danger theory³⁰) or both. Nevertheless, the finding that *C. elegans* shows distinct immune responses to different pathogens that infect via the same route and have similar levels of virulence,^{31,32} clearly supports a model of *C. elegans* specifically recognizing pathogens.

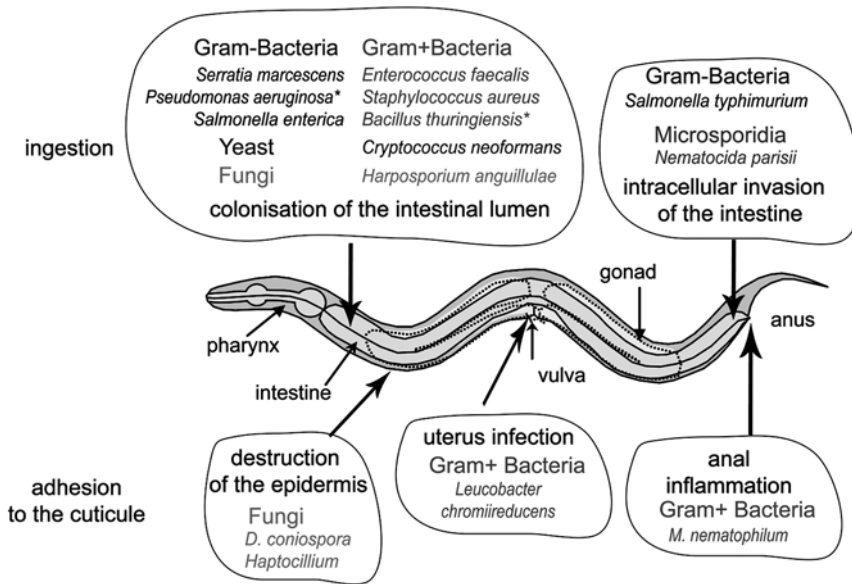


Figure 1. Pathogens of *C. elegans* and their route of infection. Most known pathogens of *C. elegans* are ingested and establish an infection in the intestinal lumen. Certain bacteria produce toxins (*) that can kill the nematode. The fungus *D. coniospora* and the bacteria *M. nematophilum* adhere to the cuticle and infect the nematode via the epidermis. Not all known pathogens of *C. elegans* are shown.

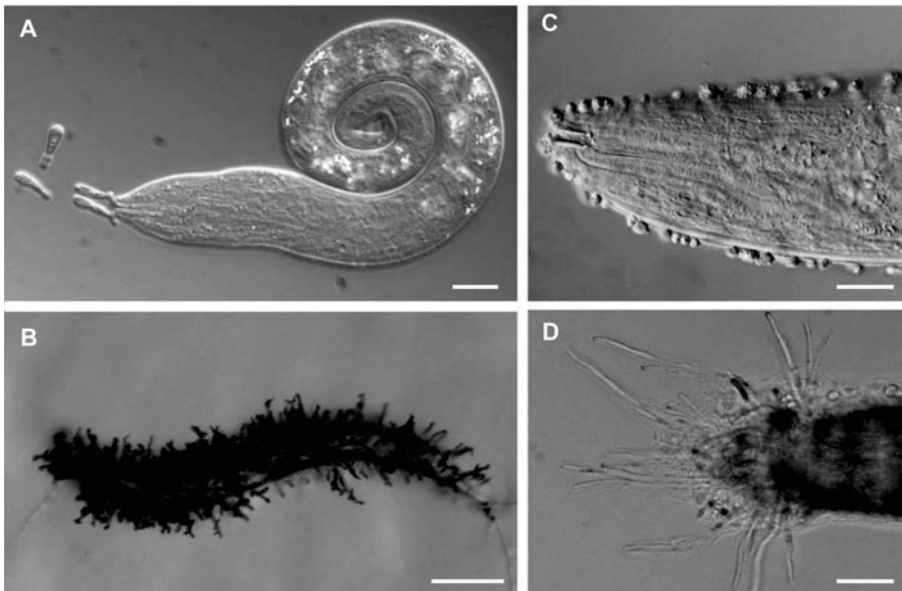


Figure 2. Fungal infection of *C. elegans*. (A and B) *D. coniospora*, (C and D) *Haptocillium*. (A and C) adhesion of the spores to the cuticle after few hours, (B and D) after 2 days fungal hyphae grow out of the worm. Scale bars are 10 μm (A), 100 μm (B) and 50 μm (C and D).

Table 1. Summary of the major signalling pathways in the *C. elegans* Immune System (updated from ref. 86)

Pathway	Tissue	Components	Homologues	References
p38 MAPK	Epidermis	GPA-12, RACK-1	G protein subunits	82
		EGL-8, PLC-3	Phospholipase C	82
		NIPI-3	Tribbles kinase	35
	Epidermis and intestine	TPA-1	Protein kinase C	82,87
		TIR-1, NSY-1, SEK-1, PMK-1	SARM MAP kinases	24,34,88 12,35
FSHR-1	Intestine	FSHR-1	G protein coupled receptor	26
ZIP-2	Intestine	ZIP-2	b-zip transcription factor	65
Insulin signalling	Nervous system	INS-7	Insulin-like peptide	76
	Intestine	DAF-2	Insulin receptor	42
		AGE-1	PI3 kinase	42
		AKT-1, AKT-2	Akt kinase	43
		DAF-16	FOXO transcription factor	42
TGF- β	Nervous system epidermis	DBL-1	TGF- β	54,55
		SMA-6	TGF- β receptor	55
		SMA-3	SMAD protein	55
Wnt/Hox	Intestine/ Hindgut	BAR-1	β -catenin	61
		EGL-5	Hox transcription factor	61,64
ERK MAPK	Hindgut	LIN-45, MEK-2, MPK-1	ERK MAP kinase	39
		EGL-8	Phospholipase C	89
		SUR-2	Mediator component	39
UPR ¹	Intestine	XBP-1	X box protein	50,52
		HSP-4	Heat shock protein	
Autophagy	Pharynx	CED-1, C03F11.3	Scavenger receptor	51
	Intestine	BEC-1, LGG-1	ATG proteins	16

¹The recent results of Richardson et al suggest that the primary function of the UPR is to protect against ER stress arising from the increase secretory response. Whether this is the case for Bt toxin⁵⁰ and for the noncanonical UPR⁵¹ remains to be seen.

SIGNALLING PATHWAYS INVOLVED IN THE IMMUNE RESPONSE

Even if the manner in which the immune response in *C. elegans* is initiated has not been fully elucidated, several signalling cascades have been described that are activated specifically by certain pathogens (Table 1) and lead to the production of effector molecules which have the potential to destroy pathogens.

Mitogen-Activated Protein Kinase (MAPK) Pathways

Mitogen-activated protein kinase (MAPK) pathways are considered to be the most ancient signal transduction cascades in immunity, found in both animals and plants. Three MAPK cascades are implicated in *C. elegans* immunity. A role for the p38 MAPK pathway in *C. elegans* defence was first revealed in a genetic screen for mutants hypersensitive to infection by *Pseudomonas aeruginosa*.¹² Since then, the p38 MAPK cascade has been shown to protect the worm against other Gram negative and positive bacteria and also fungi and seems to be one of the main signal transduction cascades in the worm's innate immune response.³³⁻³⁸

A second MAPK cascade implicated in *C. elegans* immunity is the extracellular signal-regulated kinase (ERK) pathway that is involved in the resistance of *C. elegans* to infection by the Gram positive bacterium *M. nematophilum*.³⁹ Thirdly, the MAPK kinase protein MEK-1 of the c-Jun N-terminal kinase (JNK) pathway is required for full activation of the p38 MAPK PMK-1, revealing an interaction between the different MAPK pathways.⁴⁰

DAF-2/Insulin-Like Receptor (ILR) Pathway

The DAF-2/insulin-like receptor (ILR) pathway, which involves the Foxo family transcription factor DAF-16, is also clearly important for the immune response of *C. elegans*, but its precise role is less clear. Active DAF-2 retains DAF-16/FOXO in the cytoplasm. In *daf-2* mutants, DAF-16 is predominantly in the nucleus. This results in an increase in DAF-16-dependent gene expression.⁴¹ DAF-2 is well known to be important for the control of lifespan. In addition to being long-lived, *daf-2* mutants show increased resistance to infection by several bacteria.⁴² Genetic evidence suggests, however, that the role of DAF-2 in immune signalling is distinct from its role in ageing. Downstream of DAF-2, four known serine threonine kinases, PDK-1, SGK-1, AKT-1 and AKT-2, regulate lifespan in at least 3 independent pathways. Mutants in these four kinases are long-lived, but only *akt-1* and *akt-2* mutants are more resistant to infection by *P. aeruginosa*.⁴³

It has been postulated that increased resistance of *daf-2* mutants may be linked to changes in expression for multiple antimicrobial genes.⁴⁴ But a direct comparison of the genes transcriptionally regulated by DAF-16/FOXO and the genes regulated after infection reveals a surprisingly limited overlap. Indeed most of the pathogen-induced immunity genes downstream of the PMK-1/p38 pathway are repressed by DAF-16/FOXO.^{45,46} Further, in contrast to what is seen upon exposure of *C. elegans* to several different abiotic stresses, nuclear translocation of DAF-16/FOXO has never been detected after infection. Additionally, it has been shown that DAF-16 transcriptionally regulates many genes involved in stress responses.^{44,47} It is therefore more probable that the DAF-2/DAF-16 pathway is part of a general stress response rather than a specific immune response. It should, however, be mentioned that the DAF-2/DAF-16 pathway controls multiple aspects of *C. elegans* physiology. Among other things, the DAF-2/DAF-16 pathway appears to influence the worm's pathogen avoidance behaviour, by an as yet undetermined mechanism.⁴⁸

The Unfolded Protein Response

In vertebrates, the endoplasmic reticulum (ER) unfolded protein response (UPR) is particularly important for the development and survival of highly secretory cells such

as plasma cells and exocrine gland acinar cells, which secrete immunoglobulins and digestive enzymes, respectively, as well as in dendritic cells and other antigen presenting cells.⁴⁹ In *C. elegans*, the UPR has been shown to be protective against *B. thuringiensis*. As detailed more fully below, it is activated by the poreforming toxins through the p38 MAPK pathway.⁵⁰ The UPR is also involved in the immune response to *S. typhimurium* and it appears that the scavenger receptor CED-1 is required for the activation of the UPR pathway.⁵¹

Very recently, the IRE-1-XBP-1 branch of the UPR was shown to be involved in defences against *P. aeruginosa*. Abrogation of *xbp-1* blocks part of the UPR and leads to a disruption of ER morphology. This has no major detrimental effect when worms are cultured under normal conditions, but if they are raised on *P. aeruginosa*, they are unable to complete their development and arrest as larvae. The developmental requirement for XBP-1 is bypassed in mutants of the p38/PMK-1 pathway, such that *xbp-1*; *pmk-1* double mutants can grow on *P. aeruginosa*. This led the authors to suggest that the production of antimicrobial proteins and peptides places a stress on the ER, which needs to be balanced by the activation of the UPR. In other words, the UPR may protect the host from the potentially damaging effect of its own innate immune against microbes.⁵²

TGF- β

A comparison of known targets of the developmentally important transforming growth factor β (TGF- β)/DBL-1 pathway⁵³ with those upregulated in adults upon infection with *S. marcescens* revealed a number of genes in common, including some encoding lectins and lysozymes.⁵⁴ More recently, TGF- β has been shown to be necessary for the regulation of AMP expression after a fungal infection (see below, ref. 55).

Autophagy, Apoptosis and Necrosis

A transcriptome analysis comparing the host genes affected by different bacterial infections revealed that among the genes induced by multiple bacteria were ones required for necrotic cell death. One might interpret this as indicating that necrosis could be a protective host defence mechanism. But when necrosis-defective mutants were tested, they were found to be more resistant to a bacterial infection than wild type.³² Similarly, a recent study has shown that a loss-of-function mutation in *ced-3* that encodes a caspase involved in apoptosis, also protects the worm against infection with *S. typhimurium*.¹⁶ This could be consistent with a deliberate triggering of necrotic cell death or apoptosis by pathogenic bacteria, as a strategy to increase their effective virulence.

Conversely, autophagy appears to be protective against the intracellular pathogen *S. typhimurium*. Thus *bec-1* or *lgg-1* mutants that are autophagy-defective show an increased susceptibility to infection, with an accumulation of *Salmonella* containing vacuoles (SCV) in the intestinal cell compared to wild type worms. Interestingly, these autophagy-defective mutants suppress the enhanced resistance to *S. typhimurium* infection of *daf-2/Insulin receptor* mutants and of a strain overexpressing DAF-16/FOXO,¹⁶ while at the same time increasing normal life span.⁵⁶ This suggests that increased intestinal epithelial cell autophagic activity may partially underlie the resistance of *daf-2* mutants to intracellular pathogens.¹⁶ It will be interesting to establish whether intestinal cell autophagy also contributes to the resistance of *C. elegans* to extracellular pathogens.

TRANSCRIPTION FACTORS INVOLVED IN THE IMMUNE RESPONSE

The transcription factor NF- κ B links the reception and transmission of an infection signal to the expression of effector proteins in vertebrates and insects. Therefore, its absence from the nematode genome is remarkable and opens the possibility of studying alternative mechanisms of transcriptional regulation potentially conserved in other species. For example, having shown that most of the effectors induced by *P. aeruginosa* infection in the intestine of *C. elegans* are under the control of the GATA transcription factor ELT-2, Tan and colleagues were able to show that ELT-2 increases host resistance to intestinal infection with bacterial⁴⁵ and this was subsequently also shown to be the case for intestinal fungal pathogens.⁵⁷ Another GATA transcription factor ELT-3 contributes to the proper expression in the epidermis of AMP genes. But it was also shown to be required for the expression in the epidermis of genes important for osmoregulation, not directly related to innate immunity. This led to the suggestion that this GATA TF acts as a more generic transcription factor in the epidermis.⁵⁸ This latter conclusion is in line with a study published this year showing that ELT-3 in the epidermis, but also ELT-2 in the intestine, are essential for tissue-specific activation of osmosensitive gene expression and promote survival under osmotically stressful conditions.⁵⁹

Just as the response to infection and osmotic adaptation may be controlled via regulation of common tissue-specific GATA transcription factors, so too is there a link between innate immunity and temperature adaptation. A mild heat shock has been shown to increase the resistance of *C. elegans* to infection with Gram positive and Gram negative bacteria. This resistance is independent of the p38 MAPK/PMK-1 pathway and requires the heat shock factor HSF-1 and heat shock proteins. The forkhead transcription factor DAF-16 is positively regulated by heat shock and is required for the induction of HSF-1 thus linking the heat shock pathway to the DAF-2/ILR pathway.⁶⁰

The transcriptional cofactor BAR-1/ β -catenin and the homeobox gene *egl-5* have been shown to play a role in *C. elegans* intestinal epithelial immunity and resistance to *S. aureus*,⁶¹ in addition to its established role in cell fate decision during development.⁶² EGL-5 is also necessary in the hindgut to induce swelling upon *M. nematophilum* infection.^{63,64} Interestingly, the human homologues of EGL-5, HOXA9 and HOXA10 dampen NF κ B-dependent TLR2 signalling, suggesting a conserved role in innate immune defence.⁶¹

Further insights into the complexity of innate immune signalling were obtained in a study that used a gene specifically induced by virulent *P. aeruginosa* strains called “infection response gene 1” (*irg-1*). It was chosen as its expression is independent of the PMK-1/p38 pathway. Several candidates required for the full induction of *irg-1* were identified from a screen of more than 300 transcription factors, based on RNA interference. Among them, most interest was focused on the bZIP transcription factor *zip-2*. It was shown to be required for the induction not only of *irg-1* but of several putative effector genes, in all cases independent of the PMK-1/p38 pathway and also of FSHR-1. Certain target genes, such as *irg-3*, were demonstrated to be regulated by yet another pathway, involving neither *zip-2*, nor p38, nor FSHR-1, suggesting that at least 4 independent pathways contribute to pathogen resistance in the *C. elegans* intestine upon *P. aeruginosa* infection.⁶⁵

EFFECTOR MOLECULES INVOLVED IN THE IMMUNE RESPONSE

Antimicrobial Peptides

C. elegans possesses different types of antimicrobial proteins and several classes of AMP. Among them are the mollusc defensin/mycitin-like peptides (ABF-1 to ABF-6). One of them, ABF-2 has demonstrated antimicrobial activity.⁶⁶ It is strongly upregulated upon prolonged exposure to *S. typhimurium*.⁶⁷ Additionally, there are the neuropeptide-like proteins (NLPs) and the caenacins (CNC),^{24,55,58} that are rapidly and strongly induced by fungal infection and that will be discussed in detail below.

Caenopores

Caenopores is the name given to a number of *C. elegans* proteins that contain the saposin domain, common to mammalian NK-lysin and granulysin and the protozoan amoebapores.⁶⁸ Members of this family were first identified more than a decade ago, when two among them, SPP-1 and SPP-5, were shown to have a bactericidal function.⁶⁹ SPP-5 is constitutively expressed and kills bacteria by permeabilising their membrane. Interestingly, another member of this family, SPP-3, is expressed both upon starvation and contact with certain bacteria, thus suggesting a potential link between nutrition and immunity.⁶⁸

Lysozymes

Lysozymes are another class of molecules known to be involved in immune defence in many species. In contrast to arthropods, *C. elegans* does not have C-type lysozymes, but possesses a repertoire of 15 genes, falling into 3 classes, two related to protist lysozymes and one specific to invertebrates.⁷⁰ Certain lysozymes, including *lys-7*, are induced upon bacterial challenge and their inactivation has been shown to render worms more susceptible to *M. nematophilum* and *P. aeruginosa*.^{54,71,72} The expression of other lysozymes, mainly from the invertebrate class, has been reported to be repressed upon infection. Although the exact function of these latter genes still remains to be determined, the amplification of the lysozyme family by gene duplication in the nematode is a clear example of evolutionary function diversification.⁷⁰

Lectins

Lectins are also involved in innate defences in many species and can be involved in pathogen recognition but also in immune effector functions. In *C. elegans*, there are a very large number of lectin genes, including 11 galectins, in the *lec* gene class and 265 C-type lectins in the *clec* gene class. The expression of some *lec* and *clec* genes is up-regulated by several pathogens; for others their induction appears to be relatively pathogen-specific. This differential upregulation has led to the suggestion that they might be an element conferring specificity to the immune response of *C. elegans*.^{31,32,54,72} In some cases, they have demonstrable role in host defence. Inactivation of some lectins, for example, renders worms more susceptible to *M. nematophilum*.⁷² Unfortunately, there is currently little

direct functional information about most of the large number of lectins. One exception is the glycolipid-binding galectin LEC-8 that has been shown recently to play a role in host defence against *B. thuringiensis* infection by competitively inhibiting the binding of the toxin Cry5B to its host glycolipid receptor.⁷³

Reactive Oxygen Species

In addition to its arsenal of antimicrobial proteins, *C. elegans* also has the capacity to produce bactericidal reactive oxygen species (ROS) in response to exposure to pathogens. This has been best characterised in the case of infection with the Gram-positive pathogen *Enterococcus faecalis*, which provokes ROS production via the action of the dual oxidase BLI-3. ROS are relatively unspecific in their capacity to kill cells. They affect invading micro-organisms, but can also damage host tissues. As a result, increased levels of ROS triggers a protective stress response in the host. This involves up-regulation of the superoxide dismutase SOD-3 and the catalase CTL-2, which sequentially detoxify the ROS. Both enzymes are targets of DAF-16. Indeed, their combined action is part of the mechanism underlying the increased resistance to infection of *daf-2* mutants. Consistent with a protective role for oxidative stress, the addition of compounds that scavenge ROS increase the sensitivity of *C. elegans* to infection with *E. faecalis*.^{74,75}

MODULATION OF THE IMMUNE RESPONSE BY THE NERVOUS SYSTEM

ROS can act in a relatively unspecific manner over a distance to affect cells not in direct contact with a pathogen. Several recent papers attempt to provide evidence for more refined noncell autonomous mechanisms involved in controlling the response of *C. elegans* to infection. These examples involve the nervous system. Kawli and Tan demonstrated that the release of dense core vesicles (DCVs) from neurons suppresses the intestinal immune response of *C. elegans* to *P. aeruginosa* and that this neuronal control mechanism is mediated in the intestine by the DAF-2/ILR pathway. The insulin-like peptide INS-7 has been proposed to provide the link between DCV release in neurons and the DAF-2/ILR pathway in the intestine.⁷⁶ In a second study, Stryer et al reported that NPR-1, a G-protein-coupled receptor related to mammalian neuropeptide Y receptors, functions to suppress innate immunity to *P. aeruginosa*, by acting upstream of the p38 MAPK signalling cascade.⁷⁷ It should be noted, however, that the reported changes in gene expression seen in the *npr-1* mutant are minimal compared to those seen upon infection by *P. aeruginosa*. Further, the results of Stryer et al have been contradicted by a more recent study showing that the difference in susceptibility in the *npr-1* mutant strain is due to its well-characterised behaviour of clumping, a behaviour which is linked to sensing oxygen concentration.⁷⁸ In a final example, as mentioned above and described more fully below, the expression of *cnc* AMP genes appears to involve regulation by neuronally-derived TGF- β .⁵⁵

IMMUNE RESPONSE TO PORE-FORMING TOXINS

Some bacteria, such as *B. thuringiensis*, are able to produce multiple toxins that target host cells. These pore-forming toxins (PFTs) make holes in membranes and alone can

cause the death of *C. elegans*. Not surprisingly, the nematode has evolved mechanisms to protect itself from the nefarious effects of PFTs. It has been demonstrated that the toxin directly binds glycolipids and that the major mechanism for PFT resistance in *C. elegans* entails a loss of glycolipid carbohydrates.⁷⁹ Moreover, in the case of the *B. thuringiensis* Cry5B toxin, this response involves the PMK-1/p38 and cJunN-terminal kinase-like pathways. The mechanism has an inbuilt component of amplification, since these kinases are also transcriptionally upregulated by Cry5B.³⁸ Activation of the p38 MAPK pathway by Cry5B activates the IRE-1 UPR pathway. IRE-1 induces an alternative splicing of the transcription factor *xbp-1*. The resultant infection-specific transcript then drives the expression of a number of target genes which protect against the effects of the PFT.⁵⁰ There is some cross-talk between the mechanisms involved in the response to PFTs and those required to tolerate conditions of low oxygen. Indeed, activation of the hypoxia pathway also increases resistance against PFTs. Resistance to hypoxia also involves the UPR and in common with the response to PFTs, it is mediated by the transcription factor HIF-1.⁸⁰

EPIDERMAL IMMUNE RESPONSE TO THE FUNGUS *DRECHMERIA CONIOSPORA*

Most of the bacterial or fungal pathogens described in the previous sections infect the worm through the intestinal lumen, which is primarily programmed for destroying microbes as part of normal feeding and digestion. In some instances, this can blur the distinction between an immune response and the consequence of a change of diet. Other pathogens infect worms via the cuticle. Among them, *Drechmeria coniospora* is a natural fungal pathogen of nematodes, including *C. elegans* (M.A Felix, personal communication). *D. coniospora* pierces the worm's cuticle and its hyphae proliferate first in the epidermis, then throughout the organism. This provokes a complex transcriptional response involving, among others, the upregulation of AMP genes.^{24,58} These include members of two phylogenetically-related families, the *nlp* and *cnc* genes, which are present in clusters in the genome. Phylogenetic analysis shows that these AMP genes, arose through recent duplication and diversification, have been under selective pressure during evolution and are thus likely to be important in nature for the survival of *C. elegans*.⁵⁸

As this fungal infection involves breaching the cuticle and epidermis, the question of whether *C. elegans* epidermis responds to physical injury was addressed. Needle pricking or laser wounding not only provokes an up-regulation of AMP genes, but also triggers a cellular wound-healing and scarring mechanism.³⁵ These two processes appears to be independent, but are normally kept in check by a common negative regulator, the nematode Death-associated protein kinase (DAPK).⁸¹ While infection and injury induces the expression of both *nlp* and *cnc* family genes, the molecular mechanisms that regulate each class appear to be strikingly different.

Cell-Autonomous Regulation of *nlp* Gene Expression

Through direct genetic screens, proteomics and a candidate gene approach, two signalling pathways required for the regulation of *nlp* gene expression in the epidermis have been described. One is specific for infection, the second is also activated by wounding. Both pathways converge on a protein kinase C, TPA-1, which, in turn, acts

upstream of the PMK-1/p38 pathway (Fig. 3). The proximal elements of the pathways are, however, distinct. The former requires the conserved protein kinase Tribbles, NIPI-3, while the latter involves heterotrimeric G proteins acting upstream of a phospholipase C. All of the characterised components act in a cell-autonomous manner to control *nlp* gene expression in the epidermis.^{35,82}

For the time being, the identity of the putative G-protein coupled receptor (GPCR) that activates the heterotrimeric G proteins is unknown, nor is it known how NIPI-3 is activated. There are, nonetheless, marked similarities between the molecular architecture underlying these pathways and the organisation of the signalling pathways that regulate the innate immune response both in *Drosophila* and in vertebrates. These led to the speculation that the innate immune response to *D. coniospora* arose from a GPCR-dependent mechanism

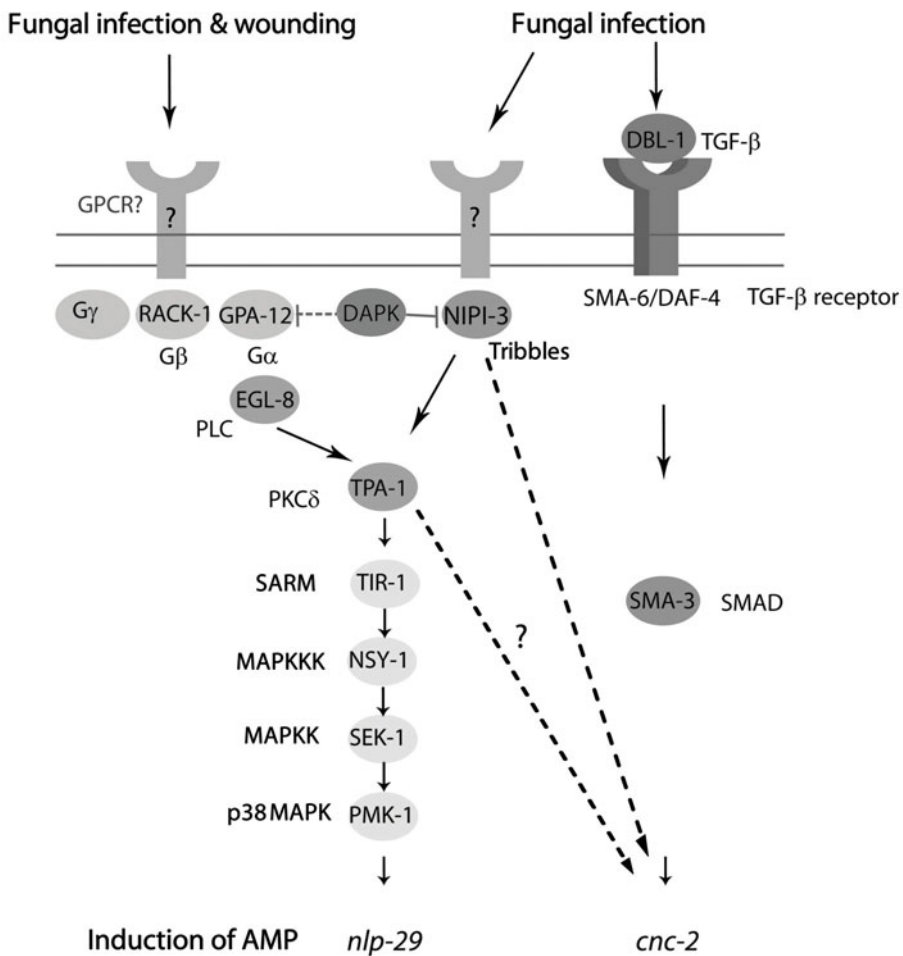


Figure 3. Schematic representation of the different signalling pathways and their components involved in the induction of antimicrobial peptides expression upon *D. coniospora* infection. Expression of the *nlp* genes is controlled by a PKC/SARM/p38 MAPK pathway and expression of *cnc* genes is controlled by a TGF- β pathway.

used to detect cellular damage, which was subsequently ameliorated by the addition of a pathogen-specific detection mechanism.

Paracrine Regulation of *cnc* Gene Expression

Surprisingly, the *cnc* gene family is regulated in an entirely different fashion, as it is largely independent of the PMK-1/p38 pathway. Induction of the *cnc* genes after infection requires the ligand DBL-1/TGF- β , produced by certain neuronal cells, which acts via its normal receptor SMA-6/DAF-4 expressed on epidermal cells (Fig. 3). The resultant signal is transduced by a noncanonical TGF- β pathway that does not involve all three SMAD proteins, hitherto considered to be indispensable for TGF- β signalling in *C. elegans*. Artificially altering the level of expression of the TGF- β modulated the strength of *cnc* induction after infection but did not affect the basal level of *cnc* gene expression, suggesting that infection triggers the conversion of an inactive precursor into an active TGF- β . This is reminiscent of the proteolytic activation of Spaetzle required for triggering the Toll pathway during the immune response in *Drosophila*. As the canonical TGF- β pathway undoubtedly existed before the appearance of the *cnc* genes, this is another clear example of the co-option and adaptation of a pre-existing signalling pathway for use by the innate immune system.⁵⁵

CONCLUSION

In the last 10 years, our knowledge of the way *C. elegans* defends itself against microbes has greatly expanded. The worm clearly shows specific immune responses to pathogens and their toxins. The signalling processes and effector molecules involved in this response have been elucidated to some extent. Interestingly, the main signalling pathways involved in the *C. elegans* immune response are conserved in other species, in part because they also play important developmental roles.

On the other hand, much remains to be learnt about the way the nematode recognizes pathogens and the receptors involved in this process, as well as the temporal and spatial dynamics of the downstream signalling processes. For the latter, techniques to visualise proteins and transcripts at the single molecule level within living cells must be improved.

It is important to note how deeply the defence mechanisms are embedded in the physiology of the organism. We described how important are several aspects of behaviour, digestion or stress resistance to defence. Other studies have shown connections with reproduction, where sterile mutants are more resistant to bacterial infection in a DAF-16/FOXO dependent manner.⁸³ The same is true for osmotic stress since several osmotic mutants are more resistant to fungal infection and some immune effectors genes are induced upon osmotic stress although apparently regulated by a dedicated pathway.^{58,84} Lastly, lipid synthesis has been shown to be required for the basal activity of the PMK-1/p38 pathway that influences resistance against *P. aeruginosa*.⁷¹ Unbiased forward genetics screens combined with global functional genomic approaches will help to unravel the complex and intricate biology that underlies successful host defences in *C. elegans*.

Another powerful approach to understand more fully how the innate immune system works is to take advantage of the fact that pathogens can specifically interfere with the defence mechanisms of the host. For the moment there are only a few examples in

C. elegans, such as the down-regulation of several intestinal immune effectors through activation of DAF-2 by *P. aeruginosa* which requires bacterial virulence factors controlled by the signalling molecule GacA.⁸⁵ The study of these interactions will also teach us more about innate immune defence and the virulence strategies that pathogens have developed to escape host immunity.

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IMMUNOCOMPETENT MOLECULES AND THEIR RESPONSE NETWORK IN HORSESHOE CRABS

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Abstract: Horseshoe crab hemocyte selectively responds to bacterial lipopolysaccharides (LPS), which depends critically on the proteolytic activity of the LPS-responsive serine protease zymogen factor C. In response to stimulation by LPS, the hemocyte secretes several kinds of immunocompetent proteins. The coagulation cascade triggered by LPS or β -1,3-D-glucans (BDG) results in the formation of coagulin fibrils that are subsequently stabilized by transglutaminase (TGase)-dependent cross-linking. Invading pathogens are recognized and agglutinated by lectins and then killed by antimicrobial peptides. Moreover, LPS-triggered hemocyte exocytosis is enhanced by a feedback mechanism in which the antimicrobial peptides serve as endogenous mediators. Factor C also acts as an LPS-sensitive complement C3 convertase. In addition, a sub-cuticular epidermis-derived protein forms a TGase-stabilized mesh at sites of injury. Horseshoe crabs have a sophisticated innate immune response network that coordinately effects pathogen recognition and killing, prophenoloxidase activation, complement activation and TGase-dependent wound healing.

INTRODUCTION AND HISTORICAL BACKGROUND

Extant horseshoe crabs comprise four species—*Limulus polyphemus*, *Tachypleus tridentatus*, *T. gigas* and *Carcinoscorpius rotundicauda*—each having a distinct geographic distribution. *L. polyphemus* is distributed along the east coast of North America and the other three species are distributed throughout Southeast Asia. In Japan, *T. tridentatus* inhabits coastal areas of the northern part of Kyushu Island as well as the Inland Sea. *T. tridentatus* is relatively long-lived; the embryo molts four

times within the fertilized eggs and after hatching it molts every year for 15 years to become a mature adult.¹

The first report of hemolymph coagulation involved *L. polyphemus* and was made by Howell in 1885.² The early studies on hemolymph coagulation and granular hemocytes were performed by Loeb and granular hemocytes were named as amebocytes after their amoeboid movement.^{3,4} In 1903, Hideyo Noguchi—who later gained fame for his studies on snake venoms and syphilology—injected horse red blood cells into *L. polyphemus* and reported strong hemagglutinating activity in hemolymph plasma.² In 1956, Bang published a landmark study on hemolymph coagulation and reported that intravascular clotting is induced by an injection of a ubiquitous marine pathogen, *Vibrio*.⁵ In 1964, Levin and Bang made the important observation that coagulation occurs when exposed to endotoxin (LPS).^{6,7} Moreover, they found that procoagulant activity and a clottable protein are located in granular hemocytes, not in hemolymph plasma.⁸ They immediately recognized the high sensitivity of LPS-mediated hemolymph coagulation and its applicability to hemocyte lysate, *Limulus* amoebocyte lysate (LAL), for assaying LPS.

In 1972, evidence for the proteolytic mediation of hemolymph coagulation was presented; LAL was fractionated, with one fraction containing the clottable protein and the other containing a coagulation accelerator with properties of proteolytic enzyme properties.⁹ The clottable protein was designated coagulogen by virtue of its functional similarity to mammalian fibrinogen.¹⁰ In 1977, Tai et al reported the mechanism underlying proteolytic conversion of coagulogen by the proclotting enzyme and they presented preliminary evidence for the activity of TGase in LAL as an unpublished observation.¹¹ Tai and Liu further demonstrated that the activation of the proclotting enzyme depends on the presence of Ca²⁺ and LPS and that the proclotting enzyme contains γ -carboxyglutamic acids, as does bovine prothrombin.¹² Later, Muta et al clearly indicated that the proclotting enzyme requires no Ca²⁺ in the proteolytic cascade and that it contains no γ -carboxyglutamic acid in the sequence.¹³ Iwanaga et al developed chromogenic or fluorogenic substrates for proteases involved in hemolymph, resulting in the first discovery of a proteolytic cascade mediated by LPS and BDG.¹⁴⁻¹⁶

In 1985, Copeland and Levin examined the morphology and the fine structure of the granular hemocyte of *L. polyphemus*, showing that the hemocyte contains two types of granules: large ones and dense ones.¹⁷ The immunocytochemical analysis of hemocytes showed that the large granules contain coagulogen and factor C, whereas the dense (or small) granules contain only an antimicrobial peptide, tachyplesin.¹⁸ Moreover, the two types of granules were separated by ultracentrifugation and their protein components were purified by reverse-phase high-performance liquid chromatography.¹⁹

Here we review the current knowledge of horseshoe crab innate immunity at the molecular level with an emphasis on the importance of immunocompetent proteins derived from hemocytes, hemolymph plasma and sub-cuticular epidermis.

THE MOLECULAR MECHANISM UNDERLYING LPS-TRIGGERED HEMOCYTE EXOCYTOSIS

In horseshoe crabs, granular hemocytes constitute 99% of all hemocytes.¹⁸ The hemocyte responds selectively to LPS but not to other pathogen-associated molecular patterns (PAMPs), such as BDG and peptidoglycans.²⁰ A variety of defense molecules are stored in the secretory granules of the hemocyte; large granules contain serine protease zymogens including factor C, factor G, factor B and the proclotting enzyme, the clottable protein

coagulogen, serine protease inhibitors (serpins), lectins and substrates for TGase, whereas dense granules contain antimicrobial peptides.²¹ In response to stimulation by LPS, these immunocompetent molecules are rapidly secreted by the hemocyte. Factor C is a unique LPS-responsive serine protease zymogen. A part of factor C is located on the hemocyte surface and acts as an LPS sensor to potentiate hemocyte exocytosis, which is localized in a punctate distribution on the hemocyte surface.^{22,23} Factor C interacts with acidic phospholipids, and it also shows tight interaction with cholesterol.²⁰ The interaction between factor C and LPS is competitively inhibited by the addition of acidic phospholipids.²⁰ In contrast, cholesterol does not inhibit this interaction, suggesting that factor C interacts with cholesterol through a binding site that is distinct from that for LPS and raising the possibility that factor C may be localized on cholesterol-rich microdomains or lipid rafts on the hemocyte membrane.²⁰ Upon activation by LPS, activated factor C initiates hemocyte exocytosis via a G-protein-dependent exocytic pathway that is also dependent on the proteolytic activity of activated factor C. In this respect, LPS-triggered hemocyte exocytosis is analogous to the thrombin-thrombin receptor (the protease-activated G protein-coupled receptor, PAR) signaling axis in mammalian platelets. Hemocyte exocytosis requires 50 mM Mg²⁺ and 10 mM Ca²⁺ ions, equivalent to the concentrations of these cations in hemolymph plasma. Exclusion of these cations from the assay buffer inhibits hemocyte exocytosis even at high concentrations of LPS. Moreover, in the absence of LPS, hemocyte exocytosis can be triggered by synthetic hexapeptides corresponding to the tethered ligands of mammalian PARs, supporting the notion of a PAR-like receptor on the hemocyte surface (Fig. 1).

Factor C is a mosaic protein that contains a Cys-rich region, an epidermal growth factor-like domain, five Sushi domains (also designated as complement control protein modules or short consensus repeats) and a C-type lectin domain, in addition to a typical serine protease domain at the COOH-terminus²¹. Factor C interacts with LPS via the N-terminal Cys-rich domain ($K_d = 7.6 \times 10^{-10}$ M).^{20,23} This contrasts with the findings of studies of the factor C ortholog from *C. rotundicauda*, which defined a region with the tandem sushi domains as important for LPS binding, using the truncated recombinant fragments, namely, sushi 123, sushi 1 and sushi 3 domains or their synthetic peptides.²⁴ However, these recombinant fragments or the peptides may not be properly folded by mixed disulfide bridges. The intermolecular disulfide bonding of the sushi 3-derived synthetic peptide, resulting in an unphysiological dimer, is practically indispensable for its interaction with LPS.²⁵

ENDOGENOUS AMPLIFICATION SYSTEM FOR HEMOCYTE EXOCYTOSIS

The binding parameter between factor C and LPS could not explain the extraordinarily high sensitivity of horseshoe crab hemolymph for LPS as low as $\sim 10^{-13}$ g/ml ($\sim 10^{-14}$ M). We found that the hemocyte has an endogenous feedback mechanism for LPS-triggered hemocyte exocytosis.²⁶ The hemolymph normally contains hemocytes at $\sim 10^6$ cells/ml. LPS-triggered hemocyte exocytosis is highly dependent on cell density, namely, an increase in cell density (cells/ml) from 0.05×10^6 to 0.8×10^6 results in a 10^6 -fold change in the apparent LPS sensitivity (from 10^{-7} to 10^{-13} g/ml of LPS). Interestingly, an antimicrobial tachyplesin in the exocytosed fluid acts as a secondary secretagogue, thereby dramatically enhancing the sensitivity of the hemocyte to LPS. Tachyplesin has structural properties in common with mastoparan, a basic tetradecapeptide from wasp venom. Mastoparan interacts directly with G proteins without direct stimulation of the upstream receptor and

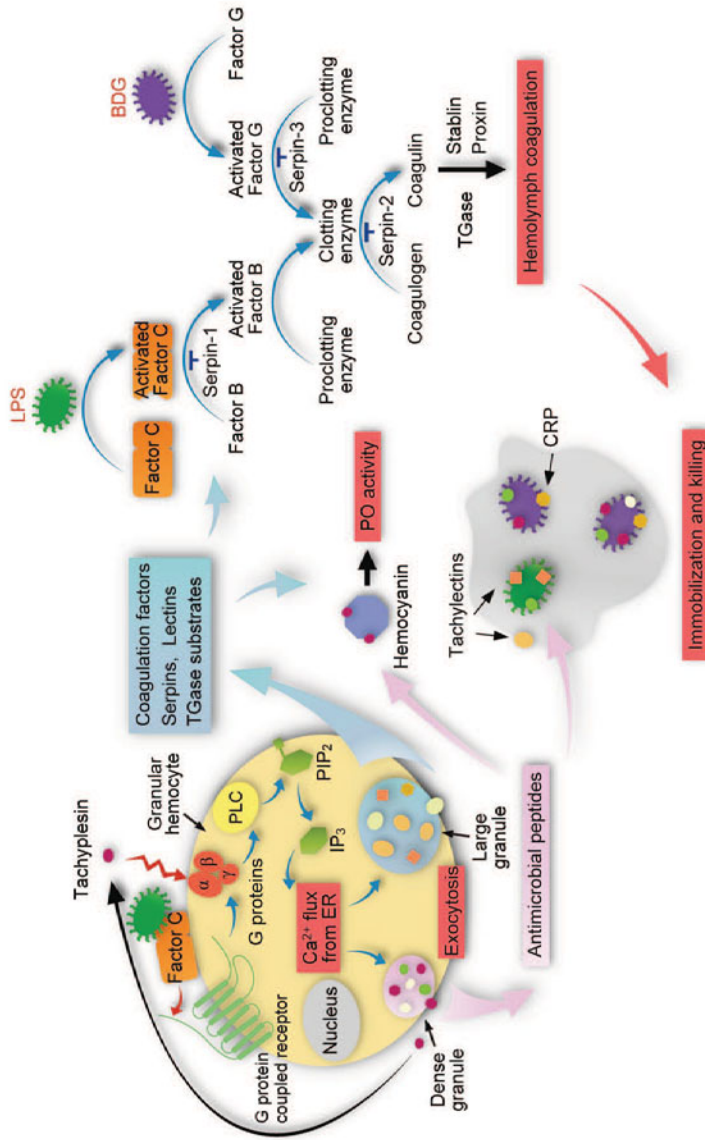


Figure 1. LPS-triggered hemocyte exocytosis and the resulting innate immune responses. In response to stimulation by LPS, granular components are rapidly secreted by the hemocyte via G protein-dependent exocytic pathway. The hemocyte stores a variety of defense molecules including coagulation factors, lectins, antimicrobial peptides and transglutaminase substrates. A serine protease zymogen, factor C, functions as an LPS sensor on the hemocyte surface and an initiator of hemolymph coagulation. Alternatively, activated factor G in the presence of β -1,3-D-glucans also triggers the coagulation cascade. Factor C and factor G independently serve to couple the recognition of pathogen-associated molecular patterns to form a physical barrier at the site of microbial invasion. LPS-induced hemocyte exocytosis leads not only to coagulation but also activates a sophisticated innate immune response network that culminates in immobilization and killing of pathogens and conversion of hemocyanin to phenoloxidase. PLC, phospholipase C; PIP₂, phosphatidylinositol-1,4,5-triphosphate; IP₃, inositol-1,4,5-triphosphate; PO, phenoloxidase.

induces exocytosis in the mast cell.²⁷ Consistent with these findings, mastoparan is able to induce hemocyte exocytosis in *T. tridentatus*.²⁰ Moreover, because tachyplesin binds to bovine G protein ($K_d = 8.8 \times 10^{-7}$ M), tachyplesin may interact with G-proteins in the hemocyte in a manner similar to that of mastoparan.²⁶ In addition, tachyplesin binds to hemocyanin ($K_d = 3.4 \times 10^{-6}$ M), the major protein in the hemolymph plasma; this suggests that hemocyanin may serve as a sink for tachyplesin, thereby spatially restricting its hemocyte-stimulating effect to the site of infection (Fig. 1).²⁸

FUNCTION OF A TOLL-LIKE RECEPTOR ON THE HEMOCYTE

Insects and mammals conserve a signaling pathway of the innate immune system through cell-surface receptors called Tolls and Toll-like receptors.^{29,30} The toll-like receptor of *T. tridentatus* is closely related to the *Drosophila* Toll in both domain architecture and overall length.^{31,32} Horseshoe crab Toll is nonspecifically expressed in all tissues examined, suggesting that it does not act as an LPS receptor on granular hemocytes.³² On the other hand, NF- κ B and I κ B homologues (CrNF- κ B and CrI κ B) have been identified in *C. rotundicauda*.³³ Gram-negative bacteria infection causes degradation of CrI κ B and nuclear translocation of CrNF- κ B, leading to upregulation of immune-related gene expression, including nitric oxide synthase and factor C, indicating that the NF- κ B/I κ B signaling cascade remains well conserved from horseshoe crabs to mammals. Although they must play a fundamental role in regulating the expression of critical immune defense molecules, further investigations are essential to reveal the physiological function of horseshoe crab Toll and its signaling.

THE COAGULATION CASCADE TRIGGERED BY LPS AND BDG

Factor C is autocatalytically activated in the presence of LPS and the resulting activated factor C activates coagulation factor B, which in turn converts the proclotting enzyme into the clotting enzyme. The clotting enzyme then promotes the proteolytic conversion of coagulogen to coagulin, which spontaneously forms an insoluble polymer (Fig. 1). Alternatively, activated factor G in the presence of BDG triggers the activation of the proclotting enzyme to the clotting enzyme. The mammalian coagulation system acts locally on the phospholipid surface in cooperation with Ca^{2+} ions at the site of vascular injury. In an analogous fashion, the horseshoe crab coagulation system is restricted to the surfaces of invading pathogens, such as Gram-negative bacteria and fungi. However, the coagulation cascade of horseshoe crabs does not have a common evolutionary origin with that of mammals. In fact, fibrinogen homologues of the horseshoe crab, tachylectins-5A and -5B, act as nonself-recognizing proteins rather than as target proteins of the coagulation cascade.³⁴ Also, coagulogen has no structural similarity or evolutionary relatedness to fibrinogen.³⁵

A protease cascade in *Drosophila* has been well characterized as the morphogenetic cascade for determining embryonic dorsal-ventral polarity, leading to the production of the Toll ligand spätzle.³⁶ The *Drosophila* Toll pathway also controls resistance to fungal and Gram-positive bacteria infections.³⁷ Spätzle belongs to the nerve growth factor family and is homologous to coagulogen.^{38,39} In addition, a clip-like domain (originally identified in the proclotting enzyme as a disulfide-knotted domain), located in the N-terminal region of coagulation factor B and the proclotting enzyme, has been identified in the proteins

snake and easter of the *Drosophila* Toll pathway.^{13,40} Therefore, the two functionally distinct cascades may have a common evolutionary origin.^{41,42}

The coagulation cascade is regulated by three types of serpins that form stable 1:1 covalent complexes with target coagulation proteases: serpins-1, -2 and -3 inhibit activated factor C, the clotting enzyme and activated factor G, respectively.⁴³⁻⁴⁵ These serpins are more closely related to mammalian serpins than they are to insect serpins: serpin-1 shows higher sequence identities to human plasminogen activator inhibitor (40%) than to an elastase inhibitor from *Manduca sexta* (29%) or silkworm antichymotrypsin (27%).⁴³ All three serpins are secreted upon hemocyte exocytosis in response to stimulation by LPS. These serpins appear to prevent diffusion of the activated forms of coagulation factors that escape into the hemolymph from the surfaces of microbes at the site of injury and thereby prevent unnecessary clot formation.

BDG RECOGNITION BY FACTOR G

In crustaceans and insects, the recognition of BDG triggers a serine protease cascade, leading to the activation of prophenoloxidase, a key enzyme in the melanization of pathogens and damaged tissues.^{46,47} In mammals, the recognition of BDG by dectin-1, a C-type lectin family member, potentiates the production of cytokines and antifungal reactive oxygen species by dendritic cells and macrophages.⁴⁸ Factor G is another pattern-recognition protein in the coagulation cascade that acts as a sensitive sensor for BDG. Factor G is a heterodimeric serine protease zymogen composed of two noncovalently associated subunits, α and β .⁴⁹ The β subunit contains a serine protease domain, whereas the α subunit acts as a pattern-recognition subunit: it comprises three types of noncatalytic glycosidase-like modules, including a single β -1,3-D-glucanase A1-like module, three tandem xylanase A-like modules and two tandem xylanase Z-like modules, Z1 and Z2. Of these three types, the C-terminal Z1 and Z2 modules act as independent binding sites for BDG.⁵⁰ This observation, taken together with the high degree of sequence identity between Z1 and Z2 (91%), suggests that duplicated binding sites for BDG may increase avidity to allow stable and specific recognition of fungi.

The Z1 or Z2 module shows the highest sequence similarity to a carbohydrate-binding module of endoglucanase 5A from the aerobic soil bacterium *Cellvibrio mixtus* (45% sequence identity). Endoglucanase 5A contains an N-terminal catalytic domain and two tandem repeats of noncatalytic family 6 carbohydrate-binding modules, *CmCBM6-1* and *CmCBM6-2*.⁵¹ Our recent structural studies of the recombinant Z2 module by NMR spectroscopy clearly indicate that the ligand-binding site in the Z2 module is located in a cleft on a β -sheet in a predicted β -sandwich structure, which is superimposed onto cleft B in *CmCBM6-2*.⁵² Pattern recognition for BDG by factor G may be accomplished by a carbohydrate-binding cleft that is evolutionarily conserved between the horseshoe crab and the bacterium. In contrast, a crystal structure of the extracellular domain of mouse dectin-1 exhibits no structural similarity to the structure of the Z2 module.⁵³

PATHOGEN RECOGNITION BY HEMOCYTE- AND PLASMA-DERIVED LECTINS

Four types of lectins have been identified in hemocytes: tachylectin-1, tachylectin-2, tachylectin-3 and tachylectin-4.⁵⁴⁻⁵⁷ Tachylectin-1 interacts with 2-keto-3-deoxyoctonate on Gram-negative bacteria, whereas tachylectin-2 binds to GlcNAc or GalNAc and recognized lipoteichoic acids of Gram-positive bacteria. In contrast, tachylectin-3 specifically recognizes a certain sugar moiety on O-antigens of S-type LPS. Tachylectin-4 also recognizes the O-antigen of *E. coli* O111:B4 and shows ligand specificity for colitose (3-deoxy-L-fucose). In addition, fibrinogen homologues, tachylectins-5A and -5B, have been identified in hemolymph plasma as acetyl-group-recognizing lectins.³⁴ Tachylectin-5A possesses extraordinarily strong hemagglutinating activity against all types of human erythrocytes (minimum agglutinating concentration ~0.004 µg/ml). The concentration of tachylectin-5A in the plasma was ~10 µg/ml, suggesting that it plays an important role in the recognition of invading pathogens.

Hemolymph plasma also contains several lectins, such as isoforms of tachylectin-1^{58,59} and three types of C-reactive proteins (CRPs) with functional and structural diversity.⁶⁰ Horseshoe crab CRP is a predominant LPS-binding protein and is upregulated at transcript levels by *Pseudomonas* infection, suggesting the importance of horseshoe crab CRP as a conserved molecule for pathogen recognition.⁶¹ CRP from *L. polyphemus* forms extended fibrillar structures that encapsulate liposomes in the presence of Ca²⁺ ions.⁶² Although tachylectin-P, an isoprotein of tachylectin-1, is present in the perivitelline space of the egg, its physiological function in innate immunity remains unknown.⁶³

The crystal structural analyses of tachylectin-2 and tachylectin-5A point out the importance of multivalency for achieving high specificity and high affinity. Tachylectin-2 contains five tandem WD repeats and adopts a five-bladed β-propeller structure with five equivalent GlcNAc/GalNAc-binding sites.⁶⁴ Each propeller blade has an independent binding site for the ligand. The specific recognition by tachylectin-2 is reinforced by the short distance between the individual binding sites, according to the pentagonal geometry. The crystal structure of tachylectin-5A is readily superimposed onto that of the C-terminal polymerization domain of the γ-chain of fibrinogen.⁶⁵ The polymerization pocket within the γ-chain structurally corresponds to the acetyl group binding site of tachylectin-5A, a finding that highlights the evolutionary connection between hemostasis and nonself recognition. Tachylectin-5A is present in oligomer in hemolymph plasma and its propeller like arrangement is evident by electron microscopy.³⁴ Given that tachylectins -2 and -5 exhibit virtually no side- or main-chain conformational changes upon ligand binding, it is likely that the polyvalent nature of these lectins underlies their avidity by allowing them to recognize specific densities or clustering ligands on pathogen surfaces.

STERILIZATION BY ANTIMICROBIAL PEPTIDES

Antimicrobial peptides are widely recognized to be important for innate immunity against invasive microbes. Several kinds of cysteine-rich peptides with antimicrobial activity are stored in the dense granules, including big defensin, tachycitin and tachystatins.⁶⁶⁻⁶⁹ The NMR structural analyses have determined the solution structures

of these antimicrobial peptides.⁷⁰⁻⁷³ For example, the N-terminal domain of big defensin possesses a more potent antimicrobial activity against Gram-positive bacteria than the C-terminal domain.⁶⁶ In contrast, the C-terminal domain homologous to mammalian β -defensins displays more potent antimicrobial activity than the N-terminal domain against Gram-negative bacteria. The structure of big defensin reveals a new class within the defensin family; the C-terminal domain adopts a β -defensin structure, whereas the N-terminal domain forms a unique globular conformation. The hydrophobic N-terminal domain, but not the C-terminal domain, undergoes a conformational change in micelle solution, possibly associated with the antimicrobial activity against Gram-positive bacteria. Interestingly, the antimicrobial activity of an antimicrobial peptide (tachycitin) is synergistically enhanced when another antimicrobial peptide (big defensin) is present.⁶⁷ Horseshoe crab antimicrobial peptides show specific chitin-binding activity, which may play an important role in innate immunity.⁶⁶⁻⁶⁹

CONVERSION OF HEMOCYANIN TO PHENOLOXIDASE BY ANTIMICROBIAL PEPTIDES AND COAGULATION FACTORS

In crustaceans and insects, the prophenoloxidase activation system is an important part of innate immunity, where it acts to detect and kill invading pathogens as well as to synthesize melanin for wound healing and encapsulation of pathogens.^{46,47} However, prophenoloxidase has not been identified in horseshoe crabs. Both prophenoloxidase and hemocyanin contain two functional copper-binding sites capable of reversibly binding an oxygen molecule.⁷⁴ Horseshoe crab hemocyanin is present at high concentrations in hemolymph plasma (~70 mg/ml) and acts as an oxygen carrier under physiological conditions. Arthropod prophenoloxidases are known to be nonenzymatically activated by treatment with detergents, lipids, or organic solvents.⁷⁵ In horseshoe crab as well, the induction of phenoloxidase activity is evident upon similar treatment.⁷⁶ Amphiphilic substances such as SDS and phosphatidylethanolamine convert horseshoe crab hemocyanin to phenoloxidase. Moreover, consistent with the amphiphilic nature of antimicrobial peptides, tachyplesin interacts with hemocyanin and induces its intrinsic phenoloxidase activity.²⁸ This tachyplesin-induced activity is inhibited by phenylthiourea, a typical inhibitor of phenoloxidase. Chemical modification of tachyplesin at Trp-2 or Tyr-8 and Try-13 on its hydrophobic face, but not modification of basic residues on its cationic face, significantly impairs its interaction with hemocyanin, implicating the hydrophobic face of tachyplesin in the functional conversion of hemocyanin to phenoloxidase. Although tachyplesin is the most effective activator of hemocyanin, other antimicrobial peptides, such as tachystatins and tachycitin, significantly induce its phenoloxidase activity.

On the other hand, the proclotting enzyme or coagulation factor B also converts hemocyanin to phenoloxidase and the resulting phenoloxidase activity reaches a plateau at a 1:1 molar ratio, whereas factor C, factor G, or trypsin does not convert hemocyanin.⁷⁷ The proteolytic cleavage of hemocyanin is not required for the functional conversion and the zymogen forms are effective activators. The clip domain of the proclotting enzyme or coagulation factor B may promote the interaction of these factors with hemocyanin to effect its functional conversion to an active phenoloxidase.

The origin of arthropod hemocyanins appears to be an ancient prophenoloxidase-like protein.⁷⁸ Under physiological conditions, arthropod prophenoloxidases require proteolytic cleavage for activation by a specific protease.⁷⁹ *Tarantula* hemocyanin expresses

phenoloxidase activity after limited proteolysis with trypsin or chymotrypsin.⁸⁰ In the crayfish *Pacifastacus leniusculus*, the antibacterial peptide astacidin 1 is released from the C-terminal part of hemocyanin by a cysteine-like protease and is upregulated by LPS or BDG injection.⁸¹ Horseshoe crab hemocyanin and human hemoglobin are also activated by microbial proteases, resulting in the production of reactive oxygen species as an antimicrobial strategy.⁸²

TGase-DEPENDENT CROSS-LINKING IN THE IMMUNE SYSTEM

In crustaceans, hemolymph coagulation depends directly on intermolecular ϵ -(γ -glutamyl)lysine cross-linking of a vitellogenin-related protein by TGase without prior proteolytic cleavage.^{83,84} Recently, in *Drosophila*, TGase and Fondue have been identified as being involved in larval hemolymph coagulation.⁸⁵ In horseshoe crabs, the coagulation cascade triggered by LPS or BDG promotes the conversion of coagulogen to coagulin, resulting in noncovalent polymerization of coagulins in a head-to-tail manner.⁸⁶ TGase of *T. tridentatus* is not present in hemolymph plasma and is restricted to cytoplasm of the hemocyte.^{87,88} Coagulogen is not a substrate for TGase and coagulin is cross-linked to other hemocyte-derived proteins, such as the cysteine-rich protein stablin and the proline-rich protein proxin, resulting in coagulin fibrils with enhanced stability.^{89,90} Stablin interacts with LPS and lipoteichoic acids and exhibits bacterial agglutinating activity against both Gram-negative and Gram-positive bacteria. Consequently, stablin colocalizes with coagulin fibrils that are cross-linked with proxin, effectively trapping bacteria. In addition, stablin binds to chitin, a major component of the arthropod cuticle. Proxin and stablin may promote not only the formation of the stable clotting fibrils but also the immobilization of invading microbes at sites of injury. Horseshoe crab cuticular proteins with chitin-binding activity have been identified as substrates for TGase.⁹¹ One of these, caraxin-1 (carapace-derived chitin-binding protein for protein cross-linking) is specifically localized to the sub-cuticular epidermis.⁹² Recombinant caraxin-1 exists as ~20-mer in solution and these oligomers are cross-linked by TGase to form an elaborate mesh of honeycomb structures that is distinguishable, by electron microscopy, from the clotting mesh triggered by LPS.

Horseshoe crab hemocytes are actively motile and one of the principal functions of these hemocytes is to seal scars in the cuticle.⁹³ According to observations on the wound repair process of *L. polyphemus*, a coagulation plug is formed within 10 min and the coagulum is then infiltrated by hemocytes to form a cellular plug within 24 h.⁹⁴ The sub-cuticular epithelial cells begin to migrate into the wound after 15 days; the epithelial cells span the wound between the cut ends of the exoskeleton by day 30 and then probably secrete cuticular components to complete the wound repair process. At the initial stage of this wound repair process, sufficient quantities of TGase may be secreted from hemocytes recruited to the site of injury and may then be immediately activated by Ca^{2+} ions in hemolymph plasma. At the same time, caraxin-1 may be secreted from the sub-cuticular epidermis. The resulting cross-linked clotting fibrils and caraxin mesh may function to seal the wound to stop bleeding, serve as a barrier to the entry of pathogens into the interior of the animal via the wound and operate as a transient extracellular matrix for the migration of epithelial cells that facilitate wound healing (Fig. 2).

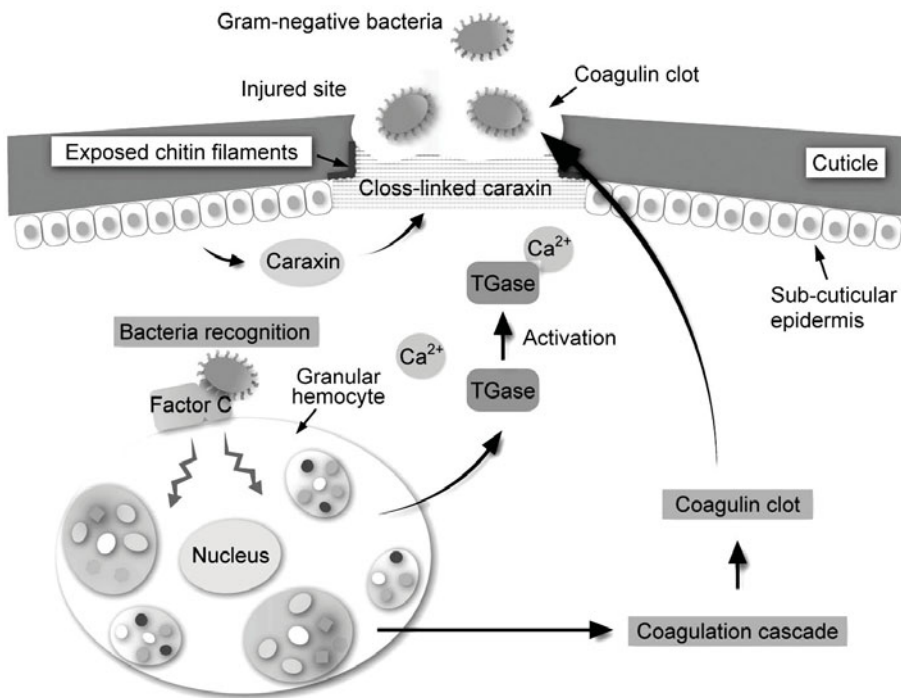


Figure 2. A hypothetical scheme for TGase-dependent cross-linking of caraxins at injured sites. In response to stimulation by LPS, TGase is secreted from the recruited hemocytes at injured sites and immediately activated by Ca^{2+} in plasma, which leads to the crosslinking of caraxins localized in the sub-cuticular epithelial cells. Caraxins may serve to provide an effective mesh to fix invading pathogens at injured sites in cooperation with the clotting mesh.

PATHOGEN RECOGNITION BY THE COMPLEMENT SYSTEM

A complement-related protein, α_2 -macroglobulin, has been previously identified in horseshoe crabs,⁹⁵ and a homolog of complement component C3 has been identified in *C. rotundicauda* (CrC3)⁹⁶ and *T. tridentatus* (TtC3),⁹⁷ indicating the presence of a complement system capable of promoting the phagocytosis of invading microbes in protostomes. CrC3 or TtC3 consists of 1,716 residues, with an overall domain structure that is identical to that of mammalian C3, including α_2 -macroglobulin domains, complement-urchin-bone domains, a thioester-containing domain, an anaphylatoxin domain and a C345C domain. The sequence identity between TtC3 and CrC3 (98%) is considerably higher than that between coagulogens from the two species (90%).⁹⁸ Horseshoe crab C3 is present as a disulfide-linked three-chain structure in hemolymph plasma, unlike mammalian C3, which is present in a two-chain form.⁹⁷

The horseshoe crab complement system promotes the deposition of C3b on the surface of Gram-negative or Gram-positive bacteria. In TtC3, LPS, but not zymosan, peptidoglycan, or laminarin, strongly induces the proteolytic conversion of TtC3 to TtC3b, highlighting the selective response of the complement system to LPS stimulation. An antifactor C antibody inhibits both the proteolytic conversion of TtC3 and the deposition

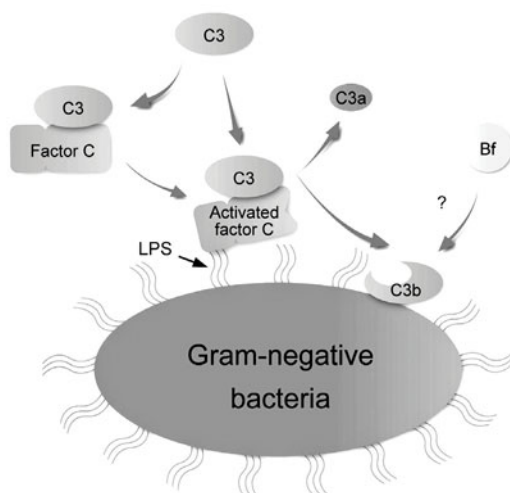


Figure 3. A proposed mechanism for the activation of horseshoe crab complement C3 by factor C localized on the surface of Gram-negative bacteria in the initial phase of the complement system. Factor C recognizes LPS on Gram-negative bacteria and recruits TtC3 by the affinity between the two proteins. The resulting activated factor C converts TtC3 to TtC3b, which is deposited on the surface of bacteria. Whether the activated factor C can convert horseshoe crab Bf leading to the formation of a vertebrate like C3 convertase (C3bBb) remains to be examined.

of TtC3b on the surface of Gram-negative bacteria. Moreover, activated factor C present on the surface of Gram-negative bacteria directly catalyzes the proteolytic conversion of TtC3, thereby promoting TtC3b deposition (Fig. 3). TtC3 interacts with factor C ($K_d = 4.9 \times 10^{-8}M$) and TtC3 is present at a concentration of at least 300 $\mu g/ml$ in hemolymph plasma, whereas the amount of factor C in hemolymph plasma is very low ($\sim 10 \mu g/ml$).⁹⁷ The relatively high concentration of TtC3 and its high affinity to factor C suggest that factor C exists in a complex with TtC3 in hemolymph plasma and that the formation of this complex is a prerequisite for the immediate activation of TtC3 by factor C on the surface of Gram-negative bacteria. In the alternative pathway of the mammalian complement system, the interaction between C3b and Bb is essential to form C3 convertase (C3bBb). A homolog of complement Bf has been identified in *C. rotundicauda*.⁹⁶ Although the physiological function of the Bf homolog in the horseshoe crab complement system remains unknown, it is likely that it may be responsible for the formation of the second C3 convertase (Fig. 3).

CONCLUSION AND FUTURE PROSPECTS IN THE HORSESHOE CRAB INNATE IMMUNE SYSTEM

In response to stimulation by LPS, hemocyte secretes several kinds of immunocompetent molecules, such as coagulation factors, lectins, antimicrobial peptides, protein substrates for TGase and also cytoplasm-derived TGase. The hemocyte possesses a feedback mechanism in which the antimicrobial peptides serve as endogenous mediators. The coagulation cascade results in the formation of coagulin fibrils that are subsequently

stabilized by TGase-dependent cross-linking of proixin and stablin. A sub-cuticular epidermis-derived caraxin forms a TGase-stabilized mesh at sites of injury. Invading pathogens are agglutinated by both hemocyte- and plasma-derived lectins. In addition, coagulation factors and tachyplesin convert hemocyanin to phenoloxidase. In the plasma, factor C acts as an LPS-sensitive complement C3 convertase. In this manner, horseshoe crabs have a sophisticated innate immune response network that coordinately effects pathogen recognition and killing, prophenoloxidase activation, complement activation and TGase-dependent wound healing.

Interestingly, the antifactor C antibody exhibits no effect on the deposition of TtC3b on *Staphylococcus aureus*, suggesting the presence of a factor C-independent pathway to initiate the opsonization of Gram-positive bacteria.⁹⁷ Complement factors required for the deposition of TtC3b on Gram-positive bacteria remain to be examined. Factor C and complement Bf from *C. rotundicauda* interact with plasma-derived lectins, such as galactose-binding protein, carcinolectin-5 and CRP.⁹⁹ Therefore, the protease-lectin complexes on the surface of Gram-positive bacteria may enhance the deposition of C3b. Although the complement-dependent clearance system of invading pathogens in horseshoe crabs remains to be examined, phagocytosis of Gram-positive bacteria by hemocytes both in vivo and in vitro is inhibited by protease inhibitors, raising the possibility that the proteolytic dependence of opsonization by C3b may underlie phagocytosis by hemocytes.⁹⁶

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TICK INNATE IMMUNITY

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Abstract: Ticks are blood feeding parasites transmitting a wide variety of pathogens to their vertebrate hosts. The vector competence of ticks is tightly linked with their immune system. Despite its importance, our knowledge of tick innate immunity is still inadequate and the limited number of sufficiently characterized immune molecules and cellular reactions are dispersed across numerous tick species. The phagocytosis of microbes by tick hemocytes seems to be coupled with a primitive complement-like system, which possibly involves self/nonself recognition by fibrinogen-related lectins and the action of thioester-containing proteins. Ticks do not seem to possess a pro-phenoloxidase system leading to melanization and also coagulation of tick hemolymph has not been experimentally proven. They are capable of defending themselves against microbial infection with a variety of antimicrobial peptides comprising lysozymes, defensins and molecules not found in other invertebrates. Virtually nothing is known about the signaling cascades involved in the regulation of tick antimicrobial immune responses. Midgut immunity is apparently the decisive factor of tick vector competence. The gut content is a hostile environment for ingested microbes, which is mainly due to the antimicrobial activity of hemoglobin fragments generated by the digestion of the host blood as well as other antimicrobial peptides. Reactive oxygen species possibly also play an important role in the tick-pathogen interaction. The recent release of the *Ixodes scapularis* genome and the feasibility of RNA interference in ticks promise imminent and substantial progress in tick innate immunity research.

INTRODUCTION

Ticks (Acari: Ixodida) are parasitic mites that suck blood from their vertebrate hosts. Systematically, they belong to the Phylum Arthropoda, Class Arachnida, Subclass Acari, Order Parasitiformes and Suborder Ixodida.¹ To date, more than 900 tick species have been described and divided into two major families—Ixodidae (generally referred to as hard ticks) and Argasidae (also known as soft ticks).² As this common nomenclature implies, the hard ticks possess a sclerotized dorsal body surface plate (scutum), whereas the soft ticks have only a leathery cuticle.³ This most conspicuous morphological feature of Ixodidae and Argasidae is just one hallmark for the other numerous biological, physiological and ecological differences between these two families,³ out of which we mention just the most relevant to the topics of this chapter. Soft ticks, represented e.g., by the genera *Argas* or *Ornithodoros*, are multi-host parasites, having several nymphal stages (2-8) that all feed rapidly on their hosts (within minutes to hours). The adult ticks can feed repeatedly and the females, which mate away from the host, deposit just a limited number of eggs (few hundreds) after each feeding. The life span of Argasidae is quite long (up to several years) and they are capable to live without food for months between individual blood meals as nymphs or adults. In contrast, hard ticks possess only three developmental stages—larvae, nymphs and adults. Most Ixodidae (e.g., genera *Ixodes*, *Amblyomma*, *Dermacentor*, *Haemaphysalis*) are three-host ticks, where each stage can feed on a different host. The hard ticks feed slowly for several days, depending on the stage and species. The adult females mate usually on the host and after full engorgement they drop off. The blood meal is digested within several weeks and the female oviposits thousands of eggs and dies. Few, veterinary important species, such as the cattle tick *Rhipicephalus* (formerly *Boophilus*) *microplus*, are one-host ticks where larvae, nymphs and adults remain and feed on the same bovine host. The obvious differences in physiology and feeding strategy strongly support the theory that adaptation to blood-feeding in hard and soft ticks has evolved independently.⁴

Ticks do harm to their hosts as obligatory ectoparasites, sometimes causing severe blood loss. More importantly, the main danger ticks present to their hosts is the enormous variety of pathogens they can transmit, including viruses, bacteria and protozoa.⁵ A comprehensive survey of tick-borne pathogens causing serious diseases to humans or domestic animals was reviewed recently.⁶ It demonstrates that the most recognized human illnesses (e.g., Lyme disease, tick-borne encephalitis, granular anaplasmosis) or animal diseases (e.g., anaplasmosis, babesiosis, theileriosis, African swine fever) represent just the tip of the iceberg of the entire problem of ticks as disease vectors.⁶

The success of pathogen transmission by ticks is due to the long period of co-evolution between pathogen and vector and also because of specific aspects of tick biology. Pathogens take advantage of the modulation of host haemostatic, inflammatory and immune responses mediated by the inexhaustible pharmacology of molecules present in tick saliva.⁷ Moreover, ingested microbes are not exposed to immediate proteolytic attack by digestive enzymes within the gut lumen, since ticks digest blood intracellularly in digestive lysosomal vesicles.³ On the other hand, it is obvious that ticks have to possess efficient defense mechanisms to eliminate microbial infections, or at least to mitigate it to a level tolerable for the vector. Despite its importance, we still know very little about how transmitted pathogens can evade or withstand the innate immune responses of ticks since, the vector-pathogen interface is the least understood site of the pathogen-tick-host interaction triangle.

Invertebrate animals lack an adaptive immune system and their defense against potential pathogens has to rely on a network of cellular immune reactions (e.g., phagocytosis, encapsulation and nodulation) and humoral factors involved in pathogen recognition and elimination (pattern recognition receptors, lectins, complement-like system, pro-phenoloxidase activation, hemolymph coagulation, antimicrobial peptides, reactive oxygen species, etc.). During the past two decades, our knowledge of innate immunity in invertebrates has developed rapidly thanks to the considerable body of work carried out on the model insect *Drosophila melanogaster*, other arthropods, such as the horseshoe crab, freshwater crayfish and prochordates such as ascidia (for review, see ref. 8 and this book). The detailed model of innate immune system based on that from *Drosophila*⁹ provided a solid basis for further investigation of immune responses and parasite transmission in insect disease vectors, such as mosquitoes¹⁰ or tse-tse flies.¹¹

In contrast to the model invertebrates and blood-sucking insects, our understanding of innate immunity in ticks is still inadequate. Only a limited number of described cellular immune reactions and sufficiently characterized immune molecules (see the overview in Table 1 and refs. 12, 13 for the recent reviews) allow extrapolation to the immune system of other model arthropods, especially the horseshoe crab, which is phylogenetically the closest relative of ticks within the subphylum Chelicerata. A few years ago, it seemed that research on tick immune system would probably remain forever at a purely descriptive level. However, several recent advances reversed this pessimistic outlook. The availability of large scale EST data sets on different tick species, including the first version of the American deer tick *Ixodes scapularis* genome, released in December 2008,¹⁴ the feasibility of RNA interference (RNAi) in ticks¹⁵ and the establishment of several tick in vitro cell culture lines as valuable models for analyzing the immunological interactions with tick-borne pathogens¹⁶ opened the gate to the postgenomic era in tick-host-pathogen interaction research.

In this chapter, we review the current knowledge on tick cellular events and molecules involved in defense against ingested and transmitted microbes. We also highlight further perspectives in the research of tick innate immunity relying mainly on the recently implemented reverse genetics and functional genomics approaches.

DEFENSE MECHANISMS IN THE TICK HAEMOCOEL

Cellular Immune Responses

Similar to other arthropods, ticks possess an open circulatory system surrounding their internal organs.³ It was demonstrated in the tick *Dermacentor andersoni* that during feeding the hemolymph volume increases linearly from about 2-3 μ l in unfed to almost 150 μ l in fully engorged females. However, the ratio of hemolymph content to the total body weight remains constant at about 23%.¹⁷

Based on morphological and physiological studies of hemolymph cells in different tick species, several hemocyte types have been described.^{3,18-20} Most reports are consistent in the description of three basic classes of tick hemocytes, namely **plasmatocytes**, **granulocytes I** and **granulocytes II**. Phagocytic activity is mainly attributed to plasmatocytes and granulocytes I, whereas granulocytes II are nonphagocytic cells. In some tick species, another hemocyte type resembling insect **spherulocytes** has been described.^{3,20} The

Table 1. Overview of characterized tick immune molecules

Defense System	Molecule	Tick Species ^a	Method ^{b/} Starting Material ^{c,d}	Molecular Characteristics	Protein Localization/ Gene Expression ^d	Function and/ or Other Characteristics	References GenBank
Lectin-complement system							
Lectins-agglutinins	Lectin (DorinM)	<i>O. moubata</i>	Classical/HI	37 kDa, 3 × glycosylated noncovalent oligomer	HI/H, SG	Lectin, sialic acid-specific Fibrinogen related protein	36,37 AY333989
	OMFREP	<i>O. moubata</i>	Reverse genetics/H	28.9 kDa	/H, SG	Fibrinogen related protein	40 AF527411
	Ixoderin A	<i>I. ricinus</i>	Reverse genetics/H	30.8 kDa	/H, SG, MG	Fibrinogen related protein	40 AY341424
	Ixoderin B	<i>I. ricinus</i>	Reverse genetics/SG	32.6 kDa	/SG	Fibrinogen related protein	40 AY643518
	Galectin (OmGa-lectin)	<i>O. moubata</i>	Reverse genetics/multiple tissues	37.4 kDa	/Multiple tissues	Intracellular protein	41 AB255165
Complement-like molecules	α2-macroglobulin (TAM)	<i>O. moubata</i>	Classical/HI	2 × 160 kDa, glycosylated noncovalent dimer	HI/H, SG	Universal protease inhibitor	42,43 AF538967
	α2-macroglobulin (IrAM)	<i>I. ricinus</i>	Reverse genetics/SG	2 × 160 kDa, glycosylated noncovalent dimer	HI/H, SG	Function in phagocytosis of <i>C. indologenes</i>	23 EU835901

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Table 1. Continued

Defense System	Molecule	Tick Species ^a	Method ^{b/} Starting Material ^{c,d}	Molecular Characteristics	Protein Localization/ Gene Expression ^d	Function and/ or Other Characteristics	References GenBank
	Factor D-like	<i>D. variabilis</i>	Reverse genetics/H	39 kDa, clip domain, protease homolog	/H	<i>E. coli</i> inducible	46 AY161279
Antimicrobial peptides							
Lysozymes	c-type lysozyme	<i>O. moubata</i>	Classical/Gut contents	14 kDa, pI 9.7	Gut contents/ MG	Active against <i>M. luteus</i> , blood-meal inducible	75,76 AAL17868
	c-type lysozyme	<i>D. variabilis</i>	Reverse genetics/H	14 kDa, pI 9.97	/H, MG, FB	<i>E. coli</i> and <i>R. montanensi</i> inducible	55,77 AY183671
	c-type lysozyme	<i>D. andersoni</i>	Reverse genetics/Cell line DEA100	14 kDa, pI 9.91			55 AY207371
Defensins	Defensin A,B,C,D	<i>O. moubata</i>	Reverse genetics/4th instar nymphs	4 kDa mature peptide	Gut contents (DefA)/MG	Active against Gram ⁺	78-80 (DefA) AB041816
	Defensin I (varisin)	<i>D. variabilis</i>	Classical/HI	4.2 kDa mature peptide	H	Active against Rickettsia	59-62 AY181027
	Defensin	<i>R. microplus</i>	Classical/H	4,291 Da mature peptide	H	Active against <i>M. luteus</i>	65 AY233213
	Defensin (amercin)	<i>A. americana</i>	Reverse genetics/H	4,050 Da mature peptide	/MG, H, FB, SG		58 DQ864986

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Table 1. Continued

Defense System	Molecule	Tick Species ^a	Method ^{b/} Starting Material ^{c,d}	Molecular Characteristics	Protein Localization/ Gene Expression ^d	Function and/ or Other Characteristics	References GenBank
	Defensin (longicin)	<i>H. longicornis</i>	Reverse genetics/MG	5,820 Da mature peptide	MG	Active against Gram ⁺ , Gram ⁻ fungi and <i>Babesia</i> sp.	81 ABI05544
	Defensin (HI-gut)	<i>H. longicornis</i>	Reverse genetics/MG	5.8 kDa pro-defensin	/MG	LPS inducible	82 EF432731
	Defensin	<i>I. persulcatus</i>	Reverse genetics/ Whole nymphs	4,199 Da mature peptide	/MG	Active against Gram ⁺	56 AB469201
	Defensin (scapularisin)	<i>I. scapularis</i>	Reverse genetics/ Mixed tissues	4,180 Da mature peptide	/MG, (H,FB)		84 AY660970
	Defensin	<i>I. ricinus</i>	Reverse genetics/ Whole females	4,496 Da mature peptide	/MG	Upregulated by Borrelia-infected blood meal	83 AY335442
Defensin-like	Noncationic defensins (ADP1, ADP2)	<i>A. hebraeum</i>	Reverse genetics/ Syngonium	4,612 mature ADP2	HI (ADP2)	Active against <i>S. aureus</i> > <i>E. coli</i> (ADP2)	63 AY437137 AY437138
(No RVRK cleavage motif)	Defensin-2	<i>D. variabilis</i>	Reverse genetics	4.7 kDa pro-defensin	/FB, MG, Ov		77 AY159879
	Defensin (HI-sal)	<i>H. longicornis</i>	Reverse genetics/SG	6.5 kDa pro-defensin	/SG	LPS inducible	82 EF432732

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Table 1. Continued

Defense System	Molecule	Tick Species ^a	Method ^b / Starting Material ^{c,d}	Molecular Characteristics	Protein Localization/ Gene Expression ^d	Function and/ or Other Characteristics	References GenBank
Other AMPs	Hebraein	<i>A. hebraeum</i>	Reverse genetics/ Syngonium	11 kDa histidine-rich	HI	Active against <i>S. aureus</i> , <i>E. coli</i> , yeast <i>C. glabrato</i>	64 AY437139
	Microplusin	<i>R. microplus</i>	Classical/HI, eggs	10,204 Da, histidine-rich	HI, Ov	Active against <i>M. luteus</i> ; divalent metal chelator	65,66,109 AY233212
	Ixodidin	<i>R. microplus</i>	Classical/H	7,103 Da	H	Active against <i>M. luteus</i> ; chy- motrypsin-elastase inhibitor	67 P83516
	Ixosin	<i>I. sinensis</i>	Classical/SG	2,870.5 Da, mature peptide	SG	Activity against <i>E. coli</i> , <i>S. aureus</i> , <i>C. albicans</i>	105 DQ100462
	Ixosin B	<i>I. sinensis</i>	Classical/SG	3,814.5 Da, mature peptide	SG	Activity against <i>E. coli</i> , <i>S. aureus</i> , <i>C. albicans</i>	106 EU047746
	IsAMP	<i>I. scapularis</i>	Classical/ Saliva	5,300 Da, mature peptide	Saliva/H, SG, FB	Activity against Gram ⁺ , Gram ⁻ , up-regulated by Borrelia infection	108 AF483734
	Hemoglobin fragment	<i>R. microplus</i>	Classical/Gut contents	3,206 Da		Activity against Gram ⁺ , fungi	68,71

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Table 1. Continued

Defense System	Molecule	Tick Species ^a	Method ^{b/} Starting Material ^{c,d}	Molecular Characteristics	Protein Localization/ Gene Expression ^d	Function and/ or Other Characteristics	References GenBank
		<i>O. moubata</i>	Classical/Gut contents			Activity against <i>S. aureus</i>	69
		<i>D. variabilis</i>	Classical/Gut contents			Activity against <i>M. luteus</i>	70
Protease inhibitors							
	Kunitz-type serine protease inhibitor (KPI)	<i>D. variabilis</i>	Reverse genetics/MG	35.5 kDa, 5 Kunitz BPTI domains	/MG	Trypsin inhibitor, anticoagulant; active against Rickettsia	89 ABY26517
	HI-cystatin 2	<i>H. longicornis</i>	Reverse genetics/MG	12.9 kDa	/MG	Cysteine peptidase inhibitor babesia-cidal	88 DQ364159
	BmSI-7, BmSI-6—subtilisin inhibitors	<i>R. microplus</i>	Classical				
			/eggs	7,408 Da (BmSI-7) 7,271 Da (BmSI-6)	Eggs/Ov, MC, SG	Active against PR1 protease from the fungus <i>Meth-arizium anisoplae</i>	110 (BmSI-7) GQ452779

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Table 1. Continued

Defense System	Molecule	Tick Species ^a	Method ^{b/} Starting Material ^{c,d}	Molecular Characteristics	Protein Localization/ Gene Expression ^d	Function and/ or Other Characteristics	References GenBank
Antioxidant system Antioxidant	Catalase	<i>R. microplus</i>	Classical/MG	inhibited by aminotriazol			95
	Glutathion peroxidase (HlPrx)	<i>H. longicornis</i>	Reverse genetics/SG	26.5 kDa	SG/SG	Antioxidant activity	99 AB038382
	Phospholipid-hydroperoxideglutath. peroxidase	<i>R. microplus</i>	Reverse genetics/cell line C34, coumaphos resistant	18.9 kDa contains selenocysteine		Increased expression in acaricide-resistant strains	102 DQ172828
	Glutathion S-transferase	<i>R. microplus</i>	Classical/Whole larvae	25.6 kDa, pI 8.53			98 AAD15991
	Glutathion S-transferase	<i>R. microplus</i>	Reverse genetics/SG	25.6 kDa	SG (MG)/SG, MG		100 AF366931
	Glutathion S-transferase	<i>D. variabilis</i>	Reverse genetics/H	24 kDa, pI 9.16 (1) 24 kDa, pI 5.27 (2)	MG, MG /MG (1); MG, Ov (2)	Blood meal inducible	97 (1) DQ224235 (2) AY241958

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Table 1. Continued

Defense System	Molecule	Tick Species ^a	Method ^{b/} Starting Material ^{c,d}	Molecular Characteristics	Protein Localization/ Gene Expression ^d	Function and/ or Other Characteristics	References GenBank
Signaling cascade Transcription factor	Akirin (Subole- sin, protective antigen 4D8)	<i>I. scapularis</i>	Reverse genetics	20.6 kDa		Intracellular trans- cription factor	115 AY652654

^a *A. Amblyomma*; *D. Dermacentor*; *H. Haemaphysalis*; *I. Ixodes*; *O. Ornithodoros*; *R. Rhipicephalus*.

^b Classical—from protein to gene, Reverse genetics—from gene to protein.

^c Tissue used for protein purification or RNA isolation, respectively.

^d H—hemocytes; HI—hemolymph; SG—salivary glands; MG—midgut; Ov—ovaries; FB—fat body.

hemocytes are believed to differentiate from stem cells called **prohemocytes**, which are only rarely detected in the hemolymph.²⁰

Phagocytosis of foreign material and microbes by tick hemocytes has been studied both in vivo and in vitro in several tick species. Fluorescent polystyrene beads inoculated into the haemocoel of *Ornithodoros moubata* were phagocytosed by plasmatocytes twice as much as by granulocytes.¹⁹ Hemocytes of *O. moubata* were also reported to phagocytose the yeast *Candida haemulonii*²¹ or the yellow pigmented Gram⁻ bacteria *Chryseobacterium indologenes*.²² Despite the active cellular response against these microbes, infection with these pathogens caused high mortality rates of these soft ticks.^{21,22} Recent RNAi experiments revealed that phagocytosis of *C. indologenes* by hemocytes of the hard tick *Ixodes ricinus* is mediated by a plasma α_2 -macroglobulin²³ and possibly also by other thioester-containing proteins²⁴ (see also the later section on complement-related molecules). Pereira et al²⁵ investigated the phagocytic activity of the *R. microplus* hemocytes upon injection of the yeast *Saccharomyces cerevisiae*. They demonstrated that plasmatocytes is the major cell type involved in phagocytosis of the yeast. Moreover, they were able to show in an in vitro assay that an oxidative burst in the phagocytic hemocytes occurred through the production of the reactive oxygen species superoxide and H₂O₂.

The process of hemocytic **encapsulation** was examined using small pieces of Epon-Araldite implanted under the cuticle of the American dog tick, *Dermacentor variabilis*.²⁶ This study showed the successive involvement of Type I and Type II granulocytes as well as plasmatocytes in the gradual forming of a multiple cell layer capsule around the implant. In addition, the authors observed Epon-Araldite particles surrounded by a fibrous matrix, which they interpreted as hemolymph coagulation. On the other hand, they did not observe any sign of melanin formation on the surface of the implant. Another report on **nodulation** in ticks was also demonstrated in *D. variabilis*.²⁷ The challenge of ticks by direct inoculation of *Escherichia coli* into the haemocoel led to the immediate clearance of bacteria from the hemolymph. A direct fluorescence assay revealed that a rapid formation of bacterial clumps surrounded by aggregated and disintegrated hemocytes is responsible for this phenomenon. The aggregated bacteria were not viable, possibly because of the involvement of other immune mechanisms, e.g., bacteriolysis or phagocytosis.²⁷

Thus far, no cell line originating from tick hemocytes has been available. However, some tick cell lines derived from embryonic cells¹⁶ display remarkable phagocytic activity and have been used as useful models for deeper insight into tick cell-pathogen interaction.^{28,29} An elegant example of this approach was provided in a recent work showing how cell lines derived from *I. scapularis* (IDE12) and *Amblyomma americanum* (AAE2) respond to infection by the entomopathogenic fungi *Metarhizium anisoplae*.³⁰

Certainly, much attention has been given to the response of tick cells against the human pathogen *Borrelia burgdorferi*, the causative agent of Lyme disease. The hemocytes from *I. ricinus* were shown to use the mechanism of “coiling” phagocytosis to engulf *B. burgdorferi* spirochetes, as performed by vertebrate phagocytic cells.³¹ Coleman et al demonstrated that *B. burgdorferi* overcome the midgut barrier of the tick *I. scapularis* via a surface bound plasmin, originating from the host blood, which facilitates spirochete dissemination into the tick hemolymph and migration towards the salivary glands.³² At least some of the spirochetes were phagocytosed and the process of their internalization by tick hemocytes was examined by confocal microscopy.³² The differences in the immune responses to the cultivated *B. burgdorferi* upon direct injection into the haemocoel were compared using the natural vector,

I. scapularis and *D. variabilis*, which is unable to transmit the pathogen.³³ The authors found that the spirochetes were rapidly cleared from the hemolymph of *D. variabilis* in contrast to *I. scapularis*, which appeared to provide a benign environment for the pathogen. The immunocompetent *D. variabilis* displayed a faster phagocytic response and a higher increase in hemocytic cells following challenge with the *B. burgdorferi*, as well as a much higher borreliacidal activity in the plasma, as compared to the immunotolerant *I. scapularis*.^{33,34} These results are also in accordance with the study of *B. burgdorferi* phagocytosis by cell lines from *I. scapularis* (IDE12) and *D. andersoni* (DAE15).²⁸

Lectins—The Potential Recognition Molecules

Research on tick lectins/hemagglutinins has been the subject of constant attention since the specific protein-carbohydrate interaction is believed to be a key to understand self/nonself recognition and how the transmitted pathogens evade tick immune responses. The hemagglutination activity, conferring the ability to agglutinate mammalian red blood cells, has been reported for a variety of tick species mainly in the hemolymph, gut and salivary glands. The binding specificity of tick lectins were mainly observed towards N-acetyl-D-hexosamines, sialic acid and glycoconjugates like fetuin (see ref. 35 for a recent review). The only sufficiently characterized tick lectin referred to as Dorin M was purified from the plasma of the soft tick *O. moubata* (Table 1).³⁶ Dorin M is mainly expressed in tick hemocytes and salivary glands and its C-terminal domain is highly homologous to the γ -chain of vertebrate fibrinogens and fibrinogen-related proteins (FREPs), such as mammalian ficolins.³⁷ Unlike ficolins, Dorin M lacks the N-terminal collagen-like domain. This feature is shared by the closely related tachylectins 5A and 5B from the horseshoe crab *Tachypleus tridentatus*,³⁸ which function as pattern recognition molecules.³⁹ Another FREP has been cloned from *O. moubata* (OMFREP) (Table 1) and found to have a similar sequence and expression profile as Dorin M.⁴⁰ By contrast, two FREPs identified in the hard tick *I. ricinus*, namely the Ixoderin A and B, are phylogenetically more distinct and differ also in their tissue expression profile. Ixoderin A gene expression was found in hemocytes, salivary glands and midgut, whereas Ixoderin B was specifically expressed in the salivary glands (Table 1).⁴⁰ Our search through the *I. scapularis* genome revealed that at least three isoforms of Ixoderin A could be identified, multiple (≥ 14) isoforms of Ixoderin B and a still uncharacterized type named Ixoderin C which was predicted to be intracellular protein (Hajdušek, unpublished results).

The gene coding for a protein related to galectins has been isolated from the whole body homogenate of *O. moubata* 4th instar nymphs and was named OmGalec (Table 1).⁴¹ Its gene was found to be expressed in all developmental stages, including eggs and in variety of tissues. The protein is likely to be present intracellularly and it lacks a transmembrane domain. OmGalec sequence contains tandem-repeated carbohydrate recognition domains. The sugar binding specificity of recombinant OmGalec was tested for a broad spectrum of glycans using an automated frontal affinity chromatography system. Its affinity profile corresponds to the general features of the galectin family, e.g., specificity for lactosamine-like disaccharides. The authors speculate that OmGalec may have multiple functions, including in innate immunity.⁴¹

Complement-Related Molecules

Two α_2 -macroglobulins (α_2 M) described in the soft tick *O. moubata* and the hard tick *I. ricinus* designated TAM and IrAM, respectively (Table 1),^{23,42,43} are the first tick representatives of the evolutionary oldest and best conserved family of thioester-containing proteins (α_2 M superfamily), which have an important role in both invertebrate and vertebrate innate immunity. This superfamily comprises in invertebrates three main, phylogenetically distinct groups (i) α_2 -macroglobulins—the universal macromolecular protease inhibitors; (ii) the components C3/C4/C5 of the complement system; (iii) insect thioester proteins (TEPs).⁴⁴ TAM and IrAM share high sequence homology, in addition to disulfide bridge and glycosylation patterns. However, several structural features differentiate them from the majority of vertebrate and invertebrate α_2 Ms: the absence of disulfide bridges covalently binding their subunits, posttranslational processing of their precursors resembling cleavage events typical for C3 and C4 complement components and variability within their bait regions (site of protease attack) generated by alternative splicing.^{23,43} A recent functional study based on RNAi-silencing linked with an in vitro phagocytic assay revealed that IrAM is involved in phagocytosis of the tick pathogen *C. indologenes* by *I. ricinus* hemocytes. The phagocytosis activity was dependent on an active metalloprotease secreted by the bacteria, indicating that interaction of tick α_2 M with a protease from an invading pathogen is linked with a cellular immune response.²³ Interestingly, phagocytosis of the Lyme disease agent *B. burgorferi* was not affected either by IrAM silencing or methylamine treatment indicating that engulfment of the spirochetes is mediated by a different mechanism independent on thioester-containing proteins.²³ We searched for other thioester-containing proteins in the *I. scapularis* genome and found nine different molecules belonging to the α_2 M-superfamily (tentatively named as IsAM1-9).²⁴ The phylogenetic analysis (Fig. 1) revealed that three IsAMs are related to α_2 Ms, three are of the C3-complement type, one is related to insect TEPs and two IsAMs belong to a fourth branch recently renamed macroglobulin complement related (MCR) proteins.⁴⁵ Hence, the tick *I. scapularis* is so far the only organism within the animal kingdom known to us to possess representatives of all major groups of invertebrate thioester-containing proteins.²⁴

Factor D-like clip-domain serine proteinase homologues have been identified from *D. variabilis* (Dv FacD-L) and also from the *D. andersoni* DEA100 cell line (Da FacD-L) (Table 1).⁴⁶ The serine residue of the catalytic triad is substituted with a glycine as in the closely related *Limulus* factor D, having a microbial activity,⁴⁷ or other arthropod clip-domain serine protease homologues playing various roles in immune response.⁴⁶ Dv FacD-L is expressed mainly in tick hemocytes and its mRNA level increased significantly upon intrahaemocoelic inoculation of *E. coli*. A clear ortholog of Dv(a) FacD-L is also present in the *I. scapularis* genome. A better understanding of the role of the immune-responsive FacD-L in tick immunity, as well as other molecules potentially related to a tick complement-like system, awaits appropriate functional genomic studies.

Do Ticks Have a Coagulation or Pro-Phenoloxidase Cascade?

Although some authors admit that tick hemolymph is capable to clotting or melanization,^{3,12,13} there are actually only a few reports dealing with this aspect of tick innate immunity. As mentioned previously, a process resembling hemolymph coagulation has

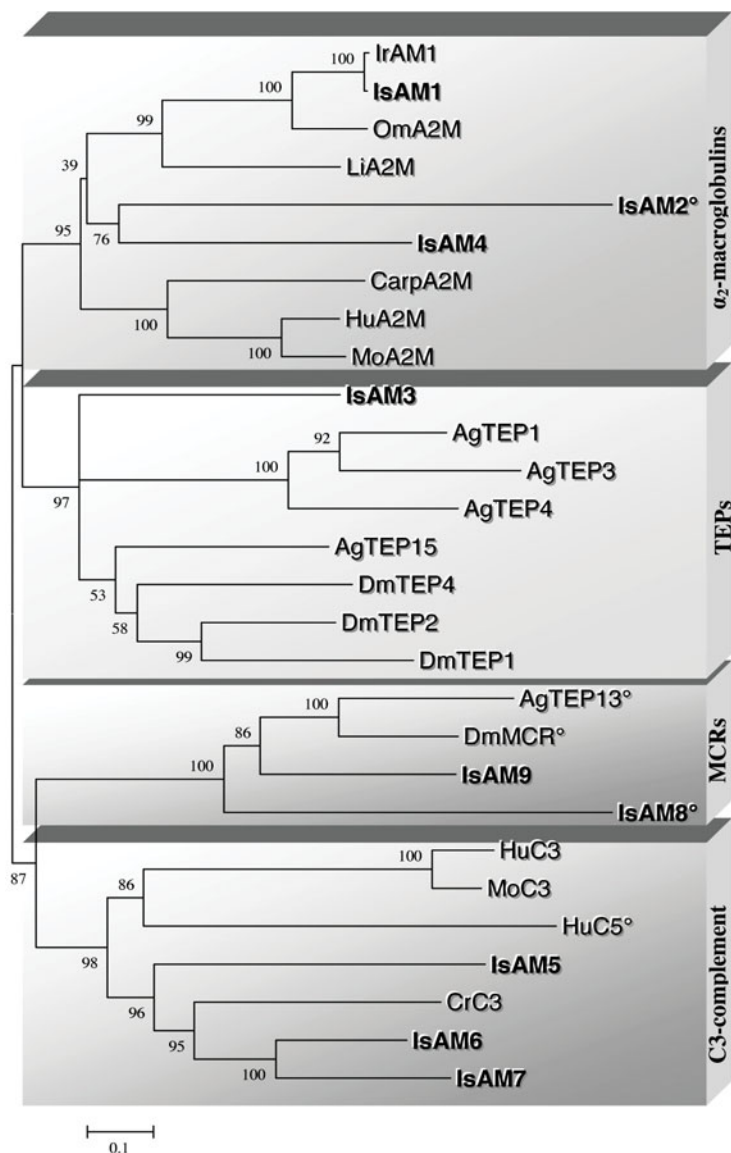


Figure 1. Phylogenetic tree of tick and selected invertebrate and vertebrate thioester-containing proteins (α_2 -macroglobulin superfamily). The tree was constructed by the neighbour-joining method using amino acid sequence of ~700 residues spanning across the conserved α_2 -M superfamily and thioester domains. Numbers on the branches display bootstrap support. **IsAM**(1-9): thioester proteins found in the *Ixodes scapularis* genome; **Ir**: *Ixodes ricinus* (hard tick); **Om**: *Ornithodoros moubata* (soft tick); **Li**: *Limulus polyphemus* (horseshoe crab); **Cr**: *Carcinoscorpius rotundicauda* (horseshoe crab); **Dm**: *Drosophila melanogaster*; **Ag**: *Anopheles gambiae*; **Carp**: *Cyprinus carpio*; **Hu**: human, **Mo**: mouse; The circlets indicate the absent thioester bond. **MCR** stands for macroglobulin-complement-related. Adapted from Burešová.²⁴

been observed in the form of a mass around Epon-Araldite particles implanted under the *D. variabilis* cuticle, although without identification of any of the molecules involved.²⁶ In addition, we were not able to find any gene related either to the horseshoe crab coagulogen or crustacean clotting protein.⁸ On the other hand, the *I. scapularis* genome contains putative transglutaminases. Therefore, a possibility of hemolymph coagulation based on crosslinking of other hemolymph proteins (e.g., the abundant heme lipoproteins^{48,49}) can not completely be ruled out. Considering melanization, there are controversial data about the PPO activity in the hemolymph of ticks. It has been shown no PPO activity in the hemolymph of the hard ticks *A. americanum*, *D. variabilis* and *I. scapularis*,⁵⁰ while such activity was detected in the hemolymph of the soft tick *O. moubata*.⁵¹ Since neither PPO nor hemocyanin, which displays PPO activity in the horseshoe crab,⁵² seem to be present in the tick genome, it would be essential to characterize the molecule responsible for this activity.

Antimicrobial Peptides in the Tick Hemolymph

The existence of a potent antimicrobial activity against various Gram⁺, Gram⁻ bacteria and rickettsiae has been described by Russian researchers in the early 1990s for a relative large set of ticks representing both the Argasidae and Ixodidae families.⁵³ The antibacterial activities were mainly attributed to the c-type **lysozymes** that were purified and partially characterized.^{53,54} Later, two c-type lysozyme orthologues were cloned and sequenced from *D. variabilis* hemocytes and the *D. andersoni* DAE100 cell line (Table 1) and their gene expression was shown to be upregulated upon *E. coli* challenge.⁵⁵

The greatest part of literature on tick antimicrobial peptides (AMPs) deals with **defensins**. This family of small cationic peptides is synthesized as prepropeptides of about 8 kDa, from which approximately 4 kDa mature peptides are cleaved C-terminally via a conserved furin cleavage motif (RVRR). Tick defensins (often having several isoforms) were so far identified in nearly twenty hard and soft tick species.^{12,56-58} In addition to the hemolymph, tick defensins were also reported to be expressed in the midgut and other tissues, including salivary glands and fat body (see below and the Table 1). Typical tick defensins with the furin cleavage site and conserved pattern of six paired cysteines within the mature active peptide can be exemplified by **varisin** from *D. variabilis*. Varisin was purified by reverse-phase high performance chromatography (RP-HPLC) from the hemolymph of *D. variabilis* females challenged with *B. burgdorferi* and was shown to be active against the Gram⁺ *Bacillus subtilis*.^{59,60} This defensin or chicken lysozyme alone have low activity against cultured borrelia spirochetes. However, their combination significantly increased borreliaicidal activity, indicating that a synergistic effect of these molecules together with efficient phagocytosis (mentioned above) is involved in the immediate clearance of spirochetes from the *D. variabilis* hemolymph.^{33,59} The role of varisin in *D. variabilis* innate immunity was also studied by RNAi. Although varisin was shown to be depleted from the hemolymph of the varisin-KD ticks, the activity against the Gram⁺ *Micrococcus luteus* was only reduced by 50%. This result indicates that varisin contributes significantly but not exclusively to the overall antimicrobial activity in the *D. variabilis* hemolymph.⁶¹ A study of the effect of varisin silencing on the transmission of a relevant pathogen, the rickettsia *Anaplasma marginale*, presented a surprising finding. In opposite to authors' expectations, the *A. marginale* infection was significantly reduced in the varisin-KD ticks for yet unknown reasons.⁶²

Two anionic **defensin-like peptides** (ADP1 and ADP2), sharing the conserved cysteine residues of typical defensins but lacking the typical furin cleavage motif, were identified as transcripts in a subtractive cDNA library originating from the syngaglion (central nervous system) dissected from fed vs unfed female *Amblyoma hebraeum* ticks (Table 1).⁶³ The authors succeeded in the purification of ADP2 from tick hemolymph using RP-HPLC and demonstrated its antimicrobial activity against both Gram⁺ and Gram⁻ bacteria but not against fungi. The same fed vs unfed *A. hebraeum* syngaglion subtraction cDNA library was used in the identification of a novel type of antimicrobial peptide, named **hebraein** (Table 1).⁶⁴ It consists of 102 amino acids, including six cysteins. However, its pattern of disulfide bridges, anionic character and secondary structure with prevailing α -helices clearly differentiate this AMP from arthropod defensins. The most striking feature of hebraein is its C-terminal histidine-rich domain. Isolated as well as recombinant hebraein were active against Gram⁺ as well as Gram⁻ bacteria (*Staphylococcus aureus* and *E. coli*) and against the yeast *Candida glabrato*, but not against *C. albicans*. The antibacterial activity of the histidine-deficient mutant was significantly reduced especially against *C. glabrato*. The authors also demonstrated that C-terminal histidine-rich fragment alone does not display any intrinsic antimicrobial activity.⁶⁴ At about the same time, a similarly histidine-rich molecule called **microplusin** (Table 1) was purified by RP-HPLC from the hemolymph of *R. microplus*.⁶⁵ Recently, the molecular structure of microplusin has been solved using NMR, leading to a model of its mode of action.⁶⁶ In addition to *M. luteus* and other Gram⁺ bacteria, microplusin was also found to be very efficient against some filamentous fungi like *Aspergillus niger* and very specific against the yeast *Candida neoformans* but not against *C. albicans*.⁶⁶ In contrast to defensins, microplusin does not affect the permeability of the microbial membrane. Instead, microplusin was found to chelate copper ions by binding them most probably to the histidine residues located at both N-terminal (H2) and C-terminal (H74) ends of the molecule (Fig. 2). The bacteriostatic effect of microplusin against *M. luteus* seems to be due to its capacity to sequester copper ions needed for bacterial respiration, a copper-dependent process. The growth of *M. luteus* could be restored by supplementing the medium with copper.⁶⁶

Besides defensin and microplusin,⁶⁵ a third cystein-rich antimicrobial peptide, designated **ixodidid** (Table 1), has been isolated by RP-HPLC from an acidic extract of *R. microplus* hemocytes.⁶⁷ Mature ixodidid possesses ten cysteins forming five internal disulfide bridges.⁶⁷ The ixodidid activity against *M. luteus* was in the submicromolar range and five times less efficient against *E. coli*. The sequence of ixodidid displayed a significant similarity with several serine protease inhibitors from different invertebrates and accordingly the peptide exerted inhibitory activity against elastase and chymotrypsin. Whether the antibacterial activity of ixodidid is due to the proteinase inhibition or to a direct effect on the bacterial membrane remains an unresolved issue.⁶⁷

IMMUNITY IN THE TICK GUT

The tick gut is the primary interface of the tick-pathogen interaction. Thus, the ability of a pathogen to survive in the gut lumen, penetrate the gut epithelia and eventually multiply before reaching the haemocoel are intuitively considered as the determinants of the tick vector competence. Ticks differ from other hematophagous arthropods in that they digest blood intracellularly and the protein-rich nutrients are only gradually absorbed by the gut epithelium.³ Since tick feeding is hardly an aseptic process, there has to be

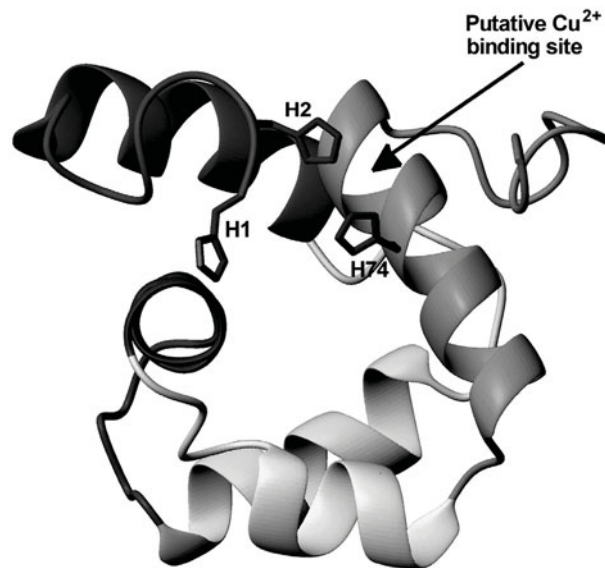


Figure 2. Ribbon representation of the microplusin structure obtained by solution NMR (PDB ID 2knj).⁶⁶ Microplusin consists of five α -helices, having histidine residues located at both N-terminal (H2) and C-terminal (H74) ends of the molecule as putative binding site for Cu^{2+} ions.

an active and efficient defense mechanism preventing bacterial growth in the tick gut lumen. The failure of such a mechanism may lead to high mortality, such as reported for the laboratory colony of the soft tick *O. moubata* accidentally infected via membrane feeding with *C. indologenes*.²² By contrast, the hard tick *I. ricinus* was quite resistant to the per os infection with this bacterium, suggesting that the families Argasidae and Ixodidae may differ also in midgut defense strategies.²²

The pivotal and constitutive role in microbial control in the tick gut of both families is apparently played by the antimicrobial activity of the host **hemoglobin fragments**. Fogaca et al⁶⁸ were the first who demonstrated that a 3,205 Da fragment of the bovine α -hemoglobin purified from the gut contents of *R. microplus* is responsible for activity against several Gram⁺ bacteria and fungi. A similar purification approach, assaying for activity against *S. aureus* or *M. luteus*, led to the identification of antibacterial hemoglobin fragments in the gut lumen of the soft tick *O. moubata*⁶⁹ and the hard tick *D. variabilis*,⁷⁰ respectively. The structural analysis of the bovine α -hemoglobin fragment (Hb 33-61) made it possible to explain how the N- and C-terminus of the molecule contribute to the disruption of the *M. luteus*⁷¹ and *C. albicans*⁷² membrane. The large hemoglobin fragments are generated in the initial phase of hemoglobin digestion, mainly by the action of an aspartic peptidase of the cathepsin D type.⁷³ However, the mechanism of their secretion from the digestive vesicles to the gut contents remains obscure. Interestingly, the hemoglobin-derived antimicrobial fragments seem to have a conserved and universal role in innate immunity since they have been also reported as defense molecules in the human placenta.⁷⁴

Apart from the antimicrobial peptides of host origin, several types of endogenous molecules demonstrably or presumably contributing to immunity within the tick gut have

been described. A c-type **lysozyme** was purified from the gut contents of the soft tick *O. moubata* showing activity against *M. luteus* (Table 1).⁷⁵ The sequence of the *O. moubata* gut lysozyme shared some motives with lysozymes found in the digestive tract of insects or stomachs of ruminants and its expression was strongly up-regulated by blood meals.⁷⁶ The lysozyme of the hard tick *D. variabilis*, originally identified in hemocytes,⁵⁵ was later found to be also markedly expressed in the midgut of this species.⁷⁷ In contrast to the *O. moubata* gene, the expression *D. variabilis* lysozyme mRNA did not increase upon feeding but seemed to be up-regulated in ticks infected *per os* with the causative agent of spotted fever, *Rickettsia montanensis*, by capillary feeding.⁷⁷

Four isoforms of typical cationic **defensin**-A, B, C, D (Table 1) that became up-regulated after blood feeding were described in *O. moubata* ticks by Nakajima et al.^{78,79} Isoforms A, B and C were preferentially expressed in the midgut and isoform D in the fat body. The authors succeeded in purification defensin A from the gut contents by RP-HPLC and demonstrated its activity against *S. aureus*.⁷⁹ The antibacterial assays performed with synthetic *O. moubata* defensin A confirmed its bactericidal activity against a panel of Gram⁺ but not Gram⁻ bacteria. The mode of action of defensin A peptide was demonstrated by the permeabilization of the bacterial membrane.⁸⁰

Of special interest is a report on a defensin-related molecule termed **longicin** (Table 1), expressed mainly in the midgut of the babesial vector *Haemaphysalis longicornis*.⁸¹ Although longicin possesses the typical furin cleavage motif, the authors have shown that the mature protein also contains the N-terminal part of typical tick pro-defensins. Recombinant longicin was active against a variety of fungi, Gram⁺ and Gram⁻ bacteria, including multidrug-resistant strains. Most strikingly, the recombinant protein exerted babesiacidal activity *in vitro*. In an *in vivo* experiment, the inoculation of longicin significantly reduced parasitemia in mice infected with *Babesia microti*. RNAi experiments performed in this work further supported the role of longicin in *Babesia* transmission and regulation of *H. longicornis* vectorial capacity. Since this work points to an exceptional activity of a defensin-related molecule against a protozoan apicomplexan parasite, which is related to the causative agent of malaria, any follow-up studies would be quite significant.

Another typical tick-defensin named HI-gut defensin (Table 1) was found to be specifically expressed in the *H. longicornis* midgut and strongly up-regulated upon intrahaemocoelic injection of LPS.⁸² Defensins predominantly expressed in the midgut tissue were also reported for the genera *Ixodes* (Table 1), namely *I. ricinus*,⁸³ *I. scapularis*⁸⁴ and *I. persulcatus*.⁵⁶ The most recent comparison of the antibacterial activity of synthetic peptides corresponding to the *I. ricinus* defensin, *O. moubata* defensin D and the above mentioned longicin from *H. longicornis*, revealed that all defensins were active against Gram⁺ *S. aureus*. On the other hand, the Lyme disease spirochete *Borrelia garinii* and the symbiotic bacteria *Stenotrophomonas maltophilia* isolated from *I. persulcatus* were resistant to the tick defensins, suggesting an adaptation of these microbes to the tick antimicrobial response.⁸⁵

In contrast to the soft ticks, neither defensins nor lysozymes have been unambiguously proven to be secreted from the midgut epithelium into the gut lumen of hard ticks. An attempt to detect these immune molecules, next to the hemoglobin antimicrobial fragments in the *D. variabilis* gut contents, was not successful.⁷⁰ Defensin and lysozyme transcripts were surprisingly absent in the *D. variabilis* midgut transcriptome.⁸⁶ These antibacterial peptides were also not found among almost 150 proteins identified by a recent extensive proteomic analysis of the midgut from partially fed *R. microplus* females.⁸⁷ Given these

data, the ultimate evidence that defensins and/or lysozymes really contribute to the control of microbial growth in the midgut of hard ticks awaits further experimental verification.

Protease inhibitors may play a role in tick midgut immunity via the inhibition of microbe-specific proteases. Zhou et al reported that the expression of a cysteine protease inhibitor of the cystatin type (Hlcyst-2) (Table 1) in the midgut of *H. longicornis* was increased in response to blood feeding, LPS injection and experimental infection with *Babesia gibsoni*. They have further shown that recombinant cystatin partially limited the growth of *Babesia bovis* in culture.⁸⁸ Another example was provided for a Kunitz-type serine protease inhibitor from *D. variabilis*, referred to as KPI.⁸⁹ In contrast to the majority of related tick Kunitz-type serine protease inhibitors expressed in the salivary glands to control homeostasis in the host, KPI was highly expressed in the midgut in response to feeding and its mRNA level was also induced after *per os* infection with rickettsia. Beside a trypsin inhibitory activity and possible role as an anticoagulant of the imbibed blood, the authors demonstrated that *D. variabilis* KPI limits rickettsial colonization of mouse fibroblasts.⁸⁹

The paradigm that *Drosophila* midgut epithelial immunity efficiently and specifically controls intestinal microbes by the fine-tuned generation and elimination of reactive oxygen species (see refs. 90, 91) raises the question as to whether **redox homeostasis** contributes to tick midgut immunity as well. Ticks defend themselves against the oxidative stress caused by an enormous amount of heme liberated from the digested hemoglobin by complex heme detoxification mechanisms (see ref. 92 for a review). Ticks get rid of the heme mainly by its aggregation into hemosomes—special organelles of digestive cells that are subsequently released to the gut lumen.⁹³ A small portion of heme binds to a heme-binding protein named HeLp,⁵³ which serves both as heme transporter and antioxidant enzyme in the hemolymph (Table 1).^{48,94} Citteli et al demonstrated that catalase, an enzyme responsible for H₂O₂ detoxification, plays an important role in regulation of oxidative stress in the *R. microplus* midgut.⁹⁵ Analysis of the *D. variabilis* midgut transcriptome revealed other putative antioxidant enzymes and radical scavengers, such as superoxide dismutase, glutathione peroxidase, thioredoxin, glutathione S-transferase (GST), selenoprotein M and others.⁸⁶ Interestingly, several of these antioxidant enzymes were reported to be up-regulated in different tick species in response to infection with relevant pathogens. In the hard tick *I. ricinus*, GST and thioredoxin peroxidase were found by the method of subtractive hybridization among the genes induced after *B. burgdorferi*-infected blood meal.⁸³ GST was also identified to be significantly up-regulated in the gut of *D. variabilis* males fed on calf infected with *A. marginale*. RNAi experiments suggested that GST is involved in the initial phase of midgut infection and possibly also in trafficking of the pathogen and its multiplication in the tick salivary glands.⁹⁶ Another study on two midgut specific *D. variabilis* GSTs demonstrated that expression of these enzymes in the tick gut did not respond to *E. coli* haemocoelic injection but was primarily up-regulated by blood feeding.⁹⁷ Several putative antioxidant enzymes have been cloned and partially characterized in different tick species, but their function in tick redox balance or immune response has not yet been demonstrated (Table 1).⁹⁸⁻¹⁰² The acquisition of *B. burgdorferi* by *I. scapularis* was reported to be facilitated in part by Salp25D, a protein homologous to glutathione peroxidases (peroxiredoxins) that are thiol-specific antioxidants.¹⁰³ The authors were able to perform a tissue-specific silencing of Salp25D, which allowed them to demonstrate that Salp25D from salivary glands but not the midgut is important for successful entry of the pathogen into the midgut. Furthermore, they showed that spirochetes exploit the salivary Salp25D to protect themselves against reactive oxygen

intermediates generated by the mammalian neutrophils at the vector-host interface.¹⁰³ This study elegantly illustrates the complexity of the molecular interplay between the host, vector and pathogen.

IMMUNE MOLECULES IN OTHER TICK TISSUES

Beside defensin and microplusin transcripts,¹⁰⁴ only three antibacterial peptides have been thus far identified among the numerous molecules secreted by tick salivary glands (Table 1). Two short peptides of different sequence, designated as Ixosin and Ixosin B were isolated from the salivary glands of the hard tick *Ixodes sinensis*.^{105,106} Both mature Ixosins lack cysteine residues and display a similar antimicrobial activity mainly against *S. aureus* and the yeast *C. albicans* and to a lesser extent against *E. coli*. Recently, a novel salivary antimicrobial peptide named IsAMP has been isolated from the saliva of *I. scapularis*.¹⁰⁷ Interestingly, it was found to belong to a family of secreted 5.3 kDa proteins previously identified to be specifically up-regulated in *B. burgdorferi*-infected nymphs.¹⁰⁸ The mature IsAMP is composed of 47 amino acid residues including six cysteine residues. IsAMP is active against Gram⁺ and Gram⁻ bacteria. In addition to salivary glands, IsAMP is expressed in hemocytes and the fat body but not in the midgut.¹⁰⁷

Esteves et al. detected several antimicrobial activities in the egg homogenate and surface material from the tick *R. microplus*.¹⁰⁹ One of the antimicrobial peptides identified in the egg homogenates was attributed to the previously described microplusin.⁶⁵ Moreover, the regulation of microplusin mRNA transcription seems to be stage-dependent, during ovary and egg development. The authors suggest that microplusin acts as a protective factor of both the female reproductive tract and embryo.¹⁰⁹ In contrast, the compound(s) responsible for antimicrobial activity on the egg surface remain to be identified. A defense role is possibly played also by two subtilisin inhibitors named as BmSI-7 and BmSI-6 (Table 1) purified from the *R. microplus* eggs since they strongly inhibit Pr1 protease from the entomopathogenic fungus *Metarhizium anisopliae*.¹¹⁰

Several transcripts of genes putatively involved in immune response were identified in phagocytic cell line BME26, derived from *R. microplus* embryos.¹¹¹ They comprise e.g., microplusin, hebraein, α_2 -macroglobulin, factor D-like protein, GST, peroxidase and NADPH oxidase. This result suggests that this cell line has the potential to be a valuable tool for further studies on the interaction of tick cells with relevant pathogens, as recently demonstrated for the *A. marginale*.¹¹²

Regulation of Immune Response in Ticks

Although several of the above mentioned immune molecules are induced by microbial challenge, there is virtually nothing known about the regulation of the immune response in ticks. The *I. scapularis* genome contains a number of genes automatically annotated as putative components of the immune sensing and signaling pathway(s),⁹ comprising e.g., putative Toll-like receptors with leucine-rich repeats domains, a nuclear factor κ B (NF- κ B) related to Dorsal and also an ankyrin-repeat containing protein resembling Cactus, a NF- κ B inhibitor. This finding suggests that the Toll signaling pathway is present in ticks. Several components putatively related to the *Drosophila* Imd pathway, such as Caspar and Relish, are also found in the *I. scapularis* genome, but an ortholog of Imd itself seems to be missing.

The only sufficiently characterized tick molecule that plays a role in regulation of immune gene transcription is **subolesin** (Table 1).¹¹³ This molecule (originally named 4D8) was discovered in *I. scapularis* as a tick protective antigen, since immunization with recombinant subolesin effectively protected hosts against infestation and had a deleterious effect on tick survival and further development.¹¹⁴ Recently, it was demonstrated that tick subolesin is an ortholog of insect and vertebrate akirin,¹¹⁵ a highly conserved nuclear protein involved in the transcription of immune genes associated with NF- κ B.¹¹⁶

CONCLUSION

Research on tick innate immunity and the molecular interplay at the tick-pathogen interface is just entering its postgenomic era. Reverse genetics and functional genomics based on RNAi will continue to substantially extend our knowledge that can potentially be exploited to combat tick-borne pathogens. In addition, the introduction of these techniques may make ticks to an additional model for the study of innate immunity, at least for the class Arachnida. Certainly, the main effort will be always targeted to understand how the transmitted pathogens evade the tick immune response. Data from other invertebrate models can be used to fish out putative functional homologues in tick genomes or EST databases and ultimately verify their immune functions. We believe that ticks will prove to be a suitable model system for discovering molecules involved in pathogen recognition, the action of the primordial complement system and regulation of antimicrobial responses, among other aspects of innate immunity. Further research on tick midgut immunity and its relation to blood digestion and/or redox homeostasis may be a key to efficient control of ticks and tick-borne pathogens.

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CHAPTER 9

BEETLE IMMUNITY

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Abstract: Genetic studies have elegantly characterized the innate immune response in *Drosophila melanogaster*. However, these studies have a limited ability to reveal the biochemical mechanisms underlying the innate immune response. To investigate the biochemical basis of how insects recognize invading microbes and how these recognition signals activate the innate immune response, it is necessary to use insects, from which larger amounts of hemolymph can be extracted. Using the larvae from two species of beetle, *Tenebrio molitor* and *Holotrichia diomphalia*, we elucidated the mechanisms underlying pathogenic microbe recognition. In addition, we studied the mechanism of host defense molecule amplification. In particular, we identified several pattern recognition proteins, serine proteases, serpins and antimicrobial peptides and examined how these molecules affect innate immunity.

INTRODUCTION

Beetles (Coleoptera) such as fireflies and bombardier beetles, are one of the most evolutionarily conserved metazoan families of insects. The red flour beetle, *Tribolium castaneum* (Coleoptera, Tenebrionidae), is a common model organism used for developmental, evolution, comparative genomics and pest science. In addition, *Tribolium* has been the subject of studies investigating host interactions with a variety of pathogenic bacteria, sporozoa, cestoda, nematoda, mites and hymenopterous parasites.¹ The *Tribolium* genome sequence has recently been determined by the Human Genome Sequencing Center, Baylor College of Medicine.² This information has made *Tribolium* an excellent model for

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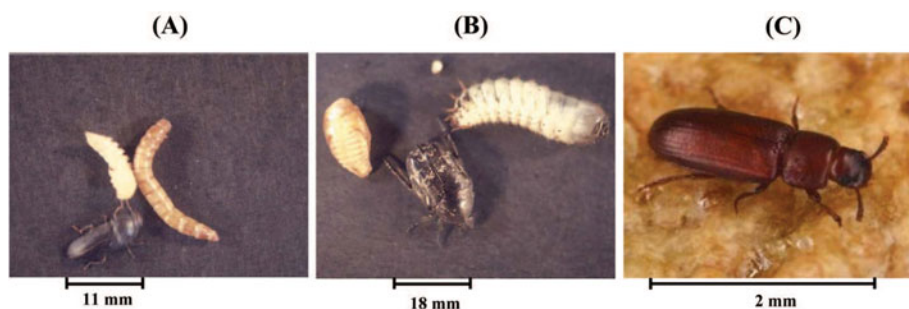


Figure 1. The pictures of beetles. (A) *Tenebrio molitor*; (B) *Holotrichia diomphalia*; (C) *Tribolium castaneum*. To image the real sizes of beetles, measures are inserted. Reproduced with permission from Iwanaga S, Lee BL. J Biochem Mol Biol 2005; 38:128-50, ©2005.³⁶

studying insect immunity. *Tenebrio molitor* (mealworm) and *Holotrichia diomphalia* are from the same family as *Tribolium* and are larger than *Tribolium* (Fig. 1). These insects reproduce rapidly and up to 500 μ l of hemolymph (insect blood) can be extracted from a single larva. Because the *Tenebrio* and *Holotrichia* larvae elicit a high antimicrobial activity when challenged with pathogenic microbes, we used *Tenebrio* and *Holotrichia* larvae as a model system to elucidate the biochemical mechanisms underlying the host defense system.

BIOCHEMICAL CHARACTERIZATION OF PATTERN RECOGNITION PROTEINS IN BEETLES

The innate immune response is a crucial host defense system to defend against microbial infection.³ The ability of a host to distinguish between self and nonself remains a central hallmark of innate immunity.⁴ The pathogenic microbes possess distinct pathogen-associated molecular patterns (PAMPs), such as lipopolysaccharides (LPSs) of Gram-negative bacteria, peptidoglycans (PGs) of Gram-positive bacteria and β -1,3-glucans of fungi.⁵ The recognition of PAMPs is achieved by a group of germ line-encoded receptors and soluble proteins.⁶ The *Drosophila* Toll receptor signaling pathway is responsible for the defense against Gram-positive bacteria and fungi, while the immune deficiency (Imd) pathway is activated primarily in response to Gram-negative bacteria. Both of these pathways lead to the expression of antimicrobial peptides (AMPs) via the activation of NF- κ B-like transcription factors.⁷⁻¹⁰ In insects, bacterial lysine (Lys)-type PGs are recognized by the PG recognition protein-SA (PGRP-SA) and Gram-negative-binding protein 1 (GNBP1).¹¹⁻¹³ These proteins are believed to mediate the activation of a serine protease (SP) cascade and ultimately, the cleavage of Spätzle pro-protein. Upon cleavage, Spätzle serves as a ligand for the Toll receptor and increases the production of AMPs.^{10,14} GNBP3 is required for the detection of fungal cell wall components.¹⁵ These recognition signals are amplified in hemolymph by a proteolytic SP cascade; this protein cascade is conserved in the vertebrate system.^{10,16} The amplification of these recognition signals results in an efficient host defense strategy in insects, which are devoid of an acquired immune system.

The prophenoloxidase (proPO) activation pathway, similar to the vertebrate complement system, is a proteolytic molecular cascade comprised of pattern recognition

proteins and several SPs and their inhibitors. The cascade terminates with the zymogen, proPO.^{17,18} Microbial carbohydrates, such as LPS, PG and β -1,3-glucan, are initially recognized by pattern recognition proteins and induce SP-dependent activation of the proPO system. The proPO-activating enzyme (PPAE) and proPO-activating factor (PPAF), which are similar to the *Drosophila* Easter-type SP, cleave proPO to generate the active enzyme, phenoloxidase (PO).¹⁹⁻²² This enzyme produces compounds that are toxic to microorganisms by the oxidation of phenols and the subsequent formation of melanin. In addition, this enzyme participates in the sclerotization of cuticles, a process that is vital for the survival of insects.²³ Numerous reports have been published about invertebrate proPO and its activation mechanism.¹⁸ A key question regarding the proPO system is: how do pattern recognition molecules activate this system in response to microbial infection?

Although information about the function of distinct PAMP recognition proteins, such as PGRP-SA and GGBP3, from genetic studies in *Drosophila* gives insight into the Toll cascade, the activation mechanism of the extracellular SP cascade in the Toll pathway has not been clearly elucidated in *Drosophila*. Because *Tenebrio* and *Holotrichia* larvae have a high antimicrobial activity against bacteria and fungi,²⁴⁻²⁸ we sought to biochemically determine the molecular activation mechanism of the Toll cascade in beetles. Initially, we purified and cloned GGBP3²⁹ from *Tenebrio* hemolymph using a 1,3- β -glucan affinity column. The purified protein bound to β -1,3-glucan but not to bacterial PG. Subsequent molecular cloning revealed that GGBP3 contains a region with a sequence similar to bacterial glucanases.²⁹ Interestingly, two catalytically important residues in the glucanases had been replaced with nonhomologous amino acids in *Tenebrio* GGBP3, suggesting that GGBP3 had evolved from an ancestral glucanase gene but retained only the ability to recognize β -1,3-glucan. Next, we purified and cloned the PGRP proteins,^{30,31} termed *Holotrichia* PGRP-SA³⁰ and *Tenebrio* PGRP-SA.³¹ The purified 20-kDa and 17-kDa *Holotrichia* proteins (referred to as PGRP-1 and PGRP-2, respectively) had a high sequence identity to *Drosophila* PGRP-SA. We purified *Tenebrio* PGRP-SA using a novel synthetic Lys-type PG peptide [(GlcNAc-Mur-Nac-L-Ala-D-isoGln-L-Lys-D-Ala)₂]. This synthetic Lys-type PG fragment is a competitive inhibitor of PG-dependent proPO cascade and thus, did not activate the proPO system. The purified *Tenebrio* PGRP-SA recognized both Lys-type PG and diaminopimel:c(DAP)-type PG. In vitro reconstitution experiments demonstrated that *Tenebrio* PGRP-SA is a common recognition molecule of Lys-PG- and DAP-PG-dependent proPO cascades.³¹

We purified a novel 40-kDa LPS recognizing protein, termed LPS recognition protein (LRP), from the cell-free plasma of *H. diomphalia* larvae.³² LRP exhibited agglutinating activity in *Escherichia coli*, but not in *Staphylococcus aureus* and *Candida albicans*. The *E. coli* agglutinating activity was preferentially inhibited by the rough-type LPS, which contains an oligosaccharide core. The LRP consisted of 317 amino acid residues and six repeats of an epidermal growth factor (EGF)-like domain. Recombinant LRP also had *E. coli* agglutination activity in vitro and was able to neutralize LPS by the inhibition of LPS-induced interleukin 6 (IL-6) production in mouse bone marrow mast cells. Furthermore, the *E. coli* coated with the purified LRP were cleared more rapidly in the *Holotrichia* larvae than in the noncoated *E. coli*, indicating that this protein participates in the clearance of *E. coli*. Three N-terminal EGF-like domains of LRP, but not the three C-terminal EGF-like domains, were involved in the LPS-binding activity. In conclusion, the data suggest that LRP is a pattern recognition protein for LPS and is a protein central to the innate immune response.

SERINE PROTEASE ZYMOGENS REGULATE THE proPO CASCADE IN BEETLES

Initially, we identified and cloned two proPO-activating factors, termed PPAF-I and PPAF-II. These proteins directly activate proPO in the *Holotrichia* hemolymph.^{19,33,34} The overall structure of PPAF-I is very similar to that of the *Drosophila* Easter SP (42.9% identity),³⁵ an essential SP zymogen in pattern formation during embryonic development. The locations of disulfide linkages in the prosegment of PPAF-I are similar to those of the proclotting enzyme from *Tachypleus*.³⁶ Furthermore, [³H]diisopropylphosphate (iPr2P)-labeled PPAF-I was produced in the crude preparation of the PPAF-I zymogen when incubated with LPS or β -1,3-glucan. However, [³H]iPr2P-labeled PPAF-I was not produced under the same conditions in the absence of these microbial polysaccharides. These results indicate that microbial polysaccharide-dependent activation of the proPO system in *Holotrichia* larvae activates the PPAF-I zymogen. The structure of the PPAF-II is similar to that of the masquerade protein,³⁷ a SP homologue expressed during embryogenesis and *Drosophila* larval and pupal development. The 45-kDa PPAF-II protein contained a trypsin-like SP domain at the C-terminus. PPAF-II differs from trypsin in that there is a substitution of a serine residue in the active site triad to a glycine and this protein has a disulfide-knotted clip domain at the N-terminus. Using in vitro reconstitution experiments, we demonstrated that PPAF-II, active PPAF-I and proPO were necessary for showing PO activity in the *Holotrichia* proPO system, suggesting that PPAF-II and PPAF-I are essential factors in the *Holotrichia* proPO cascade.

The crystal structure of PPAF-II (Fig. 2) revealed that the clip domain adopts a novel fold containing a central cleft, which is distinct from the structures of defensins that have a similar arrangement of cysteine residues.³⁸ The PPAF-II forms a homo-oligomer upon cleavage by the upstream SP and the central cleft in the clip domain of PPAF-II functions as a module for binding PO. Similarly, the clip domain in the catalytically active Easter-type SP plays an essential role in the rapid activation of the protease domain.³⁸ In addition, we

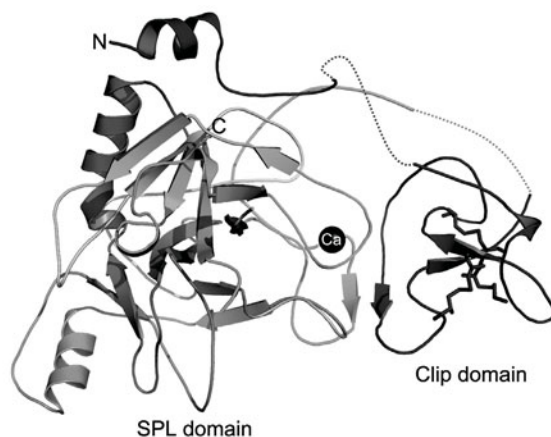


Figure 2. Overall structure of *Holotrichia* PPAF-II. A ribbon representation of PPAF-II structure. The clip domain and the PPAF-II family-specific N-terminal segment are shown in blue. Bound calcium is in black. The disordered regions are shown in dashed lines. A color version of this image is available at www.landesbioscience.com/curie.

previously reported that the crystal structure of the SP domain of the Easter-type SP is structurally similar to PPAF-I. This analysis revealed several important structural features including a Ca^{2+} binding region, an additional loop with unique disulfide linkages, a canyon-like deep active site and an exposed activation loop.³⁹ We subsequently elucidated the function of the bound Ca^{2+} and discovered that the activation loop is susceptible to proteolytic degradation, which did not involve the clip domain.

We purified and cloned the 27-kDa PPAF-III and two proPOs from the plasma of *Holotrichia* larvae.⁴⁰ The *PPAF-III* gene encoded an Easter-type SP zymogen with a mass of 40 kDa. The active form of the 27-kDa PPAF-III specifically cleaved a 55-kDa proPPAF-II in a Ca^{2+} -independent manner to generate a 45-kDa PPAF-II. Furthermore, *Holotrichia* proPO-I and proPO-II have been characterized and the activation-dependent structural changes were examined using in vitro reconstitution experiments. When the proPOs were incubated with PPAF-I, the 79-kDa proPOs were converted to 76-kDa proPOs, which did not exhibit any PO activity. However, when the proPOs were incubated simultaneously with PPAF-I, proPPAF-II and PPAF-III in the presence of Ca^{2+} , a 60-kDa protein (PO-1) with PO activity was detected in addition to the 76-kDa proPO-II protein. These results indicate that the conversion of *Holotrichia* proPOs to enzymatically active PO was accomplished by PPAF-I, PPAF-II and PPAF-III through a two-step Ca^{2+} -dependent proteolytic process. Finally, we purified and cloned the *Tenebrio* SP homologue-1 (SPH1) and proPO⁴¹ from the *Tenebrio* hemolymph.⁴² Using an in vitro reconstitution experiment, we demonstrated that the cleavage of the 55-kDa SPH1 zymogen by an unidentified SP was necessary for PO activity and that SPH1 functioned as a PPAF.⁴²

A MODULAR SERINE PROTEASE IS THE IMMEDIATE DOWNSTREAM PROTEASE OF THE PGRP-SA/GNBP1 COMPLEX

Drosophila genetic studies suggest that PGRP-SA and GNBP1 form a complex that recruits SPs and activates the Toll signaling cascade.⁴³ However, the identification of the SPs immediately downstream of the PGRP-SA/GNBP1 complex is not easily performed in *Drosophila* due to protein redundancy. Since we observed a high antimicrobial activity against a challenge with Gram-positive bacteria or fungi in *Tenebrio* larvae, we hypothesized that *Tenebrio* has the essential components necessary for the activation of the Toll signaling cascade. Therefore, we purified the immediate downstream SP of the PGRP-SA/GNBP1 complex in *Tenebrio* using a biochemical approach.¹³

We hypothesize that GNBP1 and an SP would be recruited to the PG/PGRP-SA complex when the soluble Lys-type PG/*Tenebrio* PGRP-SA complex was incubated with PGRP-SA-depleted *Tenebrio* hemolymph. Indeed, the *Tenebrio* PGRP-SA/Lys-type PG complex enriched a 50-kDa protein and a 35-kDa protein. The sequencing of the N-terminal residues identified the 50-kDa protein as *Tenebrio* GNBP1 and the 35-kDa protein as *Tenebrio* modular serine protease (MSP). However, these proteins do not contain a clip domain, which is commonly found in proteases upstream of the Toll and proPO cascades.³⁸ GNBP1 physically interacts with PGRP-SA and activates the Toll pathway in *Drosophila*.⁴³ However, an interaction between GNBP1 and PGRP-SA has not been observed in vitro. Our observation suggests that the binding of PGRP-SA to PG enhanced the interaction between PGRP-SA and GNBP1 and the active form of MSP was recruited to the Lys-type PG/PGRP-SA/GNBP1 complex. Subsequent cDNA cloning

demonstrated that *Tenebrio* MSP contains four low-density lipoprotein receptor A repeat (LDL) domains, one complement control protein (CCP) domain and a trypsin-like SP domain.⁴⁴ We demonstrated that the Lys-type PG/PGRP-SA complex recruited GGBP1 and MSP and proposed that this Lys-type PG/PGRP-SA/GGBP1/MSP complex initiates the SP cascade in the Toll and proPO pathways in beetles.

Manduca sexta hemolymph protease 14 (Ms-HP-14), which contains a domain arrangement similar to that of the *Tenebrio* MSP, initiates the activation of the proPO system.^{45,46} In addition, Lemaitre et al⁴⁷ reported that the *Drosophila* MSP plays an essential role in the activation of the Toll pathway by Gram-positive bacteria and fungi in *Drosophila*. Their analysis showed that the MSP integrates signals that originate from the circulating recognition molecules, GGBP3 and PGRP-SA and relays them to the Grass-Spätzle processing enzyme (SPE) extracellular pathway, which is upstream SPs of the Toll receptor.

THREE SERINE PROTEASE ZYMOGENS ARE INVOLVED IN LYS-TYPE PG-DEPENDENT ACTIVATION OF THE TOLL SIGNALING CASCADE

Although we found that MSP is recruited into the Lys-type PG recognition complex, the lack of information regarding the identity of the immediate downstream factor(s) of MSP limits our knowledge of the molecular details of PG recognition and the involvement of SPs in the Toll signaling pathway. Therefore, the identification and determination of the biological functions of SPs involved in the *Tenebrio* Toll signaling pathway are essential to the elucidation of the host defense system. SPE, a terminal SP that converts Spätzle pro-protein into a processed form capable of binding to the Toll receptor, was identified in *Drosophila*.⁴⁸ We purified the immediately downstream SPs of *Tenebrio* MSP and *Tenebrio* SPE in order to obtain biochemical information regarding the Lys-type PG recognition signal-dependent activation of the Toll cascade.

We purified three SPs, *Tenebrio* MSP and a 41-kDa and 44-kDa protein, to homogeneity by column chromatography and cloned their cDNAs (Fig. 3).⁴⁴ We sought to identify the SPE molecule in *Tenebrio*. Because the 44-kDa zymogen protein is similar to *Drosophila* SPE and Easter, we initially studied the active form of the 44-kDa protease. Since the cDNAs of *Tenebrio* Spätzle and the Toll proteins have not been determined, we expressed the *Tribolium* Spätzle pro-protein and the Toll ectodomain in a baculovirus system (Fig. 3). To address whether the purified active 44-kDa protease cleaves *Tribolium* Spätzle pro-protein in vitro, we incubated the Spätzle pro-protein with trypsin, the active forms of *Tenebrio* MSP and the 41-kDa and 44-kDa proteases. Only the active 44-kDa protease cleaved the Spätzle pro-protein. Under the same conditions, trypsin cleaved the Spätzle pro-protein nonspecifically and the active forms of *Tenebrio* MSP and the 41-kDa protease did not cleave the Spätzle pro-protein. In addition, we determined the N-terminal amino acid sequence of the cleaved 24-kDa Spätzle pro-protein to be: Phe-Asn-Thr-Asp-Glu-Met-Ser-Leu. This sequence is identical to residues 125-132 of the *Tribolium* Spätzle protein. Therefore, we named the 44-kDa protease as *Tenebrio* SPE.

We next tried to identify the upstream activator of *Tenebrio* SPE. Because *Tenebrio* SPE has a trypsin cleavage site, we hypothesized that the upstream SP is a trypsin-like SP. As mentioned above, the MSP and the 41-kDa zymogens were identified as chymotrypsin-like and trypsin-like SPs, respectively, suggesting that the active form of the 41-kDa protease may cleave the SPE zymogen. To test this hypothesis, we incubated the active form of

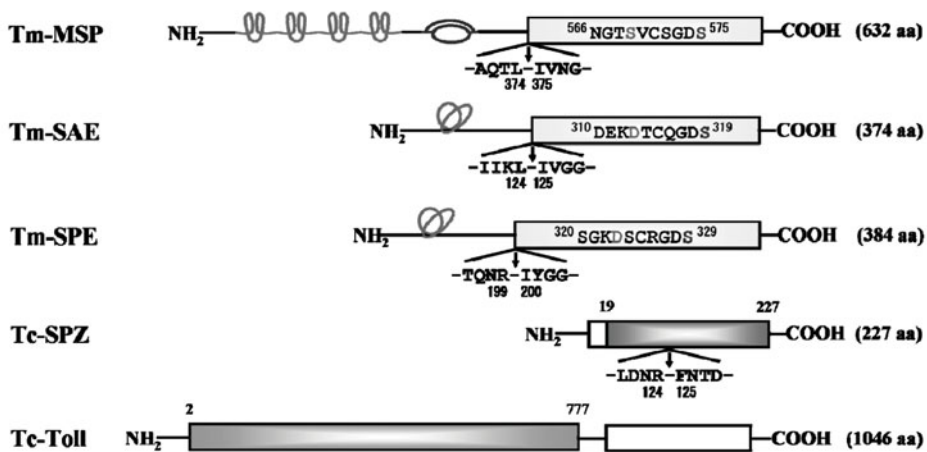


Figure 3. The domain organizations based on deduced amino acid sequences of cDNAs of *T. molitor*-MSP (Tm-MSP), Tm-41 kDa SP (Tm-SAE), Tm-44 kDa SP (Tm-SPE) and *T. castaneum*-Spätzle (TC-SPZ) and Tc-ecotodomain Toll. *Rabbit ears*, *half-double circle* and *rectangular symbols* indicate the domains of LDL, CCP and SP domains of MSP, SAE and SPE, respectively. *Moon symbols* indicate the clip domain of SPs. *Arrows* represent the cleavage sites of SP zymogens during activation. The *red* and *blue* residues in the boxes indicate the specificity-related residue and catalytic triad Ser residue, respectively. Reproduced from Kim CH et al. J Biol Chem 2008; 283:7599-07,⁴⁴ ©2008, with permission from the American Society for Biochemistry and Molecular Biology. A color version of this image is available at www.landesbioscience.com/curie.

41-kDa protein with the purified SPE zymogen. As expected, the SPE zymogen was hydrolyzed into a 35-kDa SP domain and a 15-kDa clip domain. The 35-kDa band was identified as the SP domain of SPE. Therefore, we designated the 41-kDa protease as *Tenebrio* SPE-activating enzyme (SAE).

Because the sequence of the cleavage site in the SAE zymogen is Leu-124-Ile-125, the upstream SP of SAE is probably similar to chymotrypsin. This result suggests that the SAE zymogen is cleaved by the MSP. Therefore, activated MSP was incubated with the recombinant SAE zymogen. The SAE zymogen was hydrolyzed into a 35-kDa SP domain and an 11-kDa clip domain. The N-terminal amino acid sequence of the 35-kDa band was Ile-Val-Gly-Gly-Thr-Asn. This sequence is identical to the amino acid sequence of the SAE zymogen from Ile-125 to Asn-130, demonstrating that the MSP induced a limited proteolytic cleavage between the clip domain and the catalytic SP domain of the SAE zymogen. Thus, the SAE protease is an immediate downstream target of MSP.

Finally, to confirm whether the Spätzle pro-protein is cleaved, Lys-type-PG, PGRP-SA, GNBPI, zymogens of MSP, SAE, SPE and pro-Spätzle were incubated together in the presence of Ca^{2+} and the cleavage of pro-Spätzle was detected by western blot analysis. As predicted, a 14-kDa band corresponding to cleaved Spätzle was observed (Fig. 4, column 9). However, if any one of the components was omitted from the incubation mixture, no cleavage of the pro-Spätzle occurred (columns 2-8). In conclusion, these experiments demonstrated that the PGRP-SA/GNBPI-mediated Lys-type PG recognition signal is transferred by three different SPs; the initiating enzyme is the 82-kDa chymotrypsin-like MSP and the other two enzymes are the 41-kDa SAE and 44-kDa SPE clip domain-containing trypsin-like SPs.

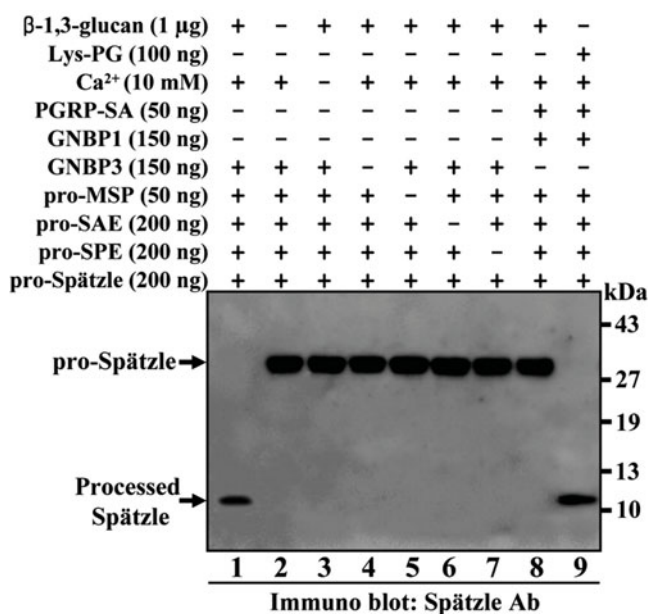


Figure 4. In vitro reconstitution experiments for the activation of pro-Spätzle by β -1,3-glucan and Lys-type PG. The mixture of *Tenebrio* pro-MSP, GNBP3, pro-MSP, pro-SAE, pro-SPE and *Tribolium* pro-Spätzle in the presence of β -1,3-glucan and Ca²⁺ was incubated for 60 min and analyzed by Western blotting with an affinity purified anti-Spätzle antibody (lane 1). The 30-kDa pro-Spätzle and the 12-kDa-processed Spätzle are indicated with arrows. As a control, when eight components, such as Lys-type PG/PGRP-SA/GNBP1/MSP/SAE/SPE/Spätzle were incubated together, the cleaved 12-kDa Spätzle was generated (lane 9). In the absence of any one of these components, pro-Spätzle was not converted to the processed Spätzle (lanes 2-8). Reproduced from Roh KB et al. J Biol Chem 2009; 284:19474-81,⁴⁹ ©2009, with permission from the American Society for Biochemistry and Molecular Biology.

IDENTIFICATION OF THE ESSENTIAL COMPONENTS REGULATING THE β -1,3-GLUCAN-DEPENDENT TOLL SIGNALING CASCADE

We identified three SPs that are involved in the Lys-type PG-dependent Toll signaling pathway. Because the activation mechanism of Toll pathway by β -1,3-glucan is not clearly characterized in insects, we aimed to identify the downstream SPs of GNBP3, a receptor for β -1,3-glucan from fungi, and their order of activation.⁴⁹ To identify the immediate downstream molecule(s) that is recruited by the β -1,3-glucan/GNBP3 complex, the insoluble β -1,3-glucan and the purified *Tenebrio* GNBP3 complex were incubated with the hemolymph fraction. A 35-kDa band was enriched from the GNBP3/insoluble β -1,3-glucan complex but not from the insoluble β -1,3-glucan lacking GNBP3. The N-terminal amino acid sequence of the 35-kDa protein was identical to the catalytic SP domain of *Tenebrio* MSP, suggesting that the MSP was recruited to the β -1,3-glucan/GNBP3 complex. *Tenebrio* MSP is an apical SP that is activated downstream of PGRP-SA/GNBP1 in response to Lys-type PG treatment.¹³

To confirm that the 35-kDa band was generated by *Tenebrio* pro-MSP in the presence of β -1,3-glucan, we performed western blot analysis using an MSP antibody

on hemolymph that was incubated with β -1,3-glucan and GNB3. *Tenebrio* pro-MSP in the hemolymph was not activated by GNB3 but was processed to the 35-kDa active form of MSP when both GNB3 and β -1,3-glucan were added to the hemolymph. These results suggest that the pro-MSP is processed into its active 35-kDa form in the presence of GNB3 and that β -1,3-glucan and MSP binds to the β -1,3-glucan/GNB3 complex. These data indicate that MSP is an apical SP that is immediately downstream of the β -1,3-glucan/GNB3 complex.

Because the *Tenebrio* MSP zymogen is activated in the presence of the β -1,3-glucan/GNB3 complex and the Lys-type PGN/PGRP-SA/GNB1 complex, we hypothesized that the SPs downstream of the β -1,3-glucan/GNB3 complex are identical to *Tenebrio* SAE and SPE, which are activated in response to Lys-type PG treatment.⁴⁴ To address this hypothesis, we performed an in vitro reconstitution experiment using five purified proteins: GNB3, pro-MSP, pro-SAE, pro-SPE and pro-Spätzle. A western blot analysis revealed that the processed Spätzle was generated upon incubation of the five proteins with β -1,3-glucan and Ca^{2+} (Fig. 4, column 1). The elimination of any of the components resulted in the loss of pro-Spätzle cleavage (columns 2-8). Processing of pro-Spätzle to the processed Spätzle was also observed when PGRP-SA, GNB1, MSP, SAE, SPE and Spätzle were incubated with Lys-type PG and Ca^{2+} . These results clearly demonstrate that GNB3, in the presence of β -1,3-glucan, activates a three-step proteolytic cascade involving MSP, SAE and SPE. The cascade activation results in the formation of processed Spätzle, which functions as a ligand for the Toll receptor. In addition, our data demonstrate that the β -1,3-glucan and Lys-type PG recognition signals share a three-step proteolytic cascade that relays their signals to the Toll receptor (Fig. 5).

Although we have provided biochemical evidence elucidating the mechanism by which the β -1,3-glucan recognition signal is transferred to the Toll receptor, we have not demonstrated whether this SP cascade is present in vivo. We hypothesized that if this cascade is present in vivo, the same AMP(s) will be produced in the insect hemolymph when the pathway molecules are injected into the *Tenebrio* larvae. Furthermore, we anticipate that AMP(s) expression will increase upon injection of Lys-type PG. To address this hypothesis, we injected four molecules, β -1,3-glucan, Lys-type PG, activated SAE and processed Spätzle, into the *Tenebrio* larvae. The hemolymph samples had high antimicrobial activities against *S. aureus*, *E. coli* and *S. cerevisiae*. We purified two AMPs, tenecin 1 and tenecin 2, by column chromatography from four hemolymph samples. Tenecin 1 had a bactericidal activity against Gram-positive bacteria and was previously identified by our group.²⁵ The amino acid sequence of tenecin 1 and its disulfide bond arrangement are similar to the defensin protein from *Drosophila*.⁵⁰ Tenecin 2 is highly homologous to the antimicrobial peptide, coleopteracin,⁵¹ which was purified from the coleopteran insect, *Zophobas atratus*. Tenecin 2 has a high sequence identity with coleopteracin and holotricin 2 (65% and 36%, respectively).²⁴ Holotricin 2, previously identified by our group, is an inducible antibacterial peptide purified from the *Holotrichia* larvae. Tenecin 2 had a bactericidal activity against *E. coli* and *S. cerevisiae*. These results demonstrated that tenecin 1 and tenecin 2 are induced by treatment with β -1,3-glucan, Lys-type PG, SAE and Spätzle. These results support the notion that β -1,3-glucan and Lys-type PG activate Toll receptors by the same three-step proteolytic cascade, which results in the production of tenecin 1 and 2 (Fig. 5).

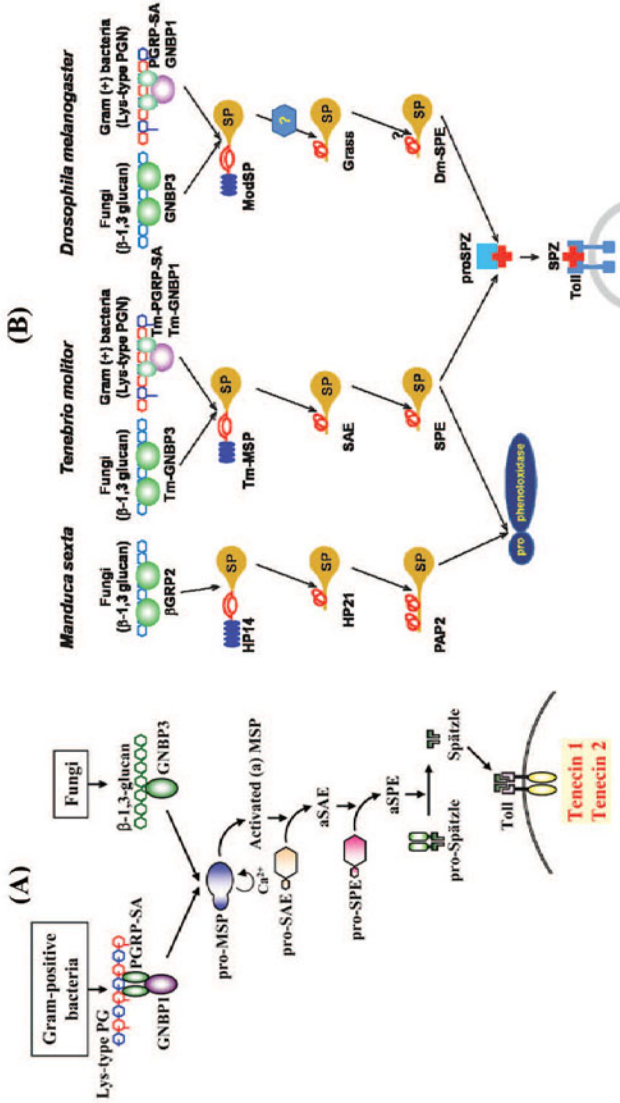


Figure 5. A model of the serine protease cascades regulating the Toll pathway in *Tenebrio* (A) and in other insects (B). A) When the Lys-type-PG of Gram-positive bacteria or β-1,3-glucan of fungi is exposed to the insect, *Tenebrio* PGRP-SA/GNBP1 complex or GNBPs binds to Lys-type PG or β-1,3-glucan, respectively and then recruits pro-MSP. In the presence of Ca²⁺, the Lys-type PG/PGRP-SA/GNBP1 or β-1,3-glucan/GNBP3 complex induces activation of pro-MSP to activated MSP (aMSP). The aMSP converts pro-SAE to activated SAE (aSAE), which subsequently converts the pro-SPE to activated SPE (aSPE). The aSPE cleaves pro-Spätzle to processed Spätzle leading to the production of AMPs such as tenecin 1 and tenecin 2 via the Toll receptor. Reproduced from Roh KB et al. J Biol Chem 2009; 284:19474-81. ©2009, with permission from the American Society for Biochemistry and Molecular Biology. B) Biochemical studies performed in *T. molitor* and *M. sexta* and genetic analysis in *D. melanogaster*, reveal striking similarities in the mechanisms underlying SP activation by pattern recognition proteins. In *M. sexta*, HP-14 is converted to a 2-chain active form in the presence of β-1,3-glucan and the β-1,3-glucan recognition protein 2 (β-GRP2, a protein related to GNBPs).^{46,69} Binding of β-1,3-glucan and β-GRP2 triggers the autoactivation of proHP14 into HP14 that then processes HP21. In turn, HP21 cleaves proPAP-2 into PAP2 that finally activates proPO. Hence, the mechanism of activation of HP14 in response to glucan is highly reminiscent of the mode of activation of *Tenebrio*-MSP, with the exception that HP14 can be activated upon direct binding to PG during bacterial infection.⁶⁹ This suggests that the PGRP-SA/GNBP1 complex involved in sensing Gram-positive bacteria has been added on to a more ancient cascade core during evolution.

THREE PROTEASE-SERPIN COMPLEXES COOPERATIVELY REGULATE THE INNATE IMMUNE RESPONSES IN BEETLES

The serpins belong to a superfamily of SP inhibitors and act as suicide substrates by binding covalently to their target proteases.⁵² Serpins are known to regulate various physiological processes and molecular defense systems in mammals.⁵³ To date, the four *Drosophila* serpins involved in innate immunity, SPN43Ac, SPN27A, SPN77Ba and SPN28D, have been extensively analyzed using a genetic approach. *SPN43Ac* mutant flies accumulate cleaved Spätzle, resulting in the constitutive activation of the Toll pathway and expression of AMPs.¹⁴ *SPN27A* and *SPN28D* are known to regulate the Toll pathway during early development⁵⁴⁻⁵⁶ and are involved in melanin biosynthesis.^{57,58} Another serpin, *SPN77Ba*, was identified as a negative regulator of melanization in the respiratory system (the trachea) in *Drosophila*.⁵⁹ Six serpins have been identified in *M. sexta* and characterized as negative regulators of the proPO cascade.⁶⁰ These serpins are thought to regulate the Toll and proPO cascades. However, the molecular identity of the serpin target SPs and the biochemical regulatory mechanisms of these serpins in relation to the Toll and proPO cascades are not clearly understood. Because we have already identified three SPs that directly activate the *Tenebrio* Toll cascade, we used the *Tenebrio* model system to identify and characterize the novel serpin targets. Recently, we identified three novel serpin-protease pairs that negatively regulate the Toll and proPO cascades.⁶¹

A hallmark of serpin-dependent SP inhibition is the formation of a covalently bonded SP-serpin complex that is visualized as a high-molecular-weight band by SDS-PAGE.⁶² The identity of this band is easily confirmed by western blot analysis using SP antibodies.⁶³ Recently, we obtained the active forms of three *Tenebrio* Toll cascade-activating SPs and raised polyclonal antibodies to these proteins.⁴⁴ These tools were used to monitor the generation of SP-serpin complexes and to purify the serpins. Three novel serpins with a molecular mass of 40 kDa, 55 kDa and 48 kDa (termed *Tenebrio* SPN40, SPN55 and SPN48, respectively) were purified and formed complexes with MSP, SAE and SPE, respectively. To examine the specificity of each serpin for its protease target in vitro, we incubated each purified serpin with the SPs and performed a western blot analysis (Fig. 6). As expected, SPN40 formed a complex with the MSP (lane 2 in Fig. 6B); it did not form a complex with SAE or SPE (lanes 3 and 4). In addition, SPN55 and SPN48 formed complexes with SAE and SPE, respectively (lanes 7 and 12). As a control, purified recombinant *Tenebrio* SPN1, which has a high sequence identity (37%) with *Drosophila* SPN1,⁶⁴ was incubated with the proteases under the same conditions. SPN1 did not bind to the SPs (lanes 14-16). These results suggest that the three purified serpins bind specifically to the Toll cascade-activating SPs. We then cloned the serpin genes and expressed the recombinant serpins with *Tenebrio* SPN1 in *E. coli*. *Tenebrio* SPN40, SPN55, SPN48 and SPN1 consisted of 392, 461, 389 and 394 amino acid residues, respectively. A sequence comparison of the four serpins determined that these proteins share a reactive center loop (RCL) domain that acts as bait for trapping the target SP. This RCL domain contains a putative cleavage site (between the P1 and P1' residues) that is targeted by the proteases during complex formation (Fig. 7).

SAE and SPE cleave after Arg residues in the SPE zymogen (Arg199-Ile200) and pro-Spätzle (Arg124-Phe125) during the activation of the *Tenebrio* Toll cascade, respectively.⁴⁴ Therefore, we hypothesized that SPN55 and SPN48 might have an Arg residue in their P1 position. However, SPN55 was cleaved between the putative P2 Tyr and P1 Met residues by SAE; the cleavage site of SPN48, which is targeted by SPE,

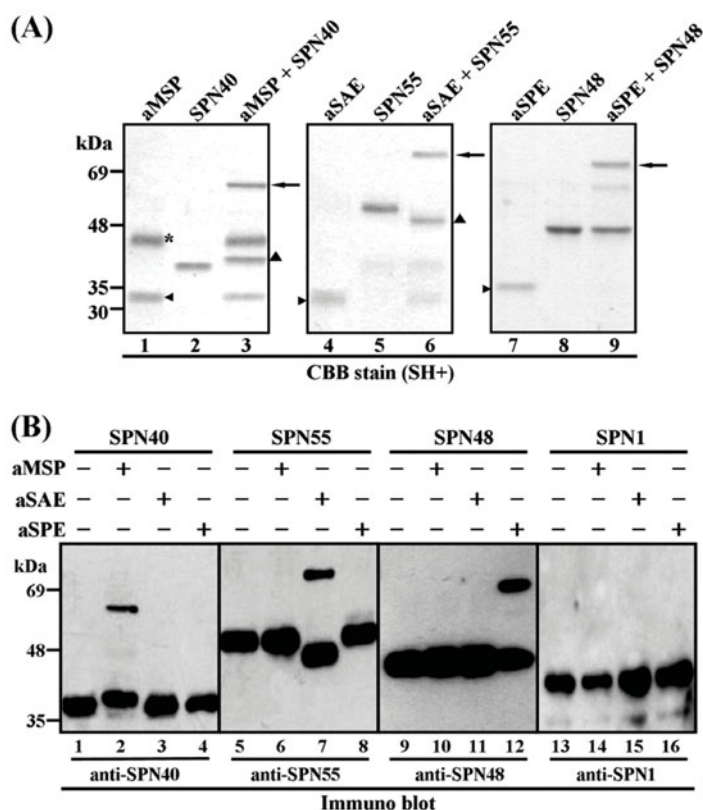


Figure 6. Three *Tenebrio* serpins formed specific complexes with their target SPs. (A) The specific SP-serpin complexes were observed using SDS-PAGE. The purified activated Toll cascade-activating SPs (500 ng, lanes 1, 4 and 7) and purified serpins (1 μ g, lanes 2, 5 and 8) were incubated for 1 h at 30°C and then analyzed by SDS-PAGE under reducing conditions. The gels were stained with a solution containing 0.1% Coomassie Brilliant Blue (CBB) R-250. Arrows in lanes 3, 6 and 9 indicate the SP-serpin complex. Star and arrowheads indicate the N-terminal domain of MSP and the C-terminal catalytic SP domains, respectively. Triangles indicate cleaved SPN40 and SPN55. (B) Each purified serpin (20 pmol), including SPN40 (lanes 1-4), SPN55 (lanes 5-8), SPN48 (lanes 9-12) and SPN1 as a control (lanes 13-16), was incubated with a Toll cascade-activating SP (5 pmol), MSP, SAE, or SPE, under the same conditions as in A. The reaction mixtures were analyzed by immunoblotting to visualize the SP-serpin complexes. Reproduced from Jiang R et al. *J Biol Chem* 2009; 284:35652-58,⁶¹ ©2009, with permission from the American Society for Biochemistry and Molecular Biology.

was between the putative P1 Glu and P2 Met residues. This unusual P1 specificity of the Arg-targeting SPs is similar to several mammalian serpins.^{52,65,66} For example, kallistatin, a serpin that inhibits tissue kallikrein, has a P1 Phe residue, despite the fact that kallikrein targets Arg-specific substrates.⁶⁶ In addition, the protein Z-dependent protease inhibitor, which has a P1 Tyr residue, is a specific inhibitor of the membrane bound factor Xa, an Arg-specific SP.⁵² These unexpected findings provide more evidence supporting the specificity and tight regulation of the Toll cascade by serpins in insects. These proteins colocalize with various digestive SPs in the insect circulatory system. The combined activity of the three serpins described in our study results in the enhancement of the Toll

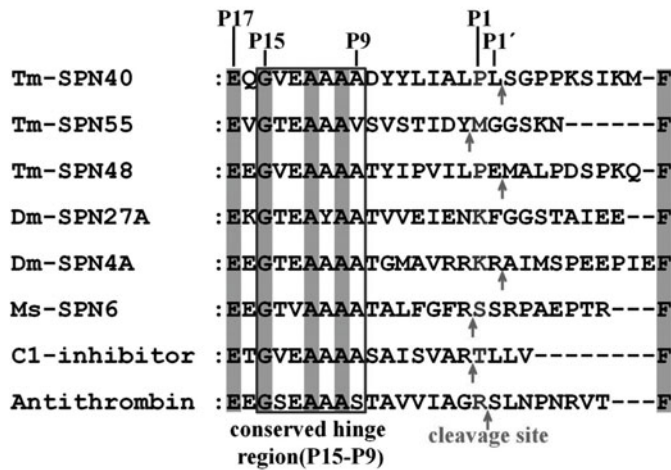


Figure 7. Sequence alignment of the putative and experimentally determined cleavage sites (P1-P1') in the serpins RCL domains. *Blue boxes* indicate the hinge region (P15-P9) of the RCL domains. *Blue highlighted residues* are the putative P1 sites cleaved by target SPs. *Red arrows* indicate the experimentally determined cleavage sites of Dm-SPN4,⁷⁰ Ms-SPN6,⁷¹ human antithrombin,⁷² and C1 inhibitor⁷³ by their target SPs. *Tm*-, *Dm*- and *Ms*- are abbreviations for *T. molitor*, *D. melanogaster* and *M. sexta*, respectively. Reproduced from Jiang R et al. J Biol Chem 2009; 284:35652-58,⁶¹ ©2009, with permission from the American Society for Biochemistry and Molecular Biology. A color version of this image is available at www.landesbioscience.com/curie.

signaling cascade in a specific and a tightly-regulated manner. Our study also provides novel insights into the regulation of pathogen recognition mechanisms by the innate immune response.

When invading pathogens are recognized by the soluble pattern recognition proteins in the hemolymph, SP zymogens, which are positioned upstream of the Toll cascade, are sequentially activated and amplify the pathogen recognition signals.^{6,44,49} We hypothesized that serpins, which are present in low abundance in un-stimulated conditions, may act as a negative feedback inhibitor coincident with Toll cascade activation. To address this hypothesis, we injected β -1,3-glucan, pro-Spätzle or processed Spätzle, a ligand of the Toll receptor, into the *Tenebrio* larvae and determined serpin expression by western blot analysis (Fig. 8). As expected, the levels of SPN40 and SPN55, but not SPN48 or SPN1, were dramatically increased 24 h after injection of β -1,3-glucan and processed Spätzle. These results support the hypothesis; the induction of SPN40 and SPN55 expression may decrease MSP and SAE activation in order to regulate the Toll cascade.

A balance between activation and inhibition of the SP-mediated innate immune response must be maintained to avoid damage to the host.⁶⁷ In our study, we identified three serpins that target three Toll cascade-activating SPs, which act as negative regulators of Toll signaling and melanin synthesis. This is the first functional study of the SP-serpin pairs, which directly regulate the pattern recognition protein-dependent Toll signaling cascade. Surprisingly, each protease in the proteolytic cascade is selectively regulated by a serpin inhibitor. It had previously been hypothesized that these cascades were regulated by a single “bottleneck” protease. There is increasing evidence, in both mammals and *Drosophila*, that the control of proteolytic signaling cascades is

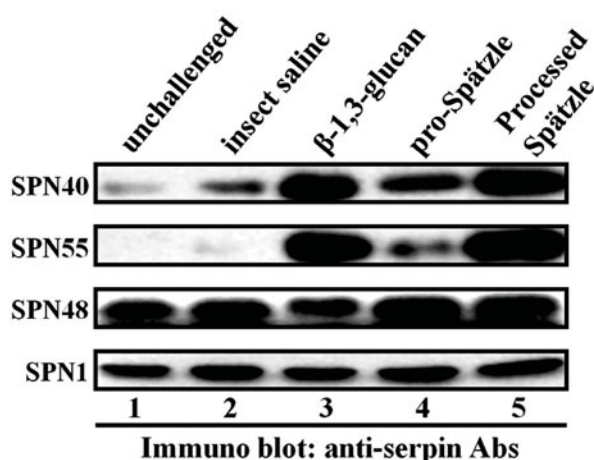


Figure 8. The amounts of SPN40 and SPN55 in the hemolymph increased after injection of β -1,3-glucan or processed Spätzle into *Tenebrio* larvae. Four microliters of insect saline (lane 2), 50 ng of β -1,3-glucan (lane 3), 60 ng of pro-Spätzle (lane 4) and 60 ng of processed Spätzle (lane 5) were injected into 10 *Tenebrio* larvae and hemolymph was collected after 24 h. A portion of each sample (40 μ g) was analyzed by immunoblotting using affinity-purified antiserpin antibodies. Reproduced from Jiang R et al. *J Biol Chem* 2009; 284:35652-58,⁶¹ ©2009, with permission from the American Society for Biochemistry and Molecular Biology.

precisely regulated. Our data indicate that each protease in a cascade may be regulated by a specific serpin. Furthermore, we have shown that SPN40 and SPN55 function as inducible negative feedback regulators *in vivo*. This study, in combination with our other reports,^{44,49,68} supports a model in which the Lys-type PG- and β -1,3-glucan-dependent Toll signaling and proPO cascades are negatively regulated by three serpins (Fig. 9). Furthermore, our study highlights the elaborate regulatory mechanism of invertebrate molecular defense systems.

CONCLUSION

During last 15 years, we have used two beetles for biochemical studies of invertebrate's innate immunity. The main reason to use beetles as a model system is because they can be used for biochemical studies since fairly large amounts of hemolymph can be obtained. A weak point is that the genomic sequence information of beetles except for *T. castaneum* is not available until now. When we purified new proteins that are involved in innate immune responses, their putative biological functions have been predicted based on amino acid sequence comparison between partial amino acid sequence of the purified beetles' proteins and amino acid sequences obtained from *Tribolium* and *Drosophila* genomic sequence. Furthermore, in order to determine the exact biological functions of these proteins, we have spent long time to obtain their cDNAs and to express the recombinant proteins. Therefore, the determination of the whole genomic sequences of *Tenebrio* larvae and adults will be essential and efficient for further understanding of invertebrate's innate immune reactions. Finally, a greater understanding of molecular

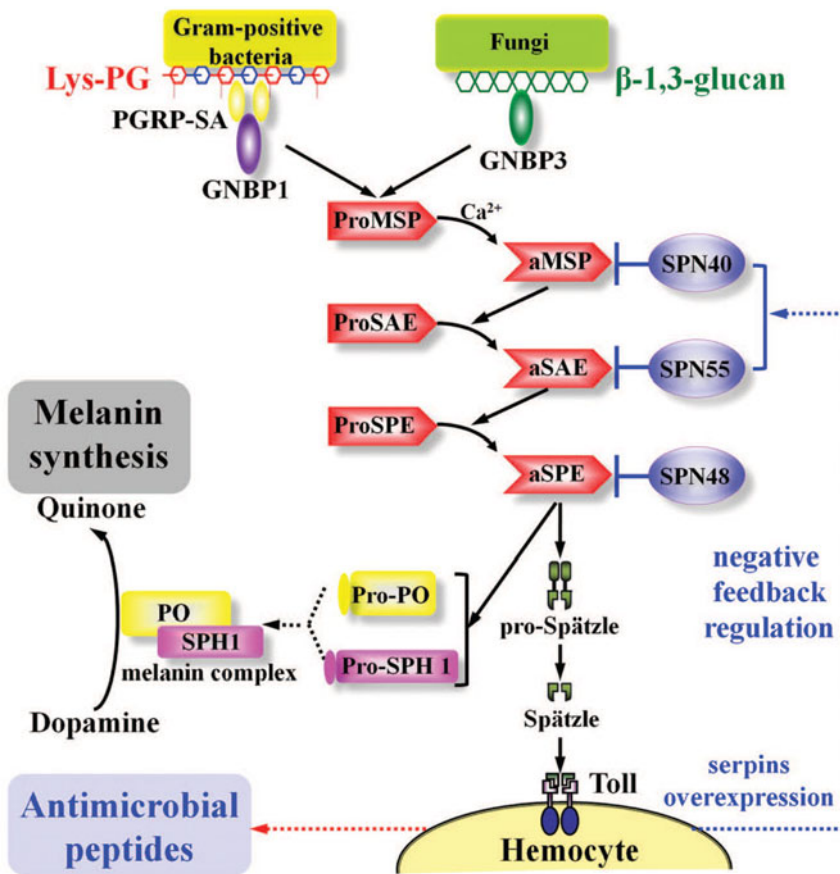


Figure 9. Model summarizing the molecular events in the regulation of the Toll and melanin synthesis cascades. The molecular activation mechanism of the *Tenebrio* Toll cascade leading to production of AMPs^{44,49} and of the pro-PO cascade leading to melanin synthesis⁶⁸ were reported. Processed Spätzle induces the production of endogenous SPN40 and SPN55 as a negative feedback regulator of the Toll cascade. Also, the three serpins blocked activation of proPO, leading to inhibition of melanin synthesis. Reproduced from Jiang R et al. J Biol Chem 2009; 284:35652-58,⁶¹ ©2009, with permission from the American Society for Biochemistry and Molecular Biology.

pathogen recognition mechanisms and their signaling pathways of proteolytic cascade reactions will also facilitate the development of novel kits to rapidly and sensitively detect bacterial PG or β-1,3-glucan in blood and food products.

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IMMUNITY IN LEPIDOPTERAN INSECTS

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Abstract: Lepidopteran insects provide important model systems for innate immunity of insects, particularly for cell biology of hemocytes and biochemical analyses of plasma proteins. Caterpillars are also among the most serious agricultural pests, and understanding of their immune systems has potential practical significance. An early response to infection in lepidopteran larvae is the activation of hemocyte adhesion, leading to phagocytosis, nodule formation, or encapsulation. Plasmatocytes and granular cells are the hemocyte types involved in these responses. Infectious microorganisms are recognized by binding of hemolymph plasma proteins to microbial surface components. This “pattern recognition” triggers phagocytosis and nodule formation, activation of prophenoloxidase and melanization and the synthesis of antimicrobial proteins that are secreted into the hemolymph. Many hemolymph proteins that function in such innate immune responses of insects were first discovered in lepidopterans. Microbial proteinases and nucleic acids released from lysed host cells may also activate lepidopteran immune responses. Hemolymph antimicrobial peptides and proteins can reach high concentrations and may have activity against a broad spectrum of microorganisms, contributing significantly to clearing of infections. Serine proteinase cascade pathways triggered by microbial components interacting with pattern recognition proteins stimulate activation of the cytokine Spätzle, which initiates the Toll pathway for expression of antimicrobial peptides. A proteinase cascade also results in proteolytic activation of phenoloxidase and production of melanin coatings that trap and kill parasites and pathogens. The proteinases in hemolymph are regulated by specific inhibitors, including members of the serpin superfamily. New developments in lepidopteran functional genomics should lead to much more complete understanding of the immune systems of this insect group.

INTRODUCTION

Moth larvae have proven to be extremely useful for experiments providing insights on the innate immune systems of insects. Many hemolymph proteins with immune functions were first studied in caterpillars by biochemical methods.¹⁻¹¹ Lepidopteran larvae have also been important for experiments aimed at characterizing immune functions of insect hemocytes.¹² Much of the research on lepidopteran immunity has made use of large moth species, including the tobacco hornworm, *Manduca sexta* and wild silkmoths such as *Hyalophora cecropia*. The domestic silkmoth, *Bombyx mori* has also provided significant discoveries about immunity in moths and with the advantage of a sequenced genome,¹³ it continues to serve as an important model organism. The wax moth, *Galleria mellonella*, was one of the earliest species used for research in insect immunity^{14,15} and is the subject of much current research. In addition to these laboratory model species, research on immunity is also underway to investigate the immune systems of moths whose larvae are among the most destructive agricultural pests worldwide, particularly including species in the family Noctuidae. In comparison, there has been relatively little research on immunity in butterflies, perhaps because they can be more difficult to rear or due to lower agricultural impact.

In this chapter, we describe selected developments in research on innate immunity in lepidopteran insects, with an emphasis on aspects discovered through biochemical and cell biological approaches and potential new insights that may be gained from functional genomics methods. Several recent reviews with a focus on lepidopteran immunity are available.¹⁶⁻²³

HEMOCYTES

With a large hemolymph volume in many lepidopteran larvae and pupae, it is possible to collect 10^4 - 10^6 hemocytes from a single individual, making feasible the use of cell biological techniques such as flow cytometry and cell sorting that have advanced mammalian immunology. The most abundant hemocyte types typically described in Lepidopteran larvae are granular cells and plasmatocytes, which are capable of adhesion and phagocytosis.¹² Nonadherent hemocyte types include oenocytoids, which synthesize prophenoloxidase (proPO), and spherule cells, whose functions are poorly understood. Monoclonal antibodies are very useful reagents for distinguishing lepidopteran hemocyte populations based on antigenicity rather than morphology,²⁴⁻²⁶ which can vary considerably, especially for plasmatocytes. Two subpopulations of plasmatocytes have been distinguished based on monoclonal antibody markers in *Pseudoplusia includens*²⁶ and in *M. sexta*,²⁷ which may indicate different stages in differentiation or functional specialization. The composition of hemocyte populations varies through larval development.^{28,29} Hematopoiesis in lepidopteran larvae^{28,31} and the embryonic origin granular cells and plasmatocytes³⁰ have been described.

Granular cells and plasmatocytes function in immunity through responses that involve adherence of cells to foreign surfaces or to other hemocytes.^{22,32} Both granular cells and plasmatocytes can be phagocytic and in different lepidopteran species either cell type may be the predominant contributor to phagocytosis as a defense.¹² A hyperphagocytic cell type, very large cells capable of phagocytosing 500 bacteria, has been described in *M. sexta*.³³ These cells are morphologically similar to a neuroglian-positive subpopulation

of plasmatocytes, which can act as a focus for attachment of hemocytes to foreign surfaces³⁴ and perhaps are the same hemocyte type.

Adhesion of granular cells and plasmatocytes leads to two related responses to infection: nodule formation, in which hemocytes cluster to entrap aggregated microorganisms, and encapsulation, in which hemocytes form a multi-layered cellular capsule around a larger eukaryotic parasite.¹² Adhesion of hemocytes to injured body wall also probably helps to seal wounds to prevent bleeding.³² Cytokines have been identified in lepidopterans, which promote these hemocyte functions. A hemocyte chemotactic peptide from *Pseudaletia separata* stimulates directed movement and aggregation of hemocytes.³⁵ This peptide is structurally similar to lepidopteran cytokines called ENF peptides, which have multiple biological activities, including the stimulation of plasmatocyte adhesion and spreading, reduced bleeding and the stimulation of oenocytoid lysis to release proPO.³⁶⁻⁴⁰ RNAi results indicate that the *M. sexta* plasmatocyte spreading peptide from this family promotes hemocyte nodule formation as a protective response to bacterial infection.⁴¹ The active ENF peptides are produced by proteolytic processing of a larger protein present in hemolymph.^{36,37} In *M. sexta* hemolymph, a serine proteinase with trypsin-like specificity is responsible for this activation, but has not yet been identified due to its instability (Kanost et al, unpublished results). Eicosanoids such as prostaglandins can also stimulate hemocytes to aggregate to form nodules.^{42,43} Prostaglandins can also elicit the lysis of oenocytoids, releasing proPO into the plasma.⁴⁴ ProPO is activated by plasma proteinases and participates in formation of melanin, which coats nodules and encapsulated objects. This response is discussed in more detail below.

Hemocyte attachment during encapsulation and nodulation is mediated by cell surface adhesion proteins. Lepidopteran hemocytes have cell surface integrins, which function as adhesion molecules.⁴⁵⁻⁵⁰ Plasmatocytes of *M. sexta* express a specific integrin that is required for efficient encapsulation.^{47,50} The adhesive properties of this hemocyte-specific integrin derive at least in part from binding to neuroglian and a tetraspanin on neighboring hemocytes.^{49,51}

RECOGNITION OF MICROORGANISMS

Plasma proteins that bind to components on the surface of microorganisms are a key component of the innate immune system of insects. Such proteins stimulate responses including phagocytosis and activation of proteinase signaling cascades. Some of these “pattern recognition” proteins were first discovered in silkworms and then found to occur in immune systems of other insect groups, whereas others appear to be unique to lepidopterans.

Hemolin is a 48 kDa plasma protein first identified in *Hyalophora cecropia*^{52,53} and *Manduca sexta*.^{54,55} Hemolin is composed of four I-set immunoglobulin (Ig) domains commonly found in cell adhesion proteins of vertebrates and invertebrates. Hemolin also exists in other lepidopteran species including *B. mori*, *Hyphantria cunea*, *Lymantria dispar*, *Antheraea pernyi*, *Antheraea mylitta*, *Plutella xylostella*, *Samia cynthia*.^{13,56-61} Hemolin has not been identified in insects from any other order, suggesting that it may have evolved after the split of the Lepidoptera from other insect groups. Hemolin shares structural features with neuroglian,⁶² a transmembrane Ig-domain protein located on the surface of *Manduca* glial and neuronal cells⁶³ and the developing embryonic prothoracic gland,⁶⁴ as well as a subpopulation of *M. sexta* plasmatocytes.^{34,51} Hemolin expression

is induced by bacteria or their surface components injected into the hemocoel. In *H. cecropia*, this transcriptional activation is controlled via an intronic enhancer that contains a κ B motif.⁶⁵ Developmental and hormonal signals (e.g., 20-hydroxyecdysone) also affect hemolin production.^{58,66,67} It has been speculated that hemolin may have an antiviral function.⁶⁸ Baculovirus exposure up-regulates hemolin transcription in *A. pernyi*,⁶⁹ but not in *B. mori*, *Helicoverpa zea*, or *Heliothis virescens*.^{13,70} A possible role for hemolin in antiviral responses remains to be established. On the other hand, several lines of evidence support the idea that hemolin functions in immune responses to bacterial infection. Hemolin binds to bacterial lipopolysaccharide (LPS) and lipoteichoic acid^{71,72} and associates with hemocytes.^{55,73,74} The horseshoe-shape structure of hemolin suggests that one or more of its Ig domains may interact through domain swapping with Ig domains of cell adhesion molecules such as neuroglian on hemocyte surfaces.⁷⁵ Interaction of hemolin with molecules on the surface of bacteria and with hemocyte membranes suggests that it may bring microorganisms to hemocyte surfaces, promoting phagocytosis or nodule formation. RNAi knockdown of hemolin expression in *M. sexta* larvae significantly reduced phagocytosis and nodulation of *E. coli*.⁷⁶

Peptidoglycan recognition proteins (PGRPs) associate with bacterial peptidoglycans through a conserved domain homologous to T4 bacteriophage lysozyme.⁷⁷ PGRPs were first discovered in lepidopterans. *B. mori* PGRP-S1 was isolated from plasma of silkworm larvae as a 19 kDa protein that binds to *Micrococcus luteus* peptidoglycan and triggers the proPO activation system.⁸ Molecular cloning of PGRPs from *Trichoplusia ni*,¹⁰ *B. mori*,^{13,78} *M. sexta*,⁷⁹⁻⁸¹ *G. mellonella*,⁸² *S. cynthia*,^{83,84} *P. xylostella*,⁶⁰ and *Ostrinia nubilalis*⁸⁵ suggests that multiple PGRPs are present in every lepidopteran species. In addition to binding to peptidoglycan, these proteins may hydrolyze peptidoglycan if the residues for Zn²⁺-binding and amidase activity are present.⁸⁶ The *B. mori* genome has six short (S) and six long (L) PGRP genes: S3 through S6 are putative amidases; L1 and L4 possess a transmembrane region; L6 is likely cytosolic.¹³ Constitutive expression of PGRP-S1 is up-regulated in larvae after injection of *Enterobacter cloacae*.⁷⁸ *E. coli* or *Bacillus subtilis* treatment increases the mRNA levels for PGRP-S1, -S2 and -S5, whereas *Staphylococcus aureus* injection enhances transcription of *B. mori* PGRP-S1, -S5, -L1, but not -S2.¹³ Microbe-induced transcription of PGRPs also occurs in other lepidopteran species.^{10,79,82,84} The *T. ni*, *S. cynthia* and *M. sexta* PGRPs bind to *Bacillus meso*-diaminopimelic acid-type peptidoglycans and *M. luteus* Lys-type peptidoglycans.^{10,83,87} This *M. sexta* PGRP can stimulate proPO activation.⁸⁸ Knockdown of *M. sexta* PGRP1 synthesis increased larval susceptibility to infection by *Photorhabdus luminescens*.^{89,90} So far, the level of knowledge on structures and functions of moth PGRPs is significantly less than for *Drosophila* PGRPs.^{91,92} Furthermore, molecular details of how peptidoglycan binding by PGRPs promotes the activation of serine proteinases involved in proPO and spätzle activation are unknown.

Insect β -1,3-glucan recognition proteins (β GRPs) and Gram-negative bacteria binding proteins (GNBPs) are a family of ~55 kDa plasma proteins with an amino-terminal glucan-binding domain and a carboxyl-terminal region similar to β -1,3-glucanases. *B. mori* β GRP1, *M. sexta* β GRP1 and β GRP2 and *Plodia interpunctella* β GRP bind to β -1,3-glucans and to bacteria and stimulate the proPO activation cascade.^{9,93-97} The amino-terminal domain of *B. mori* β GRP1 adopts an Ig-like β -sandwich fold and residues have been identified that may form a glucan-binding site.⁹⁸ *M. sexta* β GRP1 and β GRP2 gene expression in fat body is differentially regulated: β GRP1 is constitutively expressed, whereas β GRP2 transcripts become highly abundant in the early wandering stage prior

to pupation or after an immune challenge.^{94,95} *M. sexta* β GRP2 is also present in cuticle of wandering stage larvae.⁹⁵ Binding of β GRP1 or β GRP2 to curdlan and *M. sexta* hemolymph proteinase-14 precursor (proHP14) stimulates autoactivation of HP14 to initiate an immune proteinase cascade leading to proPO activation.^{99,100} *B. mori* GNBPs bind to *E. cloacae* but not *Bacillus licheniformis* or curdlan.¹⁰¹ An orthologous *M. sexta* GNBPs recognize LPS and laminarin (a β -1,3-glucan with β -1,6 branches) and initiates melanization (Y. Wang and H. Jiang, unpublished results). An active glucanase related in sequence to the GRPs was isolated from midgut extract of *Helicoverpa armigera* larvae. It hydrolyzes β -1,3-glucan but not β -1,4-glucan or glucans with mixed β -1,3 and β -1,4 linkages¹⁰² and probably functions as a digestive enzyme.

C-type lectins (CTLs) from animals are a large group of carbohydrate-recognition molecules that bind ligands in a calcium-dependent manner.¹⁰³ *B. mori* LPS-binding protein (LBP or CTL20),^{104,105} *Hyphantria cunea* Hdd15,¹⁰⁶ and *M. sexta* immuelectin-1¹⁰⁷ were among the first C-type lectins identified in plasma of lepidopteran larvae. They all contain two carbohydrate-recognition domains. CTLs with this dual-domain structure also include: *M. sexta* immuelectins-2, -3 and -4,¹⁰⁸⁻¹¹⁰ *B. mori* MBP (CTL10),¹¹¹ immuelectin (CTL11),¹¹² LEL-1 and -2,¹¹³ CTL19, CTL21,¹³ and *H. armigera* Ha-lectin.¹¹⁴ GenBank currently contains more than 30 additional similar CTL cDNA sequences from eleven other lepidopteran species in five families. The only other dual-domain CTL identified in any other insect species is TcCTL3¹¹⁵ from a beetle, *Tribolium castaneum*. Hence, such tandem domain CTL genes appear to be fairly unique to Lepidoptera and their emergence and expansion may have occurred early in evolution of this insect group. Expression of at least some of these CTL genes is induced microbial infection.^{106,108-110,114}

Many of the lepidopteran CTLs bind to bacterial LPS and some to lipoteichoic acid.^{105,106,108-110} They can cause agglutination of bacteria and yeast,^{107,109-111} presumably due to each of the molecule's two carbohydrate-binding domains binding to carbohydrates on the surface of adjacent microbial cells. This activity most often requires the presence of calcium. Such clustering of microorganisms may result in more efficient clearance of pathogens by hemocytes through nodule formation.¹⁰⁵ Experiments have demonstrated that *M. sexta* immuelectin-2 enhances clearance of *Serratia marcescens*,¹¹⁶ and suppression of immuelectin-2 expression by RNA interference reduced larval survival of a *Photorhabdus* infection.⁹⁰ In addition, the immuelectins can promote proPO activation and melanin deposition at the surface of objects encapsulated by hemocytes,^{107,108,110,117,118} which may also promote killing of pathogens and parasites.

Lipophorins are insect hemolymph proteins that transport lipids between tissues.¹¹⁹ The lipophorin particle contains two protein subunits, apolipophorin-I and apophorin-II. When the neutral lipid load is high, an exchangeable plasma protein, apolipophorin-III (apoLpIII) also associates with low density lipophorin to cover hydrophobic surfaces. In lepidopterans, apoLpIII concentration in hemolymph is generally much lower in larvae than in adults, where it functions in transporting lipids from fat body to flight muscles during prolonged flight. Lipophorins and apoLpIII have been implicated in several defense mechanisms.³² Lipophorin appears to be involved in hemolymph clotting in at least some species,²⁰ and may in this way participate in physical trapping of invading microorganisms. There may also be an association of proteins from the melanization cascade with lipophorin or other clotting components, which could contribute to defense.¹²⁰⁻¹²² The affinity of lipophorin and apoLpIII for hydrophobic ligands is consistent with their reported binding of bacterial LPS and lipoteichoic acids,¹²³⁻¹²⁸ and the partitioning of these microbial membrane lipids into complexes with lipophorin or apoLpIII may contribute to reducing their toxicity to

insect hosts. Lipophorin and apoLpIII have been reported to stimulate other humoral or cellular immune responses in *G. mellonella*,¹²⁹⁻¹³³ and research toward understanding molecular mechanisms underlying such observations is needed.

Pathogenic bacteria and fungi produce proteinases to utilize lepidopteran host proteins as a source of nutrients and to degrade immunity-related defense molecules such as antimicrobial peptides. Thermolysin-like metalloproteinases associated with entomopathogenic bacteria and fungi are essential virulence factors.¹³⁴⁻¹³⁷ However, the presence of microbial proteinases may also be recognized as a signal of infection and stimulate immune defenses in the host. Evidence for sensing of microbial proteinases and their regulation during innate immune responses has been reported from *G. mellonella*.¹³⁸ Thermolysin is a potent activator of the serine proteinase cascade that controls proPO activation leading to melanization. Thus, this virulence factor also directly triggers an immune response. In addition, the activity of microbial metalloproteinases within the body of *G. mellonella* generates peptide fragments that strongly elicit the synthesis of antimicrobial peptides.¹³⁹ The immune-stimulatory peptides were identified as collagen IV fragments containing the RGD/RGE motif, which bind to integrins of immune-competent hemocytes.¹⁴⁰ The stimulation of two innate immune responses, proPO activation and antimicrobial peptide synthesis, in response to microbial metalloproteinases provides evidence that lepidopteran innate immune responses include reaction to danger-associated molecules produced by microbial virulence factors.¹⁴¹ Furthermore, *G. mellonella* hemolymph contains an inducible metalloproteinase inhibitor (IMPI), the only specific peptidic inhibitor of thermolysin-like metalloproteinases reported to date from any animal.¹⁴² The IMPI gene encodes two distinct metalloproteinase inhibitors that putatively contribute to the regulation of metalloproteinases associated with invading pathogens.^{143,144}

Nucleic acids released from damaged or necrotic cells form another danger signal to enhance insect immune responses. Injection of synthetic oligonucleotides induced attacin expression in *B. mori* larvae.¹⁴⁵ Co-injection of purified host nucleic acids with heat-inactivated *P. luminescens* into *G. mellonella* larvae synergistically elevated the level of antimicrobial activity, reduced the total number of hemocytes (a consequence of the attachment of immune-competent cells to tissues during cellular responses) and prolonged the survival of insects infected by *P. luminescens*.¹⁴⁶ DNA and RNA released from damaged cells may interact with lipophorin to trigger clot formation and entrap invading pathogens in hemolymph. In *Pseudaletia separata*, nucleic acids as well as cytoplasmic proteins (*e.g.*, proPO) are released from oenocytoids through cell lysis induced by the growth-blocking peptide.⁴⁰

ANTIMICROBIAL PEPTIDES AND PROTEINS

Many insect antimicrobial plasma proteins were first identified and isolated from lepidopteran hemolymph. Expression of antimicrobial peptides and proteins is often induced by microbial infection, with strongest expression usually occurring in fat body, although hemocytes also contribute to the pool of antimicrobial peptides secreted into hemolymph plasma.¹⁴⁷ Antimicrobial peptides are also expressed in the midgut of lepidopteran prepupae and secreted into the lumen, perhaps as a prophylaxis against infection during metamorphosis.¹⁴⁸ Antimicrobial peptides can also be expressed in extraembryonic tissues of lepidopteran eggs, providing protection against infection for the developing embryo.¹⁴⁹

Lysozyme, the first antimicrobial protein reported from insects, was identified more than forty years ago in *G. mellonella*¹⁵⁰ and, like other insect lysozymes, shares structural similarity with C-type (chicken) lysozyme.¹⁵¹ Its activity against Gram-positive bacteria has been attributed to its ability to degrade cell wall peptidoglycan by hydrolysis of the β -1,4 linkages between N-acetylglucosamine and N-acetylmuramic acid residues.² Besides moderate activity against Gram-negative bacteria,¹⁵²⁻¹⁵⁴ lepidopteran lysozyme also exhibits antifungal activity.¹⁵⁵ Lysozyme also appears to negatively regulate activation of proPO in *M. sexta*.¹⁵⁶

Insects produce a variety of amphipathic peptides with antimicrobial activity attributed to their ability to damage cell membranes of pathogens. The first antimicrobial peptide from insects, isolated from the hemolymph of the silkworm *H. cecropia*, was named cecropin.⁴ Families of cecropin genes have now been found in many lepidopteran (and dipteran) species. Thirteen cecropin genes were identified in the *B. mori* genome.¹³ Cecropins are typically ~4 kDa basic peptides, which have an amphipathic α -helical structure. The moricins constitute another group of amphipathic α -helical antimicrobial peptides,^{157,158} first discovered in *B. mori*.¹⁵⁹ Nine moricin genes are present in the *B. mori* genome.¹³ Eight moricin homologs identified in *G. mellonella* have activity against Gram-negative and Gram-positive bacteria, as well as against yeast and filamentous fungi.¹⁶⁰

Lepidoptera possess glycine-rich AMPs (attacins and gloverins) and proline-rich AMPs (lebocins). *H. cecropia* has two 20 kDa attacin isoforms, an acidic and a basic attacin, with 80% sequence identity.^{5,161} The *B. mori* genome also contains two attacin genes,¹³ and attacin cDNAs have now been cloned from many lepidopteran species. Treatment of *E. coli* with *H. cecropia* attacins leads to an increase in outer-membrane permeability, preceding any increase in inner-membrane permeability by at least one generation time. Inhibition of outer-membrane protein synthesis is achieved on the transcriptional level and triggered by binding of attacin to the cell surface without entering the inner membrane or the cytoplasm. Primary binding occurs on LPS, explaining why basic attacin is more active against *E. coli* than the acidic form. Another family of proline-rich AMPs, the gloverins, appears to exert a similar mechanism of inhibition of outer-membrane protein synthesis.¹⁶² Expression and evolution of four *B. mori* gloverin genes have been investigated,^{163,164} and gloverins have been studied in several other lepidopteran species,^{16,165-168} but no homologs have been identified to date in insects from other orders. A family of proline-rich AMPs called lebocins has been characterized in lepidopterans.^{16,147,169-173} A somewhat puzzling aspect of this family is that the 3.5 kDa active lebocin peptide is processed from a larger precursor and in some members of the family, the amino-terminal pro-region of the protein is conserved in sequence, but the carboxyl-terminal sequence corresponding to the original antimicrobial peptide is not, suggesting that the pro-region may have a function not yet discovered.

AMPs stabilized by intramolecular disulfide bonds are widely distributed in insects. Those with three or four disulfide bonds are commonly referred to as insect defensins because of overall structural similarities to mammalian α and β defensins.¹⁷⁴ Insect defensins can be grouped into peptides with an α -helix/ β -sheet mixed structure and peptides forming triple-stranded antiparallel β -sheets. Defensin-like AMPs with antibacterial and antifungal activities from several lepidopteran species have been investigated.¹⁷⁵⁻¹⁸⁰ Two cysteine-rich defensin-like peptides from *G. mellonella* specifically inhibit growth of filamentous fungi.^{179,180} A group of atypical defensin-like peptide named x-tox identified in *G. mellonella* and two *Spodoptera* species is characterized by imperfectly conserved tandem repeats of cysteine-stabilized $\alpha\beta$ motifs, the structural scaffold characteristic of

invertebrate defensins and scorpion toxins.¹⁸¹⁻¹⁸³ They are induced upon activation of the immune system but lack detectable antimicrobial activity, suggesting that they may have an immune function yet to be discovered.

EXTRACELLULAR AND INTRACELLULAR SIGNAL TRANSDUCTION STIMULATING ANTIMICROBIAL PEPTIDE SYNTHESIS

Genetic investigations in *Drosophila* have revealed three major immune signaling pathways (Toll, Imd-JNK and JAK-STAT) that are conserved in mammals.¹⁸⁴ Genomic analyses suggest similar pathways exist in other holometabolous insects, including *B. mori*.¹³ Additional bioinformatic and experimental evidence described below supports the existence of functional Toll and Imd pathways in lepidopterans.

There are fourteen Toll-like receptor genes in the silkworm genome: six in group A with *Drosophila* Toll and eight in group B with *Drosophila* 18-wheeler.¹³ *BmToll* and *BmToll2* are group B genes highly expressed in ovary and their transcripts become more abundant in fat body after injection of microorganisms,¹⁸⁵ suggesting possible involvement in embryonic development and immune response. *BmToll3*, *BmToll4*, *BmToll9* and *BmToll10* mRNA levels in fat body also increase after injection of some microbes. In *M. sexta*, a Toll-like receptor is present in hemocytes, fat body, epidermis, midgut and Malpighian tubules.¹⁸⁶ Its mRNA level increased in hemocytes, but not in fat body, after injection of microorganisms.

Spätzle, the ligand which activates Toll, has been identified and functionally characterized in *B. mori* and *M. sexta*.^{187,188} Spätzle is synthesized as an inactive precursor, proSpätzle, which is secreted as a disulfide-linked homodimer into the hemolymph and requires proteolytic processing to form the active Toll ligand. Expression of proSpätzle is significantly greater in *M. sexta* hemocytes than in fat body.¹⁸⁸ In *B. mori*, expression was detected primarily in fat body and midgut, but hemocytes were not tested.¹⁸⁷ *B. mori* and *M. sexta* proSpätzle proteins are only ~20% identical to *Drosophila* proSpätzle, but have slightly greater similarity (~26% identity) in the carboxyl-terminal 108 residues corresponding to the active form of *Drosophila* Spätzle known to bind to Toll. Recombinant *B. mori* and *M. sexta* Spätzle were active when injected into larvae, inducing fat body expression of attacins, cecropins, gloverin, moricin and lebecin in *B. mori*¹⁸⁷ and attacin, cecropin, moricin and hemolin in *M. sexta*, with corresponding strong induction of plasma antimicrobial activity.¹⁸⁸ However, injection of proSpätzle had little effect, indicating the need for proteolytic activation of this cytokine in response to infection.

Hemolymph proteinase-8 (HP8) is a clip-domain proteinase demonstrated to activate proSpätzle in *M. sexta* by specific cleavage to produce the carboxyl-terminal 108 residue fragment, as a disulfide-linked homodimer.¹⁸⁸ The *Drosophila* proteinases most similar to HP8 are Easter and Spätzle processing enzyme, both of which function to activate proSpätzle.¹⁸⁹ Injection of active HP8 into larvae stimulates expression of attacin, cecropin, gloverin and moricin and elevates plasma antibacterial activity, consistent with a role for HP8 as an activator of proSpätzle.¹⁸⁹ HP8 is present in plasma as a zymogen, proHP8 and is activated by another clip-domain proteinase, HP6, an apparent ortholog of *Drosophila* Persephone. Injection of recombinant HP6 also promoted expression of antimicrobial peptides in larvae.¹⁸⁹ ProHP6 is activated in plasma exposed to bacteria or the β -1,3-glucan curdlan, but a hemolymph proteinase responsible for activation of HP6 has not been identified yet. It is apparent that recognition of microbial pattern

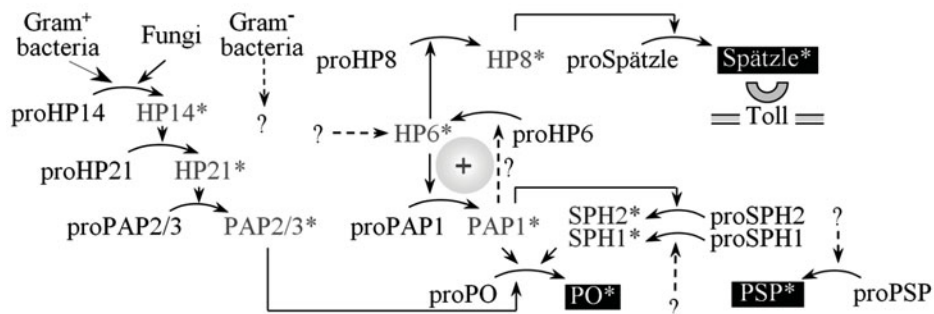


Figure 1. A current model of the hemolymph proteinase system in *M. sexta* larvae. An initiation proteinase precursor, proHP14, is autoactivated in response to Gram-positive bacterial or fungal infection. HP14 activates proHP21; HP21 activates proPAP2 or proPAP3; PAP2 or PAP3 then cleaves proPO to form active PO in the presence of SPH1 and SPH2. Activation of proPO can also be catalyzed by PAP1 when the high *M*₆ SPH complex is present simultaneously. PAP1 also activates proSPH2 directly and can indirectly lead to proHP6 activation. HP6, whose direct activator is unknown, cleaves proPAP1 and proHP8. PAP1 and HP6 form a positive feedback loop, in which PAP1 indirectly stimulates activation of HP6. HP8 activates Spätzle to induce antimicrobial peptide synthesis via Toll receptor. Active proteins, including HPs, PAPs, SPHs, PO, Spätzle and plasmatocyte-spreading peptide (PSP), are labeled “*” and unknown HPs are marked with “?”.

molecules triggers activation of a proteinase cascade to generate the cytokine Spätzle, leading to expression of a suite of antimicrobial peptides as an innate immune response in this moth (Fig. 1).

Two rel-family transcription factors participate in immunity-related gene expression in *B. mori*.¹⁹⁰⁻¹⁹² *BmRel* encodes, via alternative splicing, RelA and RelB, which are orthologous to *Drosophila* Dorsal. *BmRelish* also encodes two splicing isoforms, Relish1 and Relish2. RelA activates the lebecin-4 gene strongly and an attacin gene weakly.¹⁹⁰ RelB, lacking the first 52 residues of RelA, activates the attacin gene strongly and other genes to a lesser extent. The Rel homology domain in RelA and RelB binds specifically to κB sites in attacin and lebecin-4 genes. In transgenic silkworms whose *BmRel* expression is knocked down, expression of antibacterial peptide genes fails to be induced by *M. luteus*. Knockdown of *BmRelish* expression abolishes antimicrobial peptide production elicited by *E. coli*.¹⁹¹ Intact Relish1 and Relish2 do not activate promoters of *B. mori* attacin, cecropin B1, lebecin-3 and lebecin-4 genes. Removal of the ankyrin repeats in Relish1 is necessary for its transcriptional activation of antibacterial peptide genes. However, Relish2, which lacks the repeats and the transactivation domain, serves as a dominant negative factor to suppress the function of active Relish1. Relish1 binds to the κB sites in attacin and cecropin B1 genes, while the sites for activating lebecin-4 promoter differ between Relish1 active form and RelA. *Meso*-diaminopimelic acid- and Lys-type peptidoglycans can stimulate differential expression of antimicrobial peptide genes in the silkworm, which is affected by the level, binding affinity and transactivation activity of Relishes and Rels.¹⁹² Relish1 is inhibited by its own ankyrin repeats and RelA and RelB are negatively regulated by *B. mori* Cactus.¹⁹³ Cactus interacts with the DNA-binding domain in the Rels but not with Relish1 or Relish2. Taken together, these data strongly suggest that the Toll and Imd pathways are functional in the silkworm to regulate immunity-related gene expression.

PROPHENOLOXIDASE ACTIVATION SYSTEM

PO-catalyzed quinone and melanin formation is a universal response in arthropods for killing and entrapping pathogens or parasites.^{194,195} PO in insect hemolymph has tyrosinase-like activities, including *o*-hydroxylation of monophenols and oxidation of *o*-diphenols to quinones.¹⁸ Tyrosine, DOPA and dopamine are substrates in insect hemolymph that contribute to PO-catalyzed quinone formation and subsequent melanin synthesis. Tyrosine hydroxylase and dopa decarboxylase are upregulated in fat body after injection of bacteria and probably contribute to provision of hemolymph dopamine for the innate immune response.^{80,196-198} Oxidation of dopamine by PO leads to production of 5,6-dihydroxyindole, which has antimicrobial activity toward bacteria and fungi.¹⁹⁹ PO is produced as a zymogen that requires a specific proteolytic cleavage to gain activity. This regulatory mechanism protects the host insect from potentially harmful effects of the reactive chemicals produced by PO, as the enzyme is activated only when elicited by wounding or infection. Understanding of insect PO and its activation was pioneered through detailed biochemical investigations with *B. mori*,¹¹ and the *M. sexta* model system is providing new insights into lepidopteran PO function and regulation.^{17,19}

ProPO from *B. mori* and *M. sexta* exists in plasma as a heterodimer of two related subunits, each ~80 kDa.^{200,201} Insect proPO sequences are related to arthropod hemocyanins, and copper-binding motifs in the two groups of proteins are conserved.²⁰² ProPO is synthesized constitutively by oenocytoids.^{197,201,203} ProPOs lack secretion signal peptides and are released from oenocytoids by lysis of the cells.^{40,44} Activation of the proPO zymogen requires cleavage of a conserved Arg-Phe bond about 50 residues from the amino-terminus.^{11,18} The crystal structure of the *M. sexta* proPO heterodimer suggests that the proteolysis between Arg51 and Phe52 induces a conformational change to dislodge a specific Phe residue in each subunit and open up the active site for substrate binding.²⁰⁴ The active site contains a canonical Type-3 di-nuclear copper center, with each copper ion coordinated by three conserved His residues. Glu395 of the subunit-2 may act as a general base to deprotonate monophenols, a key step in the *o*-hydroxylation of tyrosine by PO.

Extracellular serine proteinase pathways have evolved in animals to stimulate rapid responses to tissue damage, pathogen invasion, or physiological cues.²⁰⁵ A few eliciting molecules, via specific recognition and cascade amplification, lead to sequential proteolytic activation of a large number of pathway components within minutes. This type of proteinase pathway results in activation of proPO in response to infection. Many of the proteinases that function in such cascade pathways in arthropods contain a carboxyl-terminal serine proteinase domain similar trypsin or chymotrypsin and one or two amino-terminal clip domains, which have likely regulatory functions.²⁰⁶ The proteinases from lepidopterans known to activate proPO and most of the proteinases upstream in the activation pathway are clip domain proteinases. Fifteen clip domain proteinase genes were identified in the *B. mori* genome,¹³ and fourteen such enzymes are expressed in fat body or hemocytes of *M. sexta*.²⁰⁷

In lepidopterans, proPO activating proteinases (PAPs) have been well characterized in *M. sexta*²⁰⁸⁻²¹¹ and *B. mori*.²¹² They are present in hemolymph as zymogens at low concentration in naïve larvae, and their expression in fat body is upregulated in response to injection of bacteria. *M. sexta* PAP1 contains a single clip domain, whereas PAP2 and PAP3 and *B. mori* proPO activating enzyme each contain two clip domains. The solution structure of the region of PAP2 containing the dual clip domains suggests a potential proPO-binding site, a bacteria-interacting region and a surface for activator/adaptor

docking in each domain.²¹³ Purified *M. sexta* PAP1, PAP2 and PAP3 do not efficiently generate active PO activity, even after a significant amount of proPO is cleaved at Arg51, without the presence of protein cofactors from hemolymph, identified as serine proteinase homologs (SPHs). SPHs also contain clip domains but lack proteinase activity due to substitution of the active site Ser residue with Gly.^{214,215} *M. sexta* SPH1 and SPH2 also require proteolytic processing to gain function, which leads to their assembly into the active, high *M_r* cofactor required in the reaction with proPO and PAP to generate high levels of PO activity.^{215,216} This interesting interaction, which does not seem to be required for the silkworm proPO activating enzyme,²¹² is not well understood and requires further investigation.

The *M. sexta* proPO activation system includes at least four other serine proteinases (Fig. 1). An initiating hemolymph proteinase (HP14) contains five low-density lipoprotein receptor class A repeats, one Sushi domain, one Wonton domain and one proteinase catalytic domain.²¹⁷ Adding recombinant proHP14 to larval plasma greatly enhances proPO activation in response to *M. luteus*. The HP14 proenzyme, with its first domain truncated, was isolated from plasma of larvae injected with bacteria.⁹⁹ After incubation with β -1,3-glucan and β GRP1 or β GRP2, the proHP14 was converted to a two-chain active form, which significantly enhanced plasma proPO activation. The activation of proHP14 results from an autoactivation cleavage after Leu387, occurring when proHP14 interacts with β -1,3-glucan and β GRP. Characterization of individual domains and truncation mutants of HP14 showed that the amino-terminal regulatory region of HP14 participates in the specific binding of microbial polysaccharides and β GRP1.¹⁰⁰ Proteins orthologous to *M. sexta* HP14 also function at the top of proteinase cascades in immune responses of *Drosophila* and a beetle, *Tenebrio molitor*.^{218,219}

HP14 activates a clip domain proteinase HP21, which can then activate proPAP2 and proPAP3,^{220,221} resulting in activation of proPO. ProPAP1, which differs from proPAP2 and proPAP3 in having only one clip domain, is activated by HP6.¹⁸⁹ HP6 also functions in the proSpätzle activation pathway,^{188,189} providing cross-talk between these two immune cascades in *M. sexta*. Addition of active PAP1 to hemolymph stimulates the proteolytic activation of HP6, HP8, SPH1 and SPH2.²²² PAP1 directly activates proSPH2, but processing of the other precursors is probably indirect, depending on other plasma factors. Consequently, a minute amount of PAP1 added to plasma from naïve larvae stimulates a remarkably high level of PO activity in a short period of time, as a result of a positive feedback loop (Fig. 1). Some gaps in this pathway still need to be filled. To date, it is not clear which HP generates active HP6, leading to both PAP1-mediated melanization and HP8-mediated Toll pathway activation. The proteinase which activates SPH1 has not been identified and the possible involvement of HPs in Gram-negative bacteria-induced defense responses is not yet well understood.

INHIBITORY REGULATION OF HEMOLYMPH PROTEINASES BY SERPINS

Immune responses can produce molecules that are harmful to the host. Serine proteinases and the molecules they activate have potentially toxic effects. Proteinase inhibitors of different families can exist constitutively at relatively high levels in plasma of naïve insects and may also be produced in response to physiological or pathological stimuli.²²³ Serpins are ~50 kDa proteins, many of which are irreversible inhibitors and

key modulators of immune proteinase pathways.²²⁴ Serpins occur as plasma proteins in vertebrates and invertebrates. They have been purified and studied by molecular cloning from lepidopteran insects including *B. mori*,^{225,226} *M. sexta*,^{7,227-232} *H. cunea*,⁵⁶ *Mythimna unipuncta*,²³³ and *Mamestra configurata*.²³⁴ The *B. mori* genome contains 34 serpin genes, which have been analyzed with regard to molecular evolution of this gene family.^{13,235} Serpin biochemical and physiological functions in lepidopterans have been characterized most extensively in *M. sexta*.^{17,236}

The *M. sexta* serpin-1 gene encodes twelve protein isoforms, each having the same amino-terminal 336 residues and a variable region consisting of the carboxyl-terminal ~40 residues, including the reactive center loop that interacts with a serine proteinase during an inhibition reaction. The variable region is produced by mutually exclusive alternative splicing of twelve different versions of the ninth exon of the gene,^{237,238} resulting in a group of serpin proteins with diverse inhibitory selectivity.²²⁷ Serpin genes that employ alternative splicing at the same position to generate multiple serpin isoforms have been studied in other moth species. These include *B. mori* serpin genes 1 (3 isoforms) and 28 (4 isoforms),²³⁵ *M. configurata* serpin-1 (9 isoforms),²³⁹ and *Choristoneura fumiferana* serpin-1 (at least 4 isoforms).²⁴⁰ This mechanism for expanding serpin functional diversity, first discovered in Lepidoptera, been observed in other insect orders and in nematodes.²⁴¹

Physiological functions have been identified for a few of the *M. sexta* serpin-1 isoforms. Serpin-1A, -1E and -1J can inhibit HP8, and serpin-1J appears to be a physiologically relevant regulator of HP8 activity during immune responses⁸⁷ (An, Ragan, Kanost, unpublished results). Serpin-1J also inhibits all three PAPs to regulate proPO activation^{210,216} (Jiang, unpublished data). Serpin-1I can inhibit HP14.⁹⁹ Serpin-1K was identified in hemolymph in a complex with a midgut chymotrypsin,⁸⁷ suggesting a potential role for serpin-1 proteins in protection from digestive proteinases that escape into the hemocoel.

A putative orthologous group of serpins including *M. sexta* serpin-3,²²⁹ *B. mori* serpin-3,²³⁵ and an *H. cunea* serpin⁵⁶ are synthesized in response to infection and form a clade with *Drosophila* serpin-27A and *Anopheles gambiae* and *Aedes aegypti* serpin-2, which have immune regulatory functions.²³⁵ *M. sexta* serpin-3 contains a reactive site sequence (Asn-Lys-Phe-Gly) highly similar to the proteolytic activation site (Asn-Arg-Phe-Gly) in both proPO subunits. By mimicking these natural substrates, serpin-3 acts as an efficient inhibitor of all three PAPs.²²⁹ *M. sexta* serpin-4 and serpin-5²³⁰ are closely related to each other and form a clade with *B. mori* serpins 4, 5, 7, 8, 14, 31 and 32.²³⁵ Serpin-4 suppresses proPO activation by inhibiting HPs upstream of the PAPs, such as HP1, HP6 and HP21, while serpin-5 forms complexes with HP1 and HP6.²³¹ *M. sexta* and *B. mori* serpin-6 are apparent orthologs. *M. sexta* serpin-6 can inhibit PAP3 to block proPO activation and it also inhibits HP8 to potentially regulate the Toll pathway.²⁴²

LEPIDOPTERAN IMMUNE RESPONSES TO DIFFERENT TYPES OF INFECTION

Herbivorous lepidopteran caterpillars consume enormous amounts of plant diet and are capable of increasing their body weight up to 20% per day. Plant leaves harbor microbial communities, which enter the alimentary canal with the ingested food. The midgut of caterpillars can sense bacterial contamination of the diet and trigger immune responses, which are accompanied with life history tradeoffs.^{243,244} Bacteria can also naturally

enter and infect lepidopterans through wounds. Phagocytosis by granular hemocytes or plasmatocytes, depending on the species is probably the earliest response,²⁴⁵ and when numbers of bacteria are relatively low, can efficiently clear infection, particularly with bacteria of low virulence. Larger numbers of bacteria lead to hemocyte aggregation and formation of hemocyte nodules, probably aided by plasma pattern recognition proteins that agglutinate bacteria. Activation of lepidopteran hemocytes to become adhesive involves cytokines from the ENF family²⁴⁵ and eicosanoid signaling.⁴² Hemocyte nodules often become melanized, as products of PO polymerize to form a melanin coat around the aggregated hemocytes and bacteria. This response also generates quinones and reactive oxygen species that may help to kill the entrapped bacteria. The humoral response, synthesis of antibacterial peptides, occurs more slowly than the initial hemocyte response, requiring several hours before significant concentrations of antimicrobial molecules accumulate. This broad-spectrum antibacterial activity, comprised of a mixture of different antibacterial peptides, is an effective protective response that can last up to a few days.

Entomopathogenic fungi can invade insect hosts directly via their sclerotized chitinous integument. Penetration and lateral growth within the inner part of the integument is achieved by joint action of physical pressure and secreted enzymes among which proteinases play a predominant role.²⁴⁶ Most, if not all, entomopathogenic fungi develop in the hemocoel as cells known as protoplasts or hyphal bodies, which lack a fully developed cell wall. The absence of typical fungal cell wall components such as β -1,3-glucan may allow these fungi to evade the host immune surveillance. However, hyphal bodies of the entomopathogenic fungus *Metarhizium anisopliae* are ingested by plasmatocytes in *G. mellonella* during an early phase of infection,²⁴⁷ even though they lack β -1,3-glucan on their surface. Ingested hyphal bodies are not killed, but propagate and grow within phagocytic vacuoles of the plasmatocytes, which are likely occupied as a vehicle for dispersal within the hemocoel. Survival of hyphal bodies within the hemocytes as well as overcoming of multicellular encapsulation have been attributed to fungal secondary metabolites (toxins), such as destruxins and cyclosporins which suppress cellular and humoral responses within the infected hosts.^{248,249} Similar to bacteria, fungal cells are recognized, phagocytosed or, if too large or too numerous, encapsulated by hemocytes in the hemocoel of Lepidoptera.²⁵⁰ Eicosanoids have been implicated as mediators in cellular antifungal defense.²⁵¹ The lepidopteran antifungal response also encompasses proPO activation, production of reactive oxygen species,²⁵² and synthesis of potent antifungal peptides including cecropins²⁵³ and gallerimycin.²⁵⁴ However, the humoral responses upon infection with parasitic fungi such as *Beauveria bassiana* is different from that observed after challenge with bacteria.²⁵⁵

Larger eukaryotic parasites such as nematodes and parasitic wasps provoke hemocytic encapsulation and melanization,²⁴⁵ but little is understood about molecular mechanisms for recognizing such parasites as foreign.²⁵⁶ Successful parasites are able to disrupt or suppress the host insect's immune response. Entomopathogenic nematodes have a mutualistic relationship with virulent bacterial pathogens of insects, in which the bacteria produce virulence factors that disable cellular and humoral immune responses of their insect host.²⁵⁷ Parasitoid wasps that use lepidopteran larvae as hosts inject venom and accessory fluid components that disrupt the host immune system. In braconid and ichneumonid parasitoid wasps, this adaptation includes the injection of polydnviruses, which infect host hemocytes and express immunity-disrupting proteins, but do not replicate.²⁵⁸ Among these are gene products which cause apoptosis of hemocytes, disrupt signal transduction pathways in the humoral immune response and block melanization.²⁵⁹⁻²⁶¹

Some aspects of immunity to viral infection in lepidopterans are now becoming understood.²⁶² Baculoviruses are the most commonly studied viral pathogens of these insects. These viruses enter the larva by first infecting the gut epithelial cells. A response that can protect caterpillars from these infections is apoptosis of infected midgut cells. Infected cells die before viral replication can be completed, thus preventing spread of the virus to other cells or tissues.²⁶³ Baculoviruses encode gene products that inhibit caspases responsible for initiating apoptosis, allowing the infection cycle to proceed.²⁶³ Hemocytic encapsulation of infected tracheal cells is another immune response to baculoviruses that has been observed in lepidopterans.²⁶² In addition, hemolymph PO is correlated with virucidal activity toward baculoviruses,²⁶⁴ and thus, the proPO activation cascade may help protect against baculoviral infection.

CONCLUSION

Lepidopteran insects have some important advantages as model systems for immunological research, including a depth of knowledge developed so far and the availability of large hemolymph samples from individual insects for studies of hemocytes and plasma proteins. With the recent exception of *B. mori*, studies on moths and butterflies have been hampered by a lack of genomic information, which would facilitate proteomics investigations and also lead to more ready identification of candidate genes for experimental study. This situation is likely to change dramatically in the next few years, as it is anticipated that genome sequences for several additional lepidopteran species will soon become available. Transgenic technology for silkworms is now well developed and may yield new fundamental information on immunity and perhaps strains with improved disease resistance. Furthermore, more complete understanding of lepidopteran immune responses could lead to future developments of enhanced strategies for regulating insect pest populations through use of specific pathogens and parasites.

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CHAPTER 11

FLY IMMUNITY: Recognition of Pathogens and Induction of Immune Responses

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Abstract: Despite the lack of adaptive immunity based on gene rearrangement such as that in higher vertebrates, flies are able to defend themselves from a wide array of pathogens using multiple innate immune responses whose molecular mechanisms are strikingly similar to those of the innate immune responses of other multicellular organisms, including humans. Invading pathogens passing through the epithelial barriers, the first line of self-defense, are detected by pattern recognition receptors that identify pathogen-associated molecular patterns in the hemolymph or on the immune cell surface and are eliminated by humoral and cellular responses. Some pathogens escape recognition and elimination in the hemolymph by invading the host cell cytoplasm. Some of these intracellular pathogens, however, such as *Listeria monocytogenes*, are identified by pattern recognition receptors in the cytoplasm and are eliminated by intracellular responses, including autophagy, an intracellular degradation system. Although some of these pattern recognition receptors are encoded in the germ-line as protein families, another type of receptor in the immunoglobulin-superfamily is extensively diversified by alternative splicing in somatic immune cells in *Drosophila*.

INTRODUCTION

Diseases of the silkworm, *Bombyx mori*, were already described in the classical Chinese literature “Guanzi” by Master Guan in the 7th century BCE and honeybee diseases were described in Greek mythology as early as the 6th century BCE. The analysis of insect immune responses, however, did not begin until the 19th century. In 1870, Louis Pasteur pioneered the study of insect immune responses in *Bombyx*.¹ His work was followed in

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1892 by Elie Metchnikoff's investigations on cellular immunity in insects.² Some of their observations on insect immune responses have been driven further and in some cases the molecular mechanisms explaining their observations have been elucidated.³ Studies of a model organism, the fruit fly *Drosophila melanogaster*, have greatly contributed to a deeper understanding of the detailed molecular mechanisms of innate immune responses due to the powerful genetic tools available for this organism and these investigations have revealed fundamental conservation between insect immunity and innate immunity of other multicellular organisms, including humans.

The frontline of fly host defense are the epithelial tissues, such as the epidermis, gut and trachea, which act not only as mechanical barriers but also produce defense molecules such as lysozymes, antimicrobial peptides and reactive oxygen species.⁴ Because flies have open vascular systems, pathogens passing through the epithelial barrier encounter cellular and humoral defense responses in the hemolymph.⁵ As a cellular response, immune cells called hemocytes phagocytose and encapsulate pathogens in the hemolymph. Humoral reactions are induced by the activation of cascades of constitutive proteins present in the hemolymph, such as the prophenoloxidase (proPO) cascade and the coagulation cascade and the activation of intracellular signaling pathways that produce defense proteins such as antimicrobial peptides in immune-responsive tissues and cells, such as hemocytes and the fat body, the functional equivalent of the vertebrate liver. This chapter describes the molecular mechanisms of fly immune responses, mainly the invading pathogen-recognition mechanisms in the hemolymph and in the cytoplasm of *Drosophila* immune cells. The immune responses are activated via specific recognition of pathogen-associated molecular patterns by germ-line encoding pattern recognition receptor families⁶ such as the peptidoglycan recognition protein (PGRP) family.

PRIMARY HUMORAL RESPONSES: ACTIVATION OF CONSTITUTIVE PROTEIN CASCADES IN THE HEMOLYMPH

In his study of *Bombyx* disease, Pasteur observed melanization of the insect cuticle caused by the protozoan pathogen, *Nosema bombyosis*.¹ Melanization of the hemolymph of the wax moth larvae *Galleria mellonella* has been reported by several authors, including Serge Metalnikov in 1927,⁷ who speculated that the dark pigment was formed by the action of a special enzyme on a chromogenic substrate in the insect hemolymph, which is now supported by molecular level studies. The enzyme is phenoloxidase and the chromogenic substrates in the insect hemolymph are tyrosine and dopa. Phenoloxidase is a copper-containing enzyme that catalyzes two types of reactions: the oxidation of monophenols to orthodiphenols and the oxidation of orthodiphenols to orthoquinones, which are key reactions required for melanin formation.⁸ Melanization has an important role in arthropod immune reactions such as wound healing, surrounding invading pathogens and production of intermediates toxic to invading microorganisms.⁹ Detailed biochemical mechanisms of the proPO cascade in moth, beetle and crustacean are discussed in chapters by Kanost, Lee, and Cerenius.

Phenoloxidase was purified from several arthropods, including *Bombyx mori* and is synthesized as inactive zymogens called proPO.^{10,11} *Drosophila* has three genes that encode proPO, *Black cells/Dox-A1*, *Dox-A3* and *CG8193*.^{12,13} Activation of proPO in the hemolymph is mediated by the proPO cascade, which consists of some serine proteases and some pattern-recognition receptors that recognize invading microorganisms.⁸ As

pattern recognition receptors, PGRP and β -1,3-glucan binding protein (β GRP) were first purified from *Bombyx* hemolymph by Masaaki Ashida and his colleagues.^{14,15} PGRP and β GRP specifically bind to peptidoglycan and β -1,3-glucans, cell-wall components of bacteria and fungi, respectively and activate the proPO cascade in vitro. cDNA cloning of these proteins revealed that these proteins are evolutionarily conserved; e.g., there are 13 PGRP family members in *Drosophila*, 7 in *Anopheles* and 4 in both mouse and humans.¹⁶⁻²¹ The PGRP family has a PGRP-domain in the C-terminal region that has some amino acid sequence similarity to peptidoglycan-degrading enzymes with *N*-acetylmuramyl-alanine amidase activity, such as bacteriophage lysozymes. β GRP has sequence similarity with bacterial β -1,3-glucanase in the C-terminal region, which is also in Gram-negative binding protein (GNBP) identified from *Bombyx* hemolymph.^{22,23} In *Drosophila*, one PGRP family member, PGRP-LE, which recognizes diaminopimelic acid (DAP)-containing peptidoglycans, is involved in activation of the proPO cascade upstream of a serine protease called prophenoloxidase activating enzyme (PPAE), which is regulated by a serine protease inhibitor, Serpin 27A.²⁴⁻²⁶ Two Clip domain-containing serine proteases,²⁷ melanization protease (MP)1 and serine protease (Sp) 7/MP2/PAE1, activate the proPO cascade upstream of PPAE and are also regulated by Serpin 27A.²⁸⁻³⁰ MP1 is required to activate the proPO cascade in response to both bacterial and fungal infections, whereas Sp7 is mainly involved in response to fungal infections.²⁹ Coagulation to initiate wound healing and limit hemolymph loss is also induced at the site of injury in a proPO cascade-independent manner.^{31,32} In *Drosophila*, a hemocyte expressing protein, Hemolectin and an abundant hemolymph protein, Fondue, are required for efficient coagulation.³³⁻³⁵ Transglutaminase acts in coagulation.³⁶ Hemolectin has similarity with the human von Willebrand factor, which is defective in some coagulation diseases in humans, suggesting a conserved coagulation cascade in fly and humans.³⁷

SECONDARY HUMORAL RESPONSES: INDUCTION OF SELF-DEFENSE MOLECULES

In 1906, Metalnikov observed bacteriolysis of *Mycobacterium tuberculosis* in the hemolymph of *Galleria*,³⁸ which was followed by studies on insect bacteriolysins by Andre Paillot.³⁹ We now know that bacteriolysis is explained by the activity of lysozymes and antimicrobial peptides. A lysozyme is an enzyme that catalyzes the cleavage of the β -1,4-glycoside bonds in the peptidoglycans of the bacterial cell wall, which is the main antibacterial factor in insects as well as in vertebrates. Since 1981 when antimicrobial peptides were first identified in the cecropia moth, *Hyalophora cecropia*, by Hans Boman and coworkers,⁴⁰ several hundreds of antimicrobial peptides have been isolated from various invertebrates, plants and mammals, including humans.⁴¹ *Drosophila* mutants that fail to produce antimicrobial peptides are susceptible to bacterial and fungal infections, indicating that the induction of antimicrobial peptides is crucial for host defense in fly.⁴²

Shunji Natori and his coworkers isolated several antimicrobial peptides from the flesh fly *Sarcophaga peregrina*, which is ideally suited for biochemical analysis. In response to infection or injury, four different types of antibacterial peptides are synthesized in the fat body and secreted into the hemolymph.⁴³ These are the Sarcotoxin I, II, III families and the Sapecin family. The Sarcotoxin I family has potent bactericidal activity against mainly Gram-negative bacteria.⁴⁴⁻⁴⁶ The Sarcotoxin I family comprises five subtypes, each consisting of 39 or 40 amino acids, which have slightly different target specificities.⁴⁷

Table 1. Summary of antimicrobial peptides in *Sarcophaga*

Effector Molecules	Molecular Mass	Subtypes	Primary Target Specificity	<i>Drosophila</i> Homologue
Sarcotoxin I	4000	5	Gram-negative	Cecropin
Sarcotoxin II	24000	3	Gram-negative	Attacin
Sarcotoxin III	8000	>2	Gram-negative	Diptericin
Sapecin	4000	3	Gram-positive	Defensin
Antifungal protein	7000	1	Fungi	Non

The characteristics of *Sarcophaga* antimicrobial peptides and their homologues in *Drosophila* are summarized in reference 108.

Many insect antimicrobial peptide families have several subtypes, suggesting that insects utilize antimicrobial peptide family subtypes with different target specificities to eliminate various microbes (Table 1). Sarcotoxin IA consists of two amphiphilic alpha-helical regions with a hinge region.⁴⁸ The amphiphilic helical segments are important for the expression of the antibacterial activity, suggesting that Sarcotoxin IA acts primarily on the bacterial membrane.⁴⁹ Cholesterol-containing liposomes are less sensitive to Sarcotoxin IA, explaining the selective toxicity of Sarcotoxin IA to bacteria because the membranes of the host cells contain cholesterol whereas those of bacteria do not. Although the Sarcotoxin II and III families also act on Gram-negative bacteria, Sarcotoxin families have different target specificities. The Sarcotoxin I family is similar to Cecropin, initially discovered in the hemolymph of bacteria-challenged diapausing pupae of *Hyalophora*.⁴⁰ The Sarcotoxin II and III families are similar to *Drosophila* Attacin and Diptericin, respectively (Table 1). In contrast to the Sarcotoxin families, Sapecin acts mainly on Gram-positive bacteria, including methicillin-resistant *Staphylococcus aureus*.⁵⁰ Sapecin has similarity to *Drosophila* Defensin. Sapecin comprises 40 amino acids with three internal disulfide bridges and has high affinity for cardiolipin, which is a major phospholipid of *S. aureus*. A mutant of *Escherichia coli* defective in cardiolipin synthesis is more resistant to Sapecin than wild-type *E. coli*, suggesting that cardiolipin is a target of Sapecin.⁵¹ Sapecin was originally purified from the culture medium of a *Sarcophaga* embryonic cell line and is expressed during the embryonic stage of development, suggesting a role of Sapecin in development.⁵² Consistent with this idea, Sapecin is involved in the proliferation of embryonic cells of *Sarcophaga*.⁵³ In addition to the inducible antibacterial peptides, the constitutive protein Antifungal protein, which has no similarity to other known proteins, is also present in the hemolymph.⁵⁴ The amino acid composition of Antifungal protein is unique: glycine and histidine account for more than half of the total amino acids. Antifungal protein has no bactericidal activity, but it has lethal activity against fungi such as *Schizosaccharomyces pombe*. Antifungal protein has less activity against *Candida albicans*, but the activity is greatly enhanced by the presence of Sarcotoxin IA, an inducible antibacterial peptide, suggesting that *Sarcophaga* constructs an efficient system with constitutive Antifungal protein and inducible antibacterial peptides to eliminate various microbes.

In *Drosophila*, seven classes of antimicrobial peptides have been identified. Diptericin, Attacin and Drosocin are effective against Gram-negative bacteria, whereas Defensin is active against Gram-positive bacteria. Cecropin has wide specificity, acting on both

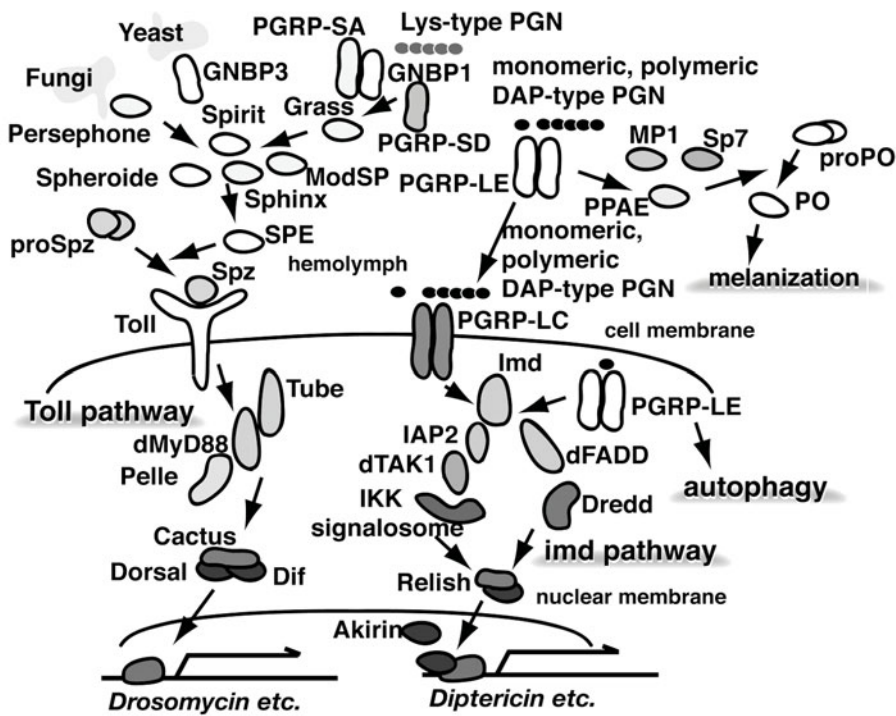


Figure 1. Recognition of various pathogens and induction of immune responses in *Drosophila*. Activation of the proPO cascade leading to melanization is induced by the cleavage of proPO to PO by a serine protease, PPAE, which is regulated by two serine proteases, MP1 and Sp7. A constitutive hemolymph protein, PGRP-LE, binds to monomeric and polymeric DAP-type peptidoglycans (PGNs), which are components of many Gram-negative and some Gram-positive bacteria and activates the proPO cascade upstream of PPAE. PGRP-LE is also involved in activating the imd pathway-dependent induction of antibacterial peptides in the hemolymph and inside the immune cells. Membrane PGRP-LC is required for monomeric and polymeric DAP-type PGN-mediated activation of the imd pathway. The imd pathway comprises Imd, IAP2, dFADD, Dredd, dTAK1, IKK signalosome, Akirin and Relish. PGRP-LE induces autophagy through intracellular recognition of DAP-type peptidoglycans in an imd independent manner. PGRP-SA in the hemolymph binds to Lys-type PGN of Gram-positive bacteria and is required for activation of the Toll pathway in cooperation with GNBPs. GNBPs are involved in yeast-mediated activation of the Toll pathway. The activation of the Toll pathway is mediated by its active ligand, Spz, cleaved from proSpz by the serine protease cascade, including the SPE, Spheroid, Spirit, Sphinx, Persephone, Grass and ModSP. The Toll pathway consists of Toll, dMyD88, Tube, Pelle, Dorsal and Dif. The transcriptional factors, Dorsal and Dif, are negatively regulated by Cactus. This figure was modified from Kurata S., Research Signpost, Copyright 2008.¹⁰⁸

bacteria and fungi. Drosomycin and Metchnikowin, identified so far only in Drosophilidae, act on fungi.⁵⁵ Induction of antimicrobial peptides is mediated by two distinct pathways, the imd pathway and the Toll pathway (Fig. 1),^{4,5} which are mechanistically similar to the mammalian tumor necrosis factor- α receptor signaling pathway and Toll-like receptor (TLR)/interleukin-1 receptor signaling pathway, respectively.⁵⁶ The imd pathway regulates the synthesis of several antibacterial peptides such as Diptericin, which is mediated by cell surface PGRP-LC, Imd adaptor protein, IAP2, dTAK1 kinase, I κ B kinase (IKK)

complex, FADD, Dredd caspase, Akirin and a Rel transcriptional factor, Relish.⁵⁷⁻⁶⁵ The tumor necrosis factor- α signaling pathway comprises receptor-interacting protein, TAK1 kinase, IKK complex, caspase-8 and nuclear factor κ B (NF- κ B), similar to the Imd, dTAK1, dIKK complex, Dredd and Relish, respectively, in the *Drosophila* imd pathway.⁵⁶ The Toll pathway predominantly regulates induction of the antifungal peptide Drosomycin, which is mediated by the Toll receptor, dMyD88 adaptor protein and Dorsal related immune factor, similar to TLR, MyD88 and NF- κ B in the mammalian TLR signaling pathway.⁶⁶⁻⁶⁹ The important roles of these signaling pathways in host defense systems are demonstrated by the fact that mutations in the genes that regulate these pathways dramatically decrease resistance to microbial infections, e.g., *Toll* mutants are susceptible to fungal infections and *Relish* mutants lose resistance to Gram-negative bacterial infections.^{63,66}

RECOGNITION OF VARIOUS PATHOGENS IN THE HEMOLYMPH

The Toll pathway is activated predominantly in response to fungal and some Gram-positive bacterial infections, whereas the imd pathway is activated predominantly in response to Gram-negative and other Gram-positive bacterial infections.⁷⁰ Therefore, *Drosophila* distinguishes different pathogens by specific mechanisms. In contrast to mammalian TLRs, which recognize pathogen components such as lipopolysaccharides,⁷¹ the ligand of the *Drosophila* Toll receptor is Spätzle (Spz), which is produced by proteolysis of an endogenous protein (proSpz) in response to infection.⁷² Therefore, *Drosophila* Toll does not act as a pattern recognition receptor, but rather mediates the downstream signaling of such receptors. In *Drosophila*, some PGRP and GGBP family members act as pattern recognition receptors upstream of both the Toll and imd pathways.^{4,73}

PGRP-SA, PGRP-SD, GGBP1 and GGBP3 act as pattern recognition receptors upstream of the Toll pathway in *Drosophila*. PGRP-SA, a hemolymph protein that was identified from a loss-of-function mutant screen, is required for activation of the Toll pathway in response to Gram-positive bacterial infections.⁷⁴ PGRP-SA cooperates with GGBP1 to activate the Toll pathway in response to Gram-positive bacterial infections.⁷⁵ PGRP-SD has some redundancy and recognizes Gram-positive bacteria with PGRP-SA and GGBP1.⁷⁶ GGBP3 is required for activation of the Toll pathway in response to fungal infections.⁷⁷ Downstream of these recognition receptors, pro-Spz is cleaved by a serine protease, Spätzle processing enzyme (SPE).⁷⁸ The SPE-mediated processing is regulated by several serine proteases such as Persephone, Spirit, Spheroid, Sphinx, Grass and ModSP.⁷⁹⁻⁸² Persephone and Grass are specifically required for the signaling from fungi and Gram-positive bacteria, respectively and other proteases are required for the signaling from both microorganisms.⁸⁰

Upstream of the imd pathway, PGRP-LC and PGRP-LE act as pattern recognition receptors. PGRP-LC, a membrane-associated PGRP-family member, is required for activation of the imd pathway in response to Gram-negative bacterial infections.⁸³⁻⁸⁵ Susceptibility of the *PGRP-LC* mutant to Gram-negative bacterial infections is less dramatic than that of an IKK-complex mutant, in which signaling is impaired downstream of imd, suggesting that there is an activator of the imd pathway in addition to PGRP-LC.⁸³ Consistent with this finding, PGRP-LE, a constitutive hemolymph protein, selectively activates the imd pathway.²⁴ The *PGRP-LC/PGRP-LE* double-mutant is much more susceptible to *E. coli* infection than either single mutant alone, suggesting that PGRP-LC and PGRP-LE are

major recognition receptors for Gram-negative bacteria.²⁵ As described above, PGRP-LE activates the proPO cascade, which branches from the PGRP-LE-mediated activation of the imd pathway upstream of PPAAE.²⁵ In addition to the extracellular functions of PGRP-LE in the hemolymph, PGRP-LE acts as a coreceptor of the membrane receptor PGRP-LC on the surface of immune cells and as an intracellular receptor in immune reactive cells to induce antibacterial peptides by activating the imd pathway.⁸⁶

Peptidoglycans are an essential cell wall component of almost all bacteria except mycoplasma, which lack a cell wall. Peptidoglycan is a polymer of β -1,4-linked *N*-acetylglucosamine (GlcNAc) and *N*-acetylmuramic acid (MurNAc) crosslinked by short stem peptides. The glycan chain is relatively conserved in all bacteria, whereas the amino acid composition of the crosslinking stem peptides and the linkages between stem peptides are diversified, depending on the bacterial species.⁸⁷ Many Gram-negative bacteria and some Gram-positive bacteria, such as *Bacillus* species, directly crosslink DAP-containing peptidoglycans and preferentially activate the imd pathway in *Drosophila*. Many Gram-positive bacteria have peptidoglycans containing lysine instead of DAP with crosslinking peptides between stem peptides and these bacteria preferentially activate the Toll pathway. Consistent with the finding that PGRP-LE selectively activates the imd pathway in vivo, PGRP-LE selectively binds DAP-type peptidoglycans, but not Lys-containing peptidoglycans in vitro.²⁴ *PGRP-LC* is required for DAP-type peptidoglycan-mediated activation of the imd pathway in vivo and in vitro.^{88,89} The minimum structure of the DAP-type peptidoglycan required for PGRP-LC-mediated activation of the imd pathway is GlcNAc-MurNAc with an internal 1,6-anhydro bond attached to a tripeptide containing DAP, known as tracheal cytotoxin.⁹⁰ These results are consistent with the findings that PGRP-LC and PGRP-LE act synergistically to resist bacteria with DAP-type peptidoglycans such as *E. coli* and *Bacillus megaterium*.²⁵ The minimum structure of the Lys-type peptidoglycan required for PGRP-SA-mediated activation of the Toll pathway comprises two units consisting of GlcNAc-MurNAc attached to Lys-containing tetrapeptides that are covalently dimerized by an interpeptide.⁹¹ Therefore, *Drosophila* PGRP family members distinguish the structural diversity of peptidoglycans and activate the appropriate immune responses. In addition to PGRPs acting as pattern recognition receptors, there are PGRP family members such as PGRP-SC and PGRP-LB that have enzyme activity that hydrolyzes the lactylamide bond between the glycan strand and the stem peptides of peptidoglycans.⁹²⁻⁹⁴ The *N*-acetylmuramyl-alanine amidase activity is consistent with the structural similarity of the PGRP-domain to *N*-acetylmuramyl-alanine amidase. The degraded peptidoglycans lose their elicitor activity, suggesting a scavenger function of the enzyme PGRP family members and negative feedback regulation of immune responses by enzyme PGRP family members.⁹⁵

RECOGNITION AND ELIMINATION OF INTRACELLULAR BACTERIA IN THE CYTOPLASM

Intracellular pathogens, a diverse group of organisms that cause serious diseases such as tuberculosis and malaria, invade the cells and escape serum and cell surface innate immune receptors. The intracellular function of PGRP-LE is suggested by the findings that the delivery of tracheal cytotoxin into S2 cells, the *Drosophila* macrophage-like cell line, induces antimicrobial peptides in a PGRP-LE dependent manner and PGRP-LE induces antimicrobial peptides in a cell autonomous manner in the Malpighian tubules,

the functional equivalent of the mammalian kidney. Consistent with these results, the *PGRP-LE* mutant is susceptible to infection by *Listeria monocytogenes*, intracellular bacteria with DAP-containing peptidoglycans recognized by PGRP-LE.⁹⁶ In contrast to Gram-negative bacterial infections, the survival rate of the double mutant *PGRP-LE/PGRP-LC* infected with *L. monocytogenes* is similar to that of the *PGRP-LE* mutant, suggesting that PGRP-LE and PGRP-LC do not have redundant functions in producing resistance to *L. monocytogenes*.⁹⁶ In vitro infection experiments using hemocytes and PGRP-LE expressing S2 cells revealed that PGRP-LE colocalizes with *L. monocytogenes* in the cytoplasm and is essential for resistance against intracellular bacterial growth, but, interestingly, inhibition of intracellular bacterial growth is not dependent on the known innate immune signaling pathways, the Toll and imd pathways.

Autophagy was initially reported as a bulk self-degradation mechanism for the turnover of proteins and organelles and is conserved from yeast to humans.⁹⁷ Recent studies, however, revealed that autophagy has a role in many biological events, such as nutrient supply upon starvation, cell death and cancer. Autophagy also functions as an innate immune response against intracellular bacteria, viruses and parasites in cultured cells.⁹⁸⁻¹⁰⁴ In *Drosophila*, autophagy is induced in hemocytes in response to *L. monocytogenes* infection in a PGRP-LE dependent manner, which is crucial for host survival against *Listeria* infection as well as for the inhibition of bacterial growth in hemocytes. PGRP-LE is also required for DAP-type peptidoglycan-mediated autophagy, but not for Lys-type peptidoglycan-mediated autophagy in hemocytes, suggesting the existence of another intracellular sensor for bacteria with Lys-type peptidoglycans. Therefore, PGRP-LE has an essential role in detecting intracellular bacteria through DAP-type peptidoglycans to induce autophagy as an innate immune response, which indicates a direct link between pathogen recognition and the induction of autophagy in the cytoplasm (Fig. 1).

EXTENSIVE SOMATIC DIVERSIFICATION OF A RECOGNITION MOLECULE

Pattern recognition receptors acting on innate immunity, such as PGRP family members and GGBP family members, are encoded in the germ line as protein families and distinguish between different types of microorganisms. On the other hand, in *Drosophila*, the diversity of an immunoglobulin-superfamily receptor Down syndrome cell adhesion molecule (*Dscam*) is generated by alternative splicing in hemocytes.¹⁰⁵ The *Dscam* gene contains three arrays of alternative exons, exons 4, 6 and 9 and two alternative transmembrane domains, in addition to 20 constant exons, which have the potential to express *Dscam* with 19008 different extracellular domains.¹⁰⁶ Many of these different isomers are expressed in the fat body, hemocytes and brain and different *Dscam* isomers bind to bacteria with different affinities, suggesting a role for *Dscam* isomers in the recognition of different types of pathogens.¹⁰⁵ Moreover, *Dscam* is required for bacterial phagocytosis by hemocytes.¹⁰⁵ The *Dscam* gene in the mosquito *Anopheles gambiae* also has a complex organization with 101 exons and the potential to express *AgDscam* with 15960 different extracellular domains.¹⁰⁷ *AgDscam* is involved in host defense against bacterial infections by producing pathogen-specific splice form repertoires.¹⁰⁷ Therefore, the diversity in insect *Dscam* and in vertebrate antigen receptors is generated by alternative splicing and gene rearrangements to achieve similar goals, which is the recognition of a multitude of pathogens with adaptive characteristics.

CONCLUSION

In fly immunity, invading pathogens are recognized in the hemolymph, on the immune cell surface, and in the host cell cytoplasm by pattern recognition receptors, such as PGRP family members that distinguish the structural diversity of bacterial peptidoglycans, and eliminated by humoral and cellular responses, such as induction of antimicrobial peptides and autophagy, respectively.

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CHAPTER 12

MOSQUITO IMMUNITY

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Abstract: Throughout their lifetime, mosquitoes are exposed to pathogens during feeding, through breaks in their cuticle and following pathogen-driven cuticular degradation. To resist infection, mosquitoes mount innate cellular and humoral immune responses that are elicited within minutes of exposure and can lead to pathogen death via three broadly defined mechanisms: lysis, melanization and hemocyte-mediated phagocytosis. This chapter reviews our current understanding of the mosquito immune system, with an emphasis on the physical barriers that prevent pathogens from entering the body, the organs and tissues that regulate immune responses and the mechanistic and molecular bases of immunity.

INTRODUCTION

Mosquitoes (Diptera: Culicidae), like all organisms, are under constant threat of infection. For the continuation of their life cycles, females of all anautogenous species are required to take a blood meal for the production of eggs. This act of blood feeding often exposes mosquitoes to blood-borne pathogens that aim to undergo complex developmental, reproductive and/or migrational processes inside a mosquito host before they can be transmitted during a subsequent blood meal. In addition to risking infection through blood feeding, mosquitoes often acquire pathogens through sugar feeding, through breaks in their cuticle that are created after physical injury and following pathogen-driven cuticular degradation. Whereas pathogen acquisition through blood feeding occurs exclusively during the adult life stage, infection through the cuticle is likely most prevalent during the aquatic developmental stages, when mosquitoes live in environments rife with bacteria.

Although culicine and anopheline mosquitoes are effective vectors of human and animal pathogens, susceptibility or resistance to infection is often the result of intricate co-evolutionary

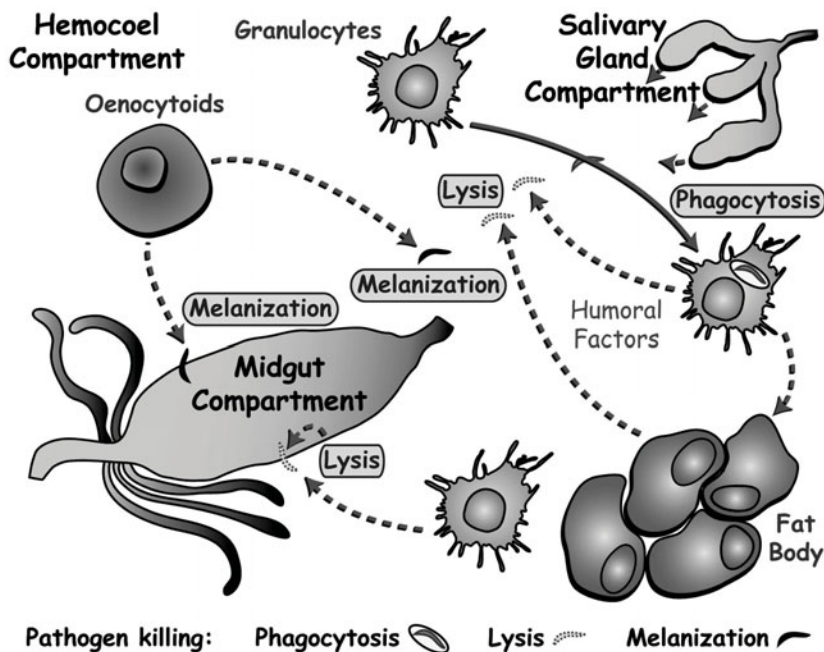


Figure 1. Mosquito immune responses in the three major immune compartments. In the hemocoel, granulocyte and oenocytoid hemocytes, as well as fat body, kill pathogens via phagocytosis, lysis and melanization. In the midgut, immune factors produced by epithelial cells, hemocytes and possibly fat body, kill pathogens via lytic and melanization pathways. Little is known about the role salivary glands play in immune responses, but they produce immune factors in response to infection.

processes in which mosquitoes and pathogens engage in counter-adaptations for survival and infection. As a consequence of this evolutionary arms race, only mosquitoes of the genus *Aedes* are capable of transmitting dengue fever virus and only *Culex* mosquitoes transmit Japanese encephalitis virus.¹ Similarly, of the greater than 3,000 known species of mosquitoes, only a subset of species from the genus *Anopheles* is capable of transmitting human malaria.² Even within the susceptible *Anopheles gambiae* species, some individuals are resistant to infection and others, while unable to eliminate the infection, are capable of drastically reducing pathogen numbers.^{3,4} The specificity of mosquito-pathogen associations also varies among species of parasites. For example, the mosquito *Armigeres subalbatus* effectively transmits the filarial nematode *Brugia pahangi* but is resistant to a close relative, *Brugia malayi*.⁵ Several factors account for the ability of pathogens to survive inside mosquitoes, including behavior (e.g., will the mosquito encounter the pathogen?) and physiological compatibility (e.g., are the correct conditions present in the host that allow the pathogen to complete its life cycle?). Another major factor that determines whether a pathogen can survive inside mosquitoes rests on the strong innate immune responses mounted by the host and on whether the pathogens have evolved mechanisms to evade these defenses.

In broad terms, pathogen killing by mosquitoes is accomplished by three primary mechanisms: cell-mediated phagocytosis, melanization and lysis (Fig. 1). Each is initiated by pattern recognition receptors and the factors leading to killing can be subdivided into cellular and humoral components (Fig. 2). The cellular response includes phagocytosis

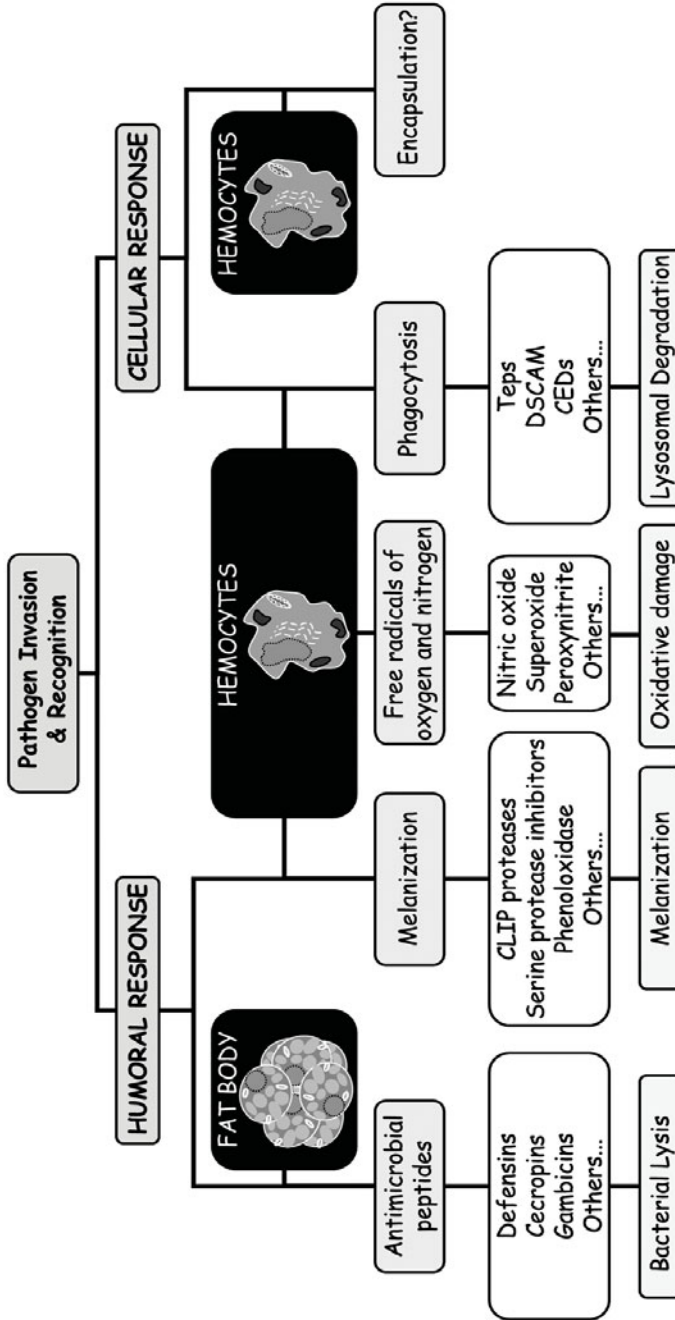


Figure 2. Immune responses in the mosquito hemocoel. Antimicrobial responses in the hemocoel are mediated by hemocytes and fat body and include antimicrobial peptides, melanization, reactive oxygen species, reactive nitrogen species and phagocytosis.

and encapsulation by hemocytes and pericardial cells.⁶⁻⁹ The humoral response includes pattern recognition receptors, inducible antimicrobial peptides, the phenoloxidase cascade system of melanization and wound healing, and reactive oxygen and reactive nitrogen intermediates.¹⁰⁻¹⁵ Regardless of this conceptual organization, the line between cellular and humoral immunity is blurred because many humoral components are produced by hemocytes and participate in cellular immune responses.^{16,17} This chapter reviews the interactions between mosquitoes and pathogens, with emphasis placed on the physical barriers that prevent pathogens from entering the body, the organs and tissues that regulate immune responses, the mechanistic manifestations of immunity and our current understanding of the molecular basis of immunity.

BIOLOGY OF PATHOGENS INSIDE MOSQUITOES

Mosquitoes are subject to infection by viral, bacterial, fungal, protozoan and metazoan pathogens and initial entry into the host generally occurs either through breaks in the cuticle or by ingestion. The biology of pathogens inside the mosquito is dependent on their mode of transmission. By and large, bacteria and fungi enter mosquitoes through wounds in their outer cuticle or through the midgut epithelium after feeding. They quickly replicate in the host's gut or hemocoel (body cavity) and can be transmitted to a subsequent host while the initial host is alive or after its death. The time between colonization and transmission can be very short; some of these infectious agents are highly pathogenic and are lethal within hours of infection.¹⁸ For that reason, bacterial and fungal pathogens are currently being used in the development of novel pest control strategies.^{19,20}

Pathogens acquired and transmitted through blood feeding, on the other hand, must undergo obligatory processes that require their interaction with multiple tissue types and require that the host survives for days or weeks before transmission can take place.²¹ *Plasmodium* parasites and arboviruses, for example, must cross the midgut epithelium, replicate, migrate through the hemocoel and invade the salivary glands before the mosquito can infect a subsequent host during her next blood feeding. Similarly, filarial nematodes must leave the midgut, develop in the thoracic musculature or Malpighian tubules and migrate to the mouthparts for the mosquito to become infectious. Unlike fungal and bacterial pathogens, the transmission of blood-borne pathogens requires mosquito viability throughout the entirety of the pathogen's life cycle inside the insect host. Nevertheless, these pathogens decrease fitness, reduce fecundity and, if acquired in large enough numbers, can be lethal to the mosquito.²²⁻²⁵

MOSQUITO COMPARTMENTS AND BARRIERS TO INFECTION

Pathogens inhabit three primary compartments in the mosquito: the midgut, the hemocoel and the salivary glands (Fig. 1). All three of these compartments include physical and physiological barriers that limit or reduce pathogen development. In addition, cells in all of these compartments produce immune factors with antimicrobial activity. These responses, though powerful, are innate and lack the properties of somatic hypermutation that are hallmarks of vertebrate adaptive immunity.²⁶

The Gut Compartment

When pathogens enter mosquitoes via ingestion, the initial barrier faced is physical destruction by the cibarial armature.²⁷ This barrier, composed of sclerotized teeth and spines that protrude into the lumen of the foregut, slices large pathogens during the initial stages of ingestion and before they reach the midgut. The cibarial armature is effective in limiting infection by large metazoan parasites such as filarial nematodes but does little to destroy protozoan, bacterial and viral pathogens. Once in the midgut lumen, pathogens must survive digestive enzymes and invade the midgut epithelium by either digesting a thick acellular chitinaceous peritrophic matrix formed in response to blood feeding or by initiating epithelium invasion prior to the formation of this matrix.²⁸⁻³⁰ The mechanisms leading to midgut penetration are not well understood, but depending on the pathogen may involve receptor-ligand interactions, physical burrowing through the epithelium and/or digestion of host cells.³¹⁻³³

Mosquitoes drastically limit pathogen development in the midgut. For *Plasmodium* parasites, the causative agents of malaria, ookinete development in the gut results in a 500 to 100,000-fold reduction in parasites numbers and the ookinete to oocyst transformation that occurs on the basal side of the midgut experiences parasite losses of 5 to 100-fold.³⁴ The bases for these parasite reductions are complex, but include lytic and melanization events that are controlled by pattern recognition receptors, serine proteases and their inhibitors, and enzymatic cascades.³⁵⁻⁴⁰ Indeed, molecular and biochemical studies have shown that the midgut rapidly produces a vast milieu of antimicrobial proteins in response to pathogen exposure.⁴¹⁻⁴⁵ In addition to these midgut-produced immune factors, infection of this organ triggers the production of immune proteins in other tissues, some of which are transported into the midgut where they exert their antimicrobial activities.^{36,38,46,47}

The Hemocoel Compartment

The hemocoel compartment is an open body cavity that contains all visceral organs and is delineated by the outer cuticle and the basal lamina surrounding internal tissues. For many reasons, including preventing infection, insects have developed a tightly-sealed hydrophobic outer cuticle that shields internal organs from the outside environment. Breaks in this cuticle form temporary openings through which pathogens may enter. While these wounds commonly occur in nature, coagulation and melanization responses involving wound contraction, hemocyte degranulation and scar formation rapidly close these lesions.^{48,49} As described above, pathogens also enter the hemocoel through ingestion followed by midgut penetration. Regardless of the mode of entry, pathogens disseminate throughout the hemocoel by either pathogen-driven active motility or the natural flow of hemolymph.⁵⁰⁻⁵³

Once in the hemocoel, pathogens are immersed in a nutrient-rich medium that contains immune cells and humoral immune factors produced by hemocytes, pericardial cells and fat body. Hemocytes are immunosurveillance cells that initiate innate immune responses and are found circulating with the hemolymph or attached to visceral tissues (Fig. 3). They are involved in the killing and sequestration of pathogens via phagocytosis, nodulation and the secretion of humoral immune factors.^{6-9,12,54} Transcriptomic analyses in several mosquito genera have repeatedly shown the broad range of immune factors produced by hemocytes, which include pattern recognition receptors, proteins involved in phagocytosis, melanization modulators and enzymes, signal transduction proteins, stress response proteins

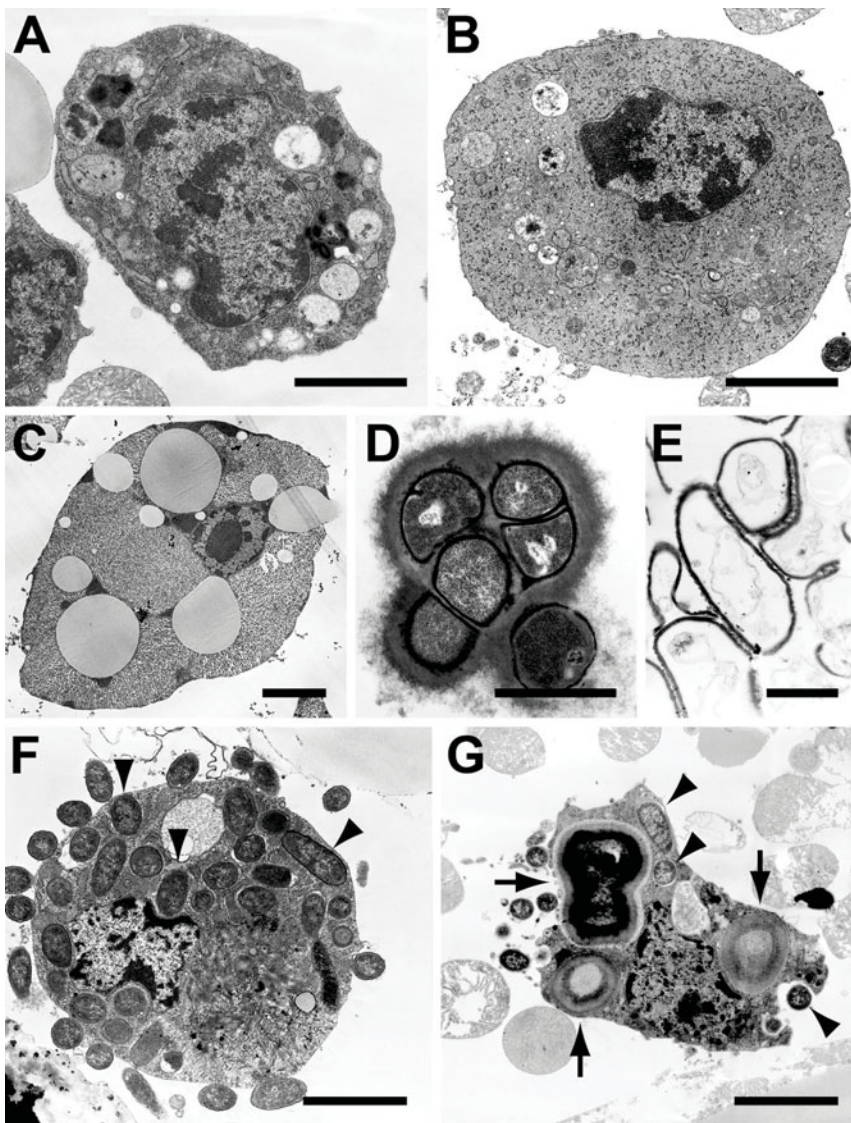


Figure 3. Transmission electron micrographs of immune cells and immune responses in the mosquito hemocoel. A) Circulating granulocyte. B) Circulating oenocytoid. C) Fat body. D) Melanization of *Staphylococcus aureus* in the hemocoel. E) Lysis of *S. aureus* in the hemocoel. F) Phagocytosis of *Escherichia coli* (e.g., arrowheads) by a circulating granulocyte. G) Phagocytosis of unmelanized *E. coli* (e.g., arrowheads) and melanized *Micrococcus luteus* (e.g., arrows) by a circulating granulocyte. Scale bars: A-B, 2 μm ; C, 5 μm ; D-E, 1 μm ; F-G, 3 μm .

and antimicrobial peptides.^{16,17,47,55} RNA interference-based transcriptional knockdown of several hemocyte-produced immunity genes results in increased susceptibility to infection with bacteria and malaria parasites, indicating that hemocytes and the effector molecules they produce are essential for efficient immune responses.^{36,38,46,47,56}

On the basis of morphology, lectin binding properties and enzymatic activity, mosquito hemocytes were initially subdivided into granulocytes, oenocytoids, adipohemocytes and thrombocytoids.⁵⁷ The latter two subtypes were later determined not to be true circulating hemocytes but instead fat body and pericardial cells also collected during the extraction process.⁹ A subsequent study confirmed the initial hemocyte classification system and postulated that a third cell type, progenitor prohemocytes, also circulates with the hemolymph.⁶

The number of circulating hemocytes in an adult mosquito is limited and mitotic activity has not been observed in these cells, suggesting that after eclosion no new hemocytes are produced. Adult mosquitoes contain slightly more than 1,000 circulating hemocytes at the time of emergence and the number of circulating cells drops with age, falling to 800 or less by their sixth day after eclosion.^{6,58} Granulocytes account for approximately 95% of the circulating hemocyte population. They contain membrane-delimited vesicles, exhibit acid phosphatase activity and strongly adhere to artificial substrates following extraction from the mosquito (Fig. 3A).^{8,57} Granulocytes also produce proteins involved in humoral immune pathways, such as nitric oxide synthase, serine proteases and serine protease inhibitors.^{6,12} However, the most striking characteristic of granulocytes is their phagocytic capacity: granulocytes engage in the phagocytosis of bacterial pathogens within 5 minutes after exposure and as infections progress individual hemocytes dramatically grow in size to accommodate the internalization of hundreds of foreign entities (Fig. 3F-G).^{9,12,54,58} Granulocytes have also been observed to phagocytose *Plasmodium* sporozoites, but the rate of sporozoite phagocytosis is low when compared to bacteria and the importance of this immune process in limiting *Plasmodium* infection remains unclear.^{8,59}

The other major hemocyte class is the oenocytoid (Fig. 3B). These cells account for approximately 5% of circulating hemocytes and produce phenoloxidase and phenylalanine hydroxylase,^{9,57,60} which are rate-limiting enzymes in the humoral melanization pathway.^{61,62} These enzymes are present in the cytosol of oenocytoids and their sequences do not contain classical signal peptides,^{60,63} indicating that enzyme release into the hemolymph must occur by either nonclassical secretion mechanisms or cell rupture. Although this process has not been resolved in mosquitoes, phenoloxidase release in lepidopterans and brachyceran dipterans has been shown to occur by cell rupture in an eicosanoid dependent manner.^{64,65}

While mosquitoes mount strong and rapid immune responses following pathogen exposure, the type of immune response can vary depending on the pathogen. For example, the primary immune mechanism against *Escherichia coli* is phagocytosis, but against *Micrococcus luteus* is melanization.^{8,9} Studies on the immune response mounted against a large panel of bacteria determined that the strength of phagocytosis versus melanization responses is not dependent on Gram-type, but the variable response illustrates that mosquitoes discriminate between pathogens.⁵⁴

Additional evidence supporting the importance of hemocytes in immunity comes from a study showing that there is age-associated mortality in *Aedes aegypti* following *E. coli* immune challenge.⁵⁸ This mortality correlates with a decrease in the number of circulating hemocytes and a decrease in the ability to kill *E. coli*, but age has no effect on the transcription of the antimicrobial peptides cecropin, defensin, or gambicin.⁵⁸ These findings suggest that the increase in susceptibility is not due to antimicrobial peptide production but instead to a decrease in the number of circulating hemocytes available to quell the infection and are in agreement with a recent report in *D. melanogaster* showing that targeted ablation of hemocytes renders flies incapable of surviving bacterial infections.⁶⁶

In addition to hemocytes, pericardial cells have been relentlessly implicated in immune surveillance, but no direct evidence has been published. Pericardial cells are large binucleate cells that flank the mosquito heart. In *Anopheles*, these cells have been reported to vary in number from 56 to greater than 300, depending on the observer, and also to uptake ammonia carmine dye.^{67,68} More recent studies have suggested that pericardial cells contain several immune-related molecules,^{69,70} but whether these cells actually produce these proteins or sequester them from the hemolymph is not known. In addition, intense phagocytic activity has been reported near the surface of the mosquito heart, an area densely populated by pericardial cells. However, it remains unclear whether this phagocytic activity is carried out by pericardial cells, sessile hemocytes, or yet unidentified immune cells.⁵⁹ Nevertheless, the location of effector cells in the vicinity of the heart is advantageous for immune surveillance and pathogen destruction, as their position in areas of high hemolymph flow increases their probability of encountering invading pathogens.⁵¹

While hemocytes and possibly pericardial cells mediate both cellular and humoral responses, the fat body's role in immunity is exclusively humoral. The fat body is a multifunctional organ consisting of loosely assembled cells that are rich in glycogen and lipids and line the mosquito integument (Fig. 3C).^{57,71} Among its many functions, fat body synthesizes vitellogenin precursors required for the production of eggs, serves in energy storage and produces numerous hemolymph components.⁷² Specifically, immune activity in the fat body includes infection-induced production of antimicrobial peptides, reactive oxygen species and reactive nitrogen species.^{12,15,73-75}

The Salivary Gland Compartment

Transmission of many viral and protozoan parasites to a vertebrate host requires their injection with the mosquito saliva during blood feeding. Hence, invasion of the salivary gland epithelium and migration into the salivary duct is a requirement for the continuation of the life cycle of these pathogens. The salivary gland epithelium forms a physical barrier that pathogens must cross and *Plasmodium* parasites as well as other pathogens have evolved proteins that drive invasion by first binding to specific mosquito salivary gland surface factors.⁷⁶⁻⁷⁸ To date, little is known about the role of the salivary glands in antimicrobial responses, but several gene expression studies have described the production of immune proteins in the salivary glands of naïve mosquitoes, *Plasmodium*-infected mosquitoes and mosquitoes that have ingested a noninfectious blood meal.⁷⁹⁻⁸⁶ Empirical evidence showing that the salivary glands serve as an active immune organ is largely lacking, with the exception of a single publication showing that a serine protease inhibitor (SRPN6) produced in the salivary epithelium limits gland invasion by *Plasmodium* sporozoites.⁸⁷

MOLECULAR BASIS OF MOSQUITO IMMUNITY

The publication of the *An. gambiae* and *Ae. aegypti* genomes has led to an explosion in the number of studies focusing on mosquito immunity.⁸⁸⁻⁹¹ Many of these studies have employed homology searches to identify putative immune genes and infer their function. These bioinformatic observations, together with gene expression data and transcriptional manipulations using RNA interference or transgenesis, have allowed researchers to conclusively identify genes that are required for pathogen suppression.

Conversely, similar approaches have been used to identify mosquito genes that facilitate pathogen survival.

Pattern Recognition Receptors

Invading pathogens are recognized by the molecular interaction between host-derived pattern recognition receptors (PRRs) and pathogen associated molecular patterns (PAMPs). Bioinformatic analysis of the *An. gambiae* genome has identified approximately 150 putative PRRs.⁹² Most are secreted proteins that contain adhesive domains capable of interacting with PAMPs and cluster as members of large gene families. While experimental evidence has shown that many are involved in immune responses, their actual role as PRRs has not been cemented, as their recognized PAMPs have not been identified.

Thioester containing proteins (TEPs) are hemolymph proteins involved in the killing of bacteria and *Plasmodium* ookinetes.^{41,46,93,94} Members of the TEP gene family share structural similarities with α 2-macroglobulins and vertebrate complement components C3, C4 and C5. By and large, most studies on mosquito TEPs have focused on the hemocyte-produced phagocytosis enhancer TEP1. This protein is secreted into the hemocoel as a single chain molecule that is activated by proteolytic cleavage.⁹³ Cleaved TEP1 is then stabilized by forming a complex with the leucine rich repeat containing proteins LRIM1 and APL1C prior to binding bacteria in the hemocoel or ookinetes in the midgut, triggering their destruction.^{36,38,93} The antiplasmodial activity of APL1 and TEP1 are further supported by studies showing that genetic variation at their respective loci has a profound effect on immune competence against *Plasmodium*.^{4,95} A recent genome-wide mapping of reciprocal crosses of mosquito strains that are susceptible or resistant to *P. berghei* demonstrated that polymorphisms in Tep1 explain a portion of the variability in *P. berghei* killing efficiency observed among laboratory mosquito colonies.⁹⁵ While this study was published years after the discovery of Tep1, APL1 was initially characterized because it is coded within a *Plasmodium*-resistance island that explains naturally occurring resistance to *P. falciparum* in field-caught *An. gambiae*.⁴ Dissection of the APL1 locus revealed that it is composed of three genes, all of which display major structural haplotypes.⁹⁶ Interestingly, APL1C is solely responsible for resistance to *P. berghei*, but distinct haplotypes in the neighboring APL1A gene are associated with various levels of resistance.⁹⁶

C-type lectins are soluble or membrane bound proteins that bind carbohydrates in a calcium-dependent manner. In *An. gambiae*, C-type lectins are both positive and negative regulators of mosquito immune responses. In the midgut, CTL4 and CTLMA2 function as negative regulators of the melanization of *Plasmodium berghei* ookinetes.³⁷ However, in the hemocoel these same C-type lectins are present in the hemolymph as disulfide-linked heterodimers that function in the killing of *E. coli* in a melanization-independent manner.⁹⁷ Transcriptional knockdown of either of these lectins increases bacterial proliferation in the hemocoel and decreases mosquito survival, indicating that they are essential players in the antibacterial response.

Gram-negative binding proteins (GNBPs) were initially identified in *An. gambiae* because they share sequence similarities with GNBPs of other insects and because they are transcriptionally upregulated following infection with bacteria and *Plasmodium* parasites.⁹⁸ Six members of this gene family are expressed in *An. gambiae* and all presumably function as PRRs by binding β -1,3-glucan and lipopolysaccharide on the surface of pathogens. GNBPs are transcribed in multiple tissues (hemocytes, midgut, salivary glands) and while

they are all upregulated following an immune challenge, they vary in their antimicrobial specificities. GNBPA4, for example, participates in the killing of *E. coli*, *Staphylococcus aureus* and *P. berghei*, but not *Plasmodium falciparum*. In contrast, GNBPA2 participates in the killing of *E. coli* and *P. falciparum*, exhibits mild activity against *P. berghei* and is ineffective against *S. aureus*.⁹⁹

The immunoglobulin superfamily consists of 138 genes in *An. gambiae*, 85 of which are upregulated following an immune challenge. Six of these genes were recently extensively characterized: two are involved in the killing of *P. falciparum* ookinetes (IRID4 and 6), two control the growth of opportunistic bacteria (IRID3 and 4) and three are involved in the killing of exogenously introduced bacteria.¹⁰⁰ Another member of this gene family, AgDSCAM, has been shown to opsonize bacteria and to kill the midgut stages of *Plasmodium*.¹⁰¹

Fibrinogen-related proteins (FREPs) represent a PRR family that has experienced massive expansion in mosquitoes.¹⁰² Fifty-nine and 37 FREPs have been identified in *An. gambiae* and *Ae. aegypti*, respectively, compared to 14 in *Drosophila melanogaster*.⁹¹ Functional studies have shown that the majority of mosquito FREPs are upregulated following an immune challenge and that many are essential for the killing of bacteria and the maintenance of immune homeostasis.^{11,103} Several also have antiplasmodial activity and one in particular (FBN9) binds ookinetes as they invade the midgut epithelium.¹¹

Immune Signaling

Microorganism recognition by PRRs can lead to pathogen destruction through constitutive effector mechanisms and/or the activation of intracellular signaling pathways that activate the transcription of effector genes. The major immune signaling pathways in mosquitoes are Toll, Imd and JAK/STAT and the path that leads to activation of these pathways may be amplified or repressed by modulatory proteins such as serine proteases and serine protease inhibitors. Most of the components in these signaling and regulatory pathways are conserved among dipteran insects.⁹¹

In mosquitoes, genes regulated by the Toll pathway are controlled by the NF- κ B transcription factor Rel1. This pathway is induced by fungi, Gram(+) bacteria, viruses and *Plasmodium*. Induction of the Toll pathway by silencing of the negative regulator of Rel1, Cactus, dramatically decreases *P. berghei* and *Plasmodium gallinaceum* infection intensity in the *Anopheles* and *Aedes* midgut, respectively.^{104,105} Co-silencing Rel1 and Cactus renders mosquitoes susceptible to infection, indicating that Cactus-mediated susceptibility is due to repression of Rel1.¹⁰⁴ Co-silencing of Cactus and LRIM1 or Tep1 also renders mosquitoes susceptible to infection, suggesting that these two effector molecules are induced through the Toll pathway. In addition to the antiplasmodial activity of genes induced through Rel1, the Toll pathway is also involved in controlling infection against entomopathogenic fungi and dengue virus.^{106,107}

The Imd pathway is controlled by the NF- κ B transcription factor Rel2. Rel2 exists as short (Rel2S) and full-length (Rel2F) forms, both of which are involved in the immune response against bacteria and *Plasmodium*.^{104,108} Interestingly, while both the Toll and Imd pathways are involved in immunity against *P. berghei*, immunity against *P. falciparum* is controlled primarily through the Imd pathway, as transcriptional knockdown of the Imd negative regulator Caspar and not the Toll negative regulator Cactus renders *An. gambiae* resistant to infection.¹⁰⁹

The least studied immune pathway in mosquitoes is the JAK/STAT pathway. In *An. gambiae*, this pathway is controlled through two STAT transcription factors that are the result of a gene duplication event. In mosquitoes, activation of the STAT pathway requires STAT-B mediated activation of STAT-A, which is regulated by a negative feedback loop controlled by the signaling suppressor protein SOCS. Activation of this pathway leads to the induction of nitric oxide synthase transcription, which is a positive regulator of *Plasmodium* infection.¹¹⁰ More recent studies have also implicated the JAK/STAT pathway in the immune response against dengue virus in *Ae. aegypti*. Here, inactivation of the JAK/STAT pathway by depletion of the receptor Domeless results in increased viral loads.¹¹¹ Conversely, hyperactivation of the pathway by depletion of the negative regulator PIAS results in a more resistant phenotype.

Antimicrobial Peptides

Antimicrobial peptides (AMPs) are secreted low molecular weight proteins that were initially identified for their antimicrobial activity *in vitro*.^{74,75,112,113} Defensins, cecropins and gambicins comprise the three main AMP gene families in mosquitoes. *In vitro* analyses of their antimicrobial spectra showed that cecropins and gambicin are cytotoxic primarily against Gram(−) bacteria and defensins are cytotoxic primarily against Gram(+) bacteria.^{74,75,112} Transcriptional regulation of AMPs occurs through the Toll and Imd pathways,^{114,115} and all are transcriptionally upregulated in fat body following exposure to viruses, bacteria, *Plasmodium* and filarial nematodes.^{16,107,116-118}

While defensins have antimicrobial activity *in vitro*, their function as essential components of the mosquito immune response continues to be debated. In *An. gambiae*, transcriptional knockdown of Defensin decreases mosquito survival following *S. aureus* infection,¹¹⁹ but RNAi-based silencing of *Ae. aegypti* Defensin has no effect on mosquito survival following challenge with three bacterial species.^{120,121} The role of cecropins in the antibacterial response *in vivo* is not known, but ectopic expression of a cecropin transgene results in increased killing of *P. berghei* ookinetes.¹²² Lastly, Gambicin exhibits antiparasitic activity in the midgut, as well as antibacterial activity in the hemocoel.⁴¹

Phenoloxidase-Based Melanization

Melanization in insects is essential for cuticle hardening, egg chorion tanning, wound healing and immunity. In mosquitoes, melanization (also known as melanotic encapsulation) is an immune effector mechanism involved in the killing of *Plasmodium*, filarial nematodes and bacteria and is visually manifested as a darkened proteinaceous capsule that surrounds invading pathogens (Fig. 3D).^{5,8,35} Melanization involves a series of reactions that include the conversion of tyrosine to melanin precursors and the cross-linking of proteins to form a layer of melanin that surrounds and sequesters invading pathogens (Fig. 4).¹⁰ Melanization often results in pathogen death but the killing mechanism remains unclear. It has been hypothesized that death may be caused by either oxidative damage brought on by unstable intermediates created during melanogenesis or by starvation, since the foreign agent becomes isolated from the nutrient-rich hemolymph.^{123,124} Besides functioning as a killing mechanism, melanization is also involved in the clearing of already dead or dying pathogens.^{37,40}

The process of melanization begins with the proteolytic cleavage of a pro-phenoloxidase zymogen into its active form. The exact sequence of events that leads

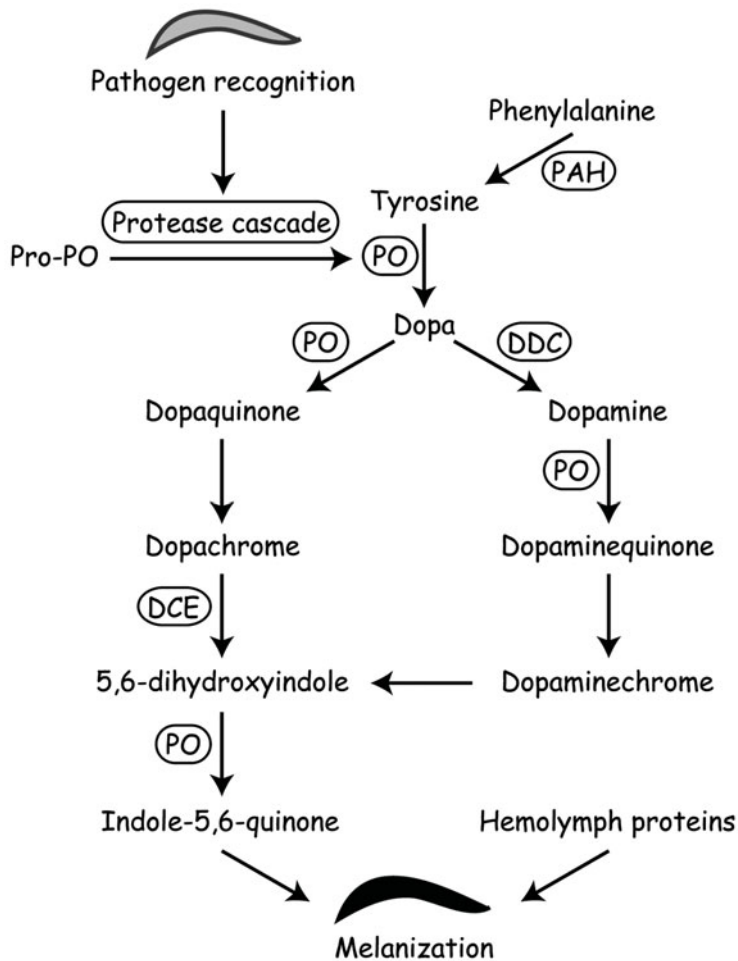


Figure 4. Proposed biochemical pathway leading to the melanization of pathogens. PAH, phenylalanine hydroxylase; PO, phenoloxidase; DDC, dopa decarboxylase; DCE, dopachrome conversion enzyme.

to this cleavage is not well understood, but involves the coordinated action of pattern recognition receptors, serine proteases and serine protease inhibitors. In the hemocoel, β -1,3-glucan recognition protein is required for the melanization of filarial nematodes and bacteria by functioning as a pattern recognition receptor.^{56,125} In the midgut, various proteins serve as promoters and inhibitors of melanization, with their exact role varying greatly between mosquito strains. For example, in *An. gambiae*, the *Plasmodium*-resistant L3-5 strain naturally melanizes *P. berghei* ookinetes but the G3 strain is susceptible to infection. Interestingly, mosquito C-type lectins function as repressors of melanization in the susceptible strain, as silencing CTL4 and CTLMA2 results in the LRIM1-dependent induction of ookinete melanization.³⁷ CLIP domain serine proteases also modulate the melanization response against *P. berghei* ookinetes and share structural similarities with pro-phenoloxidase activating enzymes from other insects.¹²⁶ However, once again, the

genetic background of the mosquito has a profound effect on the function of CLIPs.^{40,126} For example, CLIPA7 inhibits melanization in the susceptible strain but has no effect on the resistant strain. Conversely, CLIPB3 has no effect on the susceptible strain but promotes melanization in the resistant strain. Similar observations on the role of CLIPs as inhibitors and promoters of melanization have been made in the mosquito hemocoel after intrathoracic injection of sephadex beads.¹²⁷

In addition to CLIPs, mosquito serine protease inhibitors (SRPNs) modulate melanization responses.¹²⁸ Both SRPN6 and SRPN2 inhibit the melanization of *P. berghei* ookinetes in the resistant strain and SRPN2 also inhibits the spontaneous formation of melanin-based pseudotumors in the hemocoel.^{129,130} These pseudotumors drastically reduce mosquito fitness, underscoring the importance of tightly regulating this immune process. Moreover, the factors that trigger melanization in the midgut are at least partially dependent on the host-parasite combination: SRPN2, CTLs and LRIM1, molecules that regulate development of *P. berghei* in the midgut, have no detectable effect on the development of the human malaria parasite *P. falciparum*.^{131,132} These findings are not surprising given that *P. berghei* infection triggers the transcriptional regulation of over twice as many genes as *P. falciparum*.⁴¹

Once phenoloxidase becomes activated, the formation of melanin is initiated by the phenoloxidase-mediated hydroxylation of tyrosine to form dopa (Fig. 4).¹³³ Dopa is then oxidized by phenoloxidase to form dopaquinone, which is then converted to dopachrome. Dopachrome conversion enzyme converts dopachrome to 5,6-dihydroxyindole, which is oxidized into indole-5,6-quinone by phenoloxidase and then cross-linked with hemolymph proteins to form melanotic capsules. In an alternative pathway, dopa formed by the hydroxylation of tyrosine is decarboxylated by dopa decarboxylase to form dopamine, which is then converted into melanin by phenoloxidase and other enzymes. Empirical testing has shown that pathogen-induced melanin formation is accomplished via both of these pathways, as transcriptional knockdown of phenoloxidase, dopachrome conversion enzyme and dopa decarboxylase all lead to impaired melanization responses.^{62,134-136} Throughout these reactions, tyrosine remains the rate-limiting substrate and endogenous production of tyrosine is accomplished by the hydroxylation of phenylalanine by phenylalanine hydroxylase. This latter reaction is essential for the melanization of filarial nematodes but not sephadex beads,^{61,136} with the observed difference possibly related to the larger surface area that needs to be melanized following filarial worm infection. Finally, while melanization events are extracellular, many melanin-producing enzymes (e.g., phenoloxidase, dopachrome conversion enzyme, dopa decarboxylase and phenylalanine hydroxylase) are produced by circulating hemocytes,^{9,57,60,134,135} and melanized pathogens are often subsequently phagocytosed by granulocytes (Fig. 3G).^{8,9,54}

Nitric Oxide and Other Reactive Species

Nitric oxide is a multifunctional free radical created during the oxidation of L-Arginine to L-Citrulline by the enzyme nitric oxide synthase.¹³⁷ In *Anopheles*, nitric oxide synthase is a single copy gene with 18-22 distinct transcripts. Three of these transcripts are induced by *Plasmodium* infection,^{138,139} and at least one is induced by bacterial infection.¹² In the midgut of the mosquito, *Plasmodium* glycosylphosphatidylinositols and *Plasmodium*-derived hemozoin acquired with an infectious blood meal induce the transcription of nitric oxide synthase through the STAT pathway,^{43,110,140} and the resultant

nitric oxide kills *Plasmodium* ookinetes via lysis.^{14,141,142} In the hemocoel, nitric oxide synthase is transcriptionally upregulated following bacterial infection and the production of nitric oxide is required for bacterial killing and mosquito survival during systemic infections with *E. coli*.¹²

Reactive oxygen species (ROS) kill *Plasmodium* ookinetes in the midgut and bacteria in the hemocoel.^{15,143,144} While the exact mechanism of action against bacteria remains unknown, ROS kill *Plasmodium* through both lytic and melanization pathways.^{15,143,144} The *P. berghei*-resistant L3-5 strain of *An. gambiae* lives in a constant state of oxidative stress that promotes melanization of ookinetes as they traverse the midgut epithelium.¹⁴³ Conversely, the susceptible G3 strain kills ookinetes via a lytic mechanism that is dependent on infection-induced oxidative stress that is maintained by the repression of catalase, an enzyme that breaks down hydrogen peroxide into oxygen and water.¹⁵ Furthermore, reactive oxygen and reactive nitrogen species are intimately linked in mosquito immunity: peroxidases in the mosquito midgut use nitrite and hydrogen peroxide to synthesize highly reactive nitrogen dioxide and hydrogen peroxide triggers the transcriptional induction of nitric oxide synthase.^{145,146}

Phagocytosis

Phagocytosis is an evolutionarily conserved immune process used for the killing and sequestration of small microorganisms. In this process, a particle is recognized, bound by proteins in the plasma membrane and internalized into a membrane-delimited phagosome. The phagosome then fuses with a lysosome and hydrolytic enzymes digest the particle. In mosquitoes, the granulocyte subpopulation of hemocytes uses this immune process to sequester and kill bacteria as early as 5 minutes after exposure.^{8,9} Approximately 95% of circulating hemocytes are phagocytic and it has been estimated that individual hemocytes are capable of phagocytosing over 1,000 bacteria within 24 hours of infection.⁵⁸

Several studies have investigated the molecular basis of phagocytosis by visualizing the uptake of fluorescently labeled dead bacteria by mosquito immortal cell lines or by low magnification fluorescence microscopy of whole mosquitoes.^{93,94,147} Identified regulators of phagocytosis include pattern recognition receptors, transmembrane receptors and intracellular signaling proteins.

Thioester containing proteins TEP1, TEP3 and TEP4 are involved in the phagocytosis of Gram(+) and Gram(-) bacteria in the mosquito hemocoel.^{93,94} The modes of action of TEP3 and TEP4 are not known, but proteolytically activated TEP1 opsonises bacteria by thioester-mediated binding, which in turn initiates phagocytosis. In addition to the above TEPs, the leucine rich repeat containing protein LRIM1 is also required for phagocytosis.⁹⁴ The exact mechanism by which LRIM1 functions has not been resolved, but TEP1-based antimicrobial activity in the midgut is dependent on the complexing of TEP1 with LRIM1 in the hemocoel.³⁶

AgDSCAM is a hypervariable immunoglobulin that is encoded by 101 exons, which can be transpliced into over 31,000 variants.¹⁰¹ AgDSCAM binds bacteria in vitro and triggers their phagocytosis by an immortal hemocyte-like cell line. In vivo transcriptional knockdown of AgDSCAM increases bacterial proliferation in the hemocoel and decreases mosquito survival, suggesting that AgDSCAM mediated phagocytosis or humoral killing is required for effective antibacterial responses in the hemocoel.

Several transmembrane receptors have also been implicated in phagocytosis and may function by either directly recognizing pathogens or by recognizing pathogens that

have been opsonized by hemolymph proteins. Among these are a β integrin (BINT2), a peptidoglycan recognition protein (PGRPLC) and a low-density lipoprotein receptor-related protein (LRP1).^{94,147}

Lastly, several intracellular proteins trigger the internalization of bacteria. Transcriptional knockdown of *An. gambiae* CED2, CED5, or CED6 reduces phagocytosis efficacy by up to 80%.⁹⁴ Epistatic analyses then showed that TEP1, TEP3, LRIM1 and LRP1-mediated phagocytosis occurs through the CED6 pathway and TEP4 and BINT2-mediated phagocytosis occurs through the CED2/CED5 pathway. The involvement of mosquito CEDs in the phagocytosis of foreign bodies through two genetically independent pathways is significant but not unexpected. Cell death abnormal genes were initially discovered in *Caenorhabditis elegans* during screens aimed at identifying genetic factors required for the phagocytosis of apoptotic bodies in the developing worm and these factors function through two independent but partially redundant pathways: the CED-1/CED-6/CED-7 pathway and the CED-2/CED-5/CED-10/CED-12 pathway.¹⁴⁸⁻¹⁵⁰

CONCLUSION

The field of mosquito immunity has experienced unprecedented growth in the past decade. The publication of the *An. gambiae* and *Ae. aegypti* genomic sequences has led to the bioinformatic identification of numerous putative immunity genes.⁸⁸⁻⁹¹ Technical advances such as RNA interference, paratransgenesis and transgenesis have then allowed researchers to empirically test their function in effecting or regulating immune responses against diverse groups of pathogens.^{119,151,152} As a result of these studies, it has become apparent that the mosquito immune system shares numerous similarities with vertebrate immune systems. Immune responses in both vertebrates and invertebrates are initiated by microbe recognition events that trigger signaling pathways and effector mechanisms. Moreover, similar to vertebrates, mosquitoes recognize pathogens using complement-like cascades, possess phagocytic cells that circulate with the blood and transcribe effector molecules through Toll and JAK/STAT pathways. Further studies on mosquito immunity will continue to shed light on the evolution of these complex and essential responses.

In addition to the evolutionary conservation of immune components, obtaining a better understanding of the mosquito immune system may translate into novel public health interventions. Mosquitoes are cosmopolitan pests and disease vectors.¹⁵³ Because of their global importance, these organisms are subjects of constant study in efforts to uncover mechanisms that reduce their population densities as well as their ability to transmit disease. One aspect of mosquito biology that continues to receive considerable attention is their ability to fight infectious agents, as it has been hypothesized that understanding how mosquitoes kill microbial pathogens may allow us to exploit weaknesses that increase the population-reduction effectiveness of biological control strategies, or to strengthen immune responses such that otherwise susceptible mosquitoes are rendered resistant to infection, halting disease transmission cycles.^{19,20,154,155} Much must be done before the feasibility of these approaches can be further considered, including the continued expansion of work in immunity to wild mosquito populations and to a broader range of mosquito species in order to ensure that our current understanding of mosquito immunity is representative of natural host-pathogen interactions.

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CRUSTACEAN IMMUNITY

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Abstract: This chapter provides a review of recent progress in the elucidation of innate immune mechanisms in crustaceans. Mainly due to the importance of crustacean aquaculture interest in this field is large and the subject for extensive research efforts. Here, we provide detailed data on the molecular characterisation of lectins, antiviral reactions, hemocyte formation and differentiation and on the regulation of innate immune pathways.

INTRODUCTION

Crustaceans are relatively well investigated with respect to immune reactions when compared to most other invertebrates except fruit flies. This is of course due to their large size and to the intensive fishing and aquaculture of shrimps and some other decapod crustaceans. Although they are (at least until now) less amenable for genetical experiments they are relatively easy to keep in aquaria and to bleed and, therefore, considerable amounts of plasma and hemocytes can be collected for work at molecular or cellular level. Most researchers in the field are using shrimps but many pioneering studies have been carried out on other crustaceans, in particular freshwater crayfish. In this chapter we will cover recent advances in crustacean immunity with emphasis on pattern recognition and lectins, hemocytes and hematopoiesis, prophenoloxidase activating system and on antiviral mechanisms. For earlier work in general we refer to references 1 and 2, and for reviews on crustacean antimicrobial peptides to references 3 and 4 and on antiviral immunity to reference 5.

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PATTERN RECOGNITION

Pathogens that manage to break through the outer protective parts of the animal such as the resistant chitin-containing cuticle (or through the midgut which lacks chitin) will encounter an array of hemocyte- and plasma-derived immune factors. The activities of these factors are triggered by molecular signatures typically present on or released by different microorganisms. Lipopolysaccharides, β -1,3-glucans and to some extent peptidoglycans, i.e., polysaccharides from the microbial cell-wall, are known to initiate immune reactions in crustaceans. Double-stranded RNA derived from some viruses is another inducer of such reactions. It is possible that enzymes (i.e., proteinases) produced by microorganisms, or the damage on host tissue caused by such enzymes are efficient triggers of the defence as has been shown in *Drosophila*,⁶ but this is less known in crustaceans.

A number of pattern recognition proteins have been isolated from crustaceans and characterised in detail. Most known crustacean pattern recognition proteins were originally purified and cloned in freshwater crayfish but have subsequently been found in shrimps and other crustaceans. These include β -glucan-binding protein^{7,8} (BGBP also abbreviated β GBP), lipopolysaccharide- and glucan-binding protein⁹ (LGBP) some masquerade-like proteins/serine proteinase homologues^{10,11} (SPHs) and a large number of lectins (see below). The LGBPs and the BGBPs will bind β -1,3-glucans and after this binding they will trigger immune reactions such as proPO-activation. LGBP is probably the main vehicle in crustaceans for the recognition of these glucans and thus for mediating defence reactions directed against fungi and oomycetes.⁹ Also BGBP, which is not structurally related to LGBP, is capable of binding β -1,3-glucans and mediating immune reactions.^{7,8} The affinity of binding to the glucans is lower for crayfish BGBP than for LGBP and the latter is as mentioned therefore likely to be more important for mediating glucan-triggered immune reactions. However, the plasma concentration of BGBP is high and it is therefore possible that this protein is important in removing excess glucans, if present. LGBP is also rendered active by gram-negative bacteria since it is capable of binding LPS and thereafter mediate the activation of the proPO-system. Insect homologues of LGBP (variously called β -1,3-glucan binding proteins or gram-negative bacteria binding proteins, i.e., GNBPs) have been shown to trigger the Toll pathway, melanisation and other immune reactions in the presence of bacteria or fungi.¹² This family of proteins appears to have gone through a large expansion in the crustacean *Daphnia pulex* since 11 genes coding for LGBP-like proteins were detected in its genome.¹³

No obvious peptidoglycan recognition protein (PGRP) candidates were found in the recently released *Daphnia* genome, nor has yet any PGRP gene been cloned from any crustacean. PGRPs are present in many vertebrate and invertebrate species so their apparent absence in crustaceans is a surprise. Still, peptidoglycans have been reported to stimulate immune responses from crustaceans¹⁴ although the molecular mechanisms behind this need to be clarified.

SPHs are known as activators/regulators of proPO in some insects¹⁵ but in crustaceans, they have mainly been implicated in pattern recognition and as opsonins so far. The first crustacean SPH characterised, the masquerade-like protein¹⁰ is binding to gram-negative bacteria.¹¹ This binding is triggering a proteolytic processing of the protein that produces four different subunits.¹¹ Crayfish masquerade-like protein was demonstrated experimentally to be an efficient opsonin and important in the clearance of *E. coli*. A black tiger SPH that could bind lipopolysaccharides and intact *V. harveyi* and acting as opsonin has been described recently.¹⁶

LECTINS

Lectins are proteins or glycoproteins normally without catalytic activity that can recognise and noncovalently bind to specific sugar moieties and thereby agglutinate cells by binding to cell surface glycoproteins and glycoconjugates.¹⁷ Lectins, therefore, are considered important pattern recognition proteins in innate immunity and play significant roles in nonself-recognition and clearance of invading microorganisms, either as cell surface receptors or as soluble proteins existing in circulating fluids.^{18,19} C-type lectins are the most diverse and well studied among the lectin families. The term C-type lectin was originally used to distinguish a group of Ca²⁺-dependent (C-type) carbohydrate-binding proteins from the other types of lectins.²⁰ The structures of C-type lectins were defined and found to contain a conserved single module of approximately 150 amino acid residues (carbohydrate recognition domain, CRD).²⁰⁻²² This domain contains a characteristic double-loop stabilised by two highly conserved disulphide bridges and four Ca²⁺-binding sites where the Ca²⁺ binding site 2 is involved in carbohydrate binding.²³ The CRD's usually have a key motif, either QPD (Gln-Pro-Asp) or EPN (Glu-Pro-Asn), which has been predicted to be ligand-binding specific for galactose or mannose, respectively.²³ Recently many C-type lectins containing nonstandard CRDs, which do not bind Ca²⁺ have been identified. These are considered to interact with noncarbohydrate ligands²⁴ and for these CRDs the term C-type lectin-like domain (CTLD) was introduced.²⁵

Although C-type lectins have been well studied in vertebrates for many years, they have not been well characterised in invertebrates. Recently, genes containing CTLDs have been found to be abundant in the *Daphnia pulex* (6 genes) genome, *Drosophila melanogaster* genome (34 genes) and in *Caenorhabditis elegans* genome (278 genes), respectively.^{13,26} This suggests that there is a high potential for generating many C-type lectins, perhaps with different ligand specificities. C-type lectins are the largest group of immune-function ESTs found in the hepatopancreas of the shrimps *Litopenaeus vannamei* and *L. setiferus*.²⁷ It has become clear that vertebrate C-type lectins have a broad range of biological functions including cell adhesion, endocytosis, pathogen neutralisation, glycoprotein clearance, phagocytosis.^{17,28,29} In invertebrates, lectins have been reported to contribute in innate immune responses, including prophenoloxidase activation,^{30,31} enhancement of encapsulation,^{19,32,33} nodule formation of hemocytes,³⁴ opsonin formation,³⁵ antibacterial activity,³⁶ antifungal activity³⁷ and maybe contribute to injury healing.³⁸

A large number of natural lectins have been purified and characterised by biochemical methods from hemolymph of crustaceans (for reviews see refs. 39,40). Compared to vertebrate lectins, the molecular features and functions of lectins in crustaceans are just at the beginning of becoming understood. Here focus will be placed on those lectins which have been sequenced and whose functional properties have been determined using e.g., recombinant proteins (listed in Table 1).

Structure of the Shrimp C-Type Lectins

All lectins listed in the Table 1 contain a CTLD in the putative protein indicating that these lectins fall into the C-type lectin family. The four or six cysteine residues important in the formation of the CRD disulphide bonds are conserved. PmAV, PmLec and Fc-Lec4 contain a single CRD with a QPD motif that has a predicted ligand-binding specificity for galactose, while Fc-hsL, LvLec and LvCTL1 contain a single CRD with an EPN motif with predicted ligand-binding affinity toward mannose. C-type lectin-1

Table 1. Shrimp lectins characterised by molecular methods

		Amino Acids	CRD	Sugar-Binding Motif	Biological Activity	GenBank Accession Number	Challenged	Tissue Specific	Reference
C-type lectin-1	<i>P. stylirostris</i>	168	1	EPK	-	-	WSSV	Hepatopancreas	141
PmAV	<i>P. monodon</i>	170	1	QPD	Antiviral activity	AY302750	WSSV	-	50
PmLec	<i>P. monodon</i>	182	1	QPD	Binding to LPS Agglutinating, opsonic effect	DQ078266	-	-	44
LvLT	<i>L. vannamei</i>	345	2	QPD and EPD		DQ871245	WSSV	Hepatopancreas	142
Fclectin	<i>F. chinensis</i>	287	2	2 QPD		AY871270	Bacteria/ WSSV	Hemocyte	43
PmLT	<i>P. monodon</i>	333	2	QPD and EPN	WSSV binding, Enhance	DQ871244	Bacteria/ WSSV	Hepatopancreas	46
Fc-hsL	<i>F. chinensis</i>	159	1	EPN	encapsulation Agglutinating, binding, antimicrobial activity	DQ167572	Bacteria/ WSSV	Hepatopancreas	47
Fc-Lec2	<i>F. chinensis</i>	333	2	QPD and EPN	Agglutinating activity	EU834289	Bacteria/ WSSV	Hepatopancreas	41
FcLec3	<i>F. chinensis</i>	158	1	EPS	Agglutinating		Bacteria/ WSSV	Hepatopancreas	49
FcLec4	<i>F. chinensis</i>	237	1	QPD	Agglutinating, binding activity to bacteria, clearance of bacteria	EU834293	Bacteria	Hepatopancreas, gills and stomach	42

continued on next page

Table 1. Continued

	Amino Acids	CRD	Sugar-Binding Motif	Biological Activity	GenBank Accession Number	Challenged	Tissue Specific	Reference
LvLec	157	1	EPN	Agglutinating, Mannose binding	EF583939	-	Brain, hemocytes and hepatopancreas	45
LvCTLI	156	1	EPN	Hemagglutinating, sugar binding, Binding to WSSV envelope proteins	DQ858900	WSSV	Hepatopancreas	48
PtLP	164	1	-	-	ACC86854	-	-	143

and Fc-Lec3 also contain a single CRD, but with EPK and EPS, respectively instead of the usual EPN motif. PmLT, Fc-Lec2 and LvLT consist of two CRDs, the N-terminal CRD1 contains a QPD motif and the C-terminal contains an EPN motif, but LvLT has here an EPD instead. Those mutated motifs and their binding specificities need further studies. Fclectin contains two CRDs, each with a QPD motif. The number of CRDs identified in each species is variable and it seems likely that lectins with multiple CRDs have stronger affinity for binding to their ligands.²³

According to the phylogenetic tree made by Zhang et al⁴¹ lectins from shrimp, scallop and the arctic lamprey *Lethenteron japonicum* form one large cluster, those from mammals and fishes form the second cluster and those from insect belongs to the third cluster. Fc-Lec2, PmLT and LvLT are belonging to same subcluster, while Fc-hsL and LvLec are grouped in another subcluster. This could indicate that shrimp lectins may have a closer relationship with mammalian lectins than insect lectins. Thereafter was FcLec4, however, found to be closely related to insect lectins.⁴²

Tissue-Specific Expression of Shrimp Lectins

Hepatopancreas and hemocytes of crustacean are regarded as the most important tissues involved in crustacean immunity.^{1,27} Most shrimp lectins characterised to date have been isolated from hepatopancreas or hemocyte cDNA libraries. The Fclectin transcript was detected exclusively in hemocytes,⁴³ whereas the transcripts of LvLT, PmLT, Fc-hsL, Fc-lec2, Fclec3 and LvCTL1 were found specifically expressed in hepatopancreas. The detailed localisation of PmLT and Fc-Lec2 was demonstrated by immunohistochemistry to be in the F (fibrillar) cells of the hepatopancreas. In contrast, the FcLec4 transcripts were distributed in diverse tissues, mainly in the hepatopancreas, gill, stomach and a lower level could also be detected in intestine.⁴² LvLec is unusual in having its highest expression in the brain; an interesting finding that needs further functional studies.

Functional Studies

By the use of recombinant proteins functional studies of several crustacean lectins have been carried out. The lipopolysaccharide-binding lectin PmLec can function as an opsonin that enhances hemocytic phagocytosis.⁴⁴ Recombinant protein of Fc-hsL has no hemagglutinating activity, but a Ca²⁺ dependent agglutinating activity against several Gram-positive and Gram-negative bacteria. Similarly, Fc-Lec2 and its two individual CRDs did not have hemagglutination activity, but had agglutinating activity and binding activity to some bacteria in a Ca²⁺-dependent and Ca²⁺-independent manner, respectively.⁴¹ Their studies also suggest that two CRDs have synergistic effect. Recombinant LvLec has agglutinating activity to *E. coli* JM 109 depending on Ca²⁺ and the agglutination could be inhibited by mannose and EDTA.⁴⁵ Another role in immunity encapsulation, was demonstrated with PmLT by using agarose beads coated with the lectin.⁴⁶ Unlike other shrimp lectins, Fc-hsL has antimicrobial activity against several bacteria and fungi.⁴⁷

A possible effect of LvCTL1 on virus defence is indicated by the binding of the lectin to WSSV virions and the interaction in a pull-down assay with several envelope proteins of WSSV including VP95, 28, 26, 24, 19 and 14.⁴⁸ Also FcLec3 was shown to interact with a major envelope protein of WSSV, VP28.⁴⁹ PmAV, a C-type lectin

identified from WSSV-resistant shrimp, has a strong antiviral activity in inhibiting virus-induced cytopathic effect in fish cell cultures. Surprisingly, neither recombinant nor native PmAV has agglutination activity⁵⁰ so the mechanism for its antiviral activity needs to be determined.

Expression Profile of Lectins after Challenge with Bacteria or WSSV

Almost all C-type lectins are synthesised in the hepatopancreas or hemocytes, tissues that are important tissues in immunity.^{1,27} Thus, expression of mRNA and protein might be affected by bacterial and viral infection. *PmLec* was isolated from hepatopancreas libraries⁴⁴ and it was found to be highly expressed in the midgut of *P. monodon* challenged by an immersion with *V. harveyi*.⁵¹ *Fc-hsL*, *Fc-Lec2* and *FcLec3* were constitutively expressed in the hepatopancreas of normal shrimp and were highly up-regulated following challenge with either bacteria or WSSV.^{42,45,47} The expression pattern of *PmLT* mRNA, specifically expressed in hepatopancreas, was decreased initially and then gradually increased after treatment with WSSV extract either in vivo or in vitro using a hepatopancreas tissue fragment.⁴⁶ Zhao et al⁴⁸ also reported that *LvCTL1* specifically expressed in the hepatopancreas, was induced in the shrimp hemolymph after WSSV infection. Moreover, the binding of rLvCTL1 to WSSV could protect shrimp from viral infection and prolonged the survival of shrimp against WSSV infection.

FcLec4, which is distinct from other shrimp C-type lectins, is expressed in hepatopancreas, gills and stomach and intestine. A significant up-regulation of FcLec4 transcripts in gills and stomach and higher level of protein in gill stomach and hemolymph was observed after challenge with *V. anguillarum*.⁴⁹ Fclectin expression in hemocytes increased on exposure with inactive mixed bacteria of *V. anguillarum* and *S. aureus* as well as with WSSV. Similar results were observed with in vitro experiments, which showed that Fclectin expression was gradually increased in cultured hemocytes stimulated by LPS.⁴³

Apart from shrimp, a C-type lectin (PtLP) was also isolated from the swimming crab *Portunus trituberculatus*, but with unknown function. PtLP is phylogenetically related to PmAV, but no perfect QPD motif was found and its mRNA levels were very high in hepatopancreas but lower in gills, hemocytes and ovary of unchallenged animals.

In addition to C-type lectins two isoforms of Tachylecin5-like genes (*PmTL5*) have been found in *P. monodon*. It is interesting that the first PmTL5 isoform was mainly expressed in the hindgut and was induced during immersion with *V. harveyi*, while the second was expressed at a very high level in all parts of shrimp intestine and hemocytes.⁵¹

There is a rising list of putative C-type lectin genes which have been successfully cloned and characterised in different shrimps. Many lectins are up-regulated during infection and since there are some data on lectins showing that they are promoting bacterial agglutination, phagocytosis, encapsulation and other immune reactions, a role for these proteins in defence seems likely. In future, more efforts need to be concentrated on biochemical characters, regulatory mechanisms, evolution and precise function of shrimp C-type lectins as well as searching for other types of lectins in addition to C-type lectins.

HEMOCYTES AND HEMATOPOIESIS

Phagocytosis is likely to be of great importance in crustacean immunity and there are numerous studies demonstrating efficient uptake of bacteria and other particles by the circulating hemocytes or fixed phagocytic cells in these animals. Compared to vertebrates little is known about which proteins are regulating and accomplishing this uptake. A few crustacean proteins with opsonic properties have been characterised, though. One example is the masquerade-like protein mentioned earlier. Another very important protein is peroxinectin originally purified⁵² and cloned from freshwater crayfish⁵³ and since then found in shrimps, other crustaceans and many other organisms. Hemocytes, in the presence of e.g., β -1,3-glucans or other triggers of immune reactions, will release peroxinectin whereupon the protein, by limited proteolysis, will gain a strong cell adhesion activity. Peroxinectin acts as an opsonin during phagocytosis and in promoting cellular encapsulation of foreign objects. The *D. pulex* genome contains six putative scavenger receptors¹³ hinting at the possible existence of a cellular uptake mechanism via such receptors in crustaceans.

In crustaceans, the circulating hemocytes are crucial in protecting the animal against invading microorganisms by participating in recognition, phagocytosis, melanisation and cytotoxicity.¹ In most crustaceans three morphologically different classes of hemocytes, hyaline cells (HC), semigranular cells (SGCs) and granular cells (GCs) are observed within the hemolymph and all of them are important in immobilising or destroying invasive pathogens.^{54,55} The hemocyte separating technique developed by Söderhäll and Smith⁵⁶ made it possible to study the function of individual hemocyte types. In freshwater crayfish and shore crab HCs were then shown to be phagocytic, while SGCs act in early detection of pathogens.⁵⁵ The GCs contain within their granules several immune factors such as the proPO-activating system, the cell adhesion protein peroxinectin and crustin antimicrobial peptides.⁵⁷ Exocytosis is induced in both SGCs and GCs as a response to microbial polysaccharides, resulting in the release of these immune proteins. Similar hemocyte types are identified in other arthropods although with slight variations.⁵⁸

The continuous formation of new hemocytes (hematopoiesis) is essential for survival of the animals and this process is tightly regulated by factors released from circulating hemocytes. Arthropod hemocyte development has mainly been studied in the fruit fly, *D. melanogaster*⁵⁹ and in the freshwater crayfish *Pacifastacus leniusculus*.⁶⁰⁻⁶⁵ In *D. melanogaster* mature hemocytes are formed in the early embryonic head mesoderm and at larval stage in a specialised organ called the lymph gland, while no new hemocytes are produced in the adult flies. In contrast the crayfish hematopoietic development is an ongoing process throughout the animals whole life.

As early as 1891 the observation of a structure in the Atlantic ditch shrimp, *Palaemonetes varians*, named “the dorsal blood sinus” was reported by Weldon.⁶⁶ Later Allen⁶⁷ suggests in a more detailed study of this organ, “the dorsal sac”, to be blood cell producing. Since then, several studies of crustacean hematopoietic tissues (HPTs) have been presented and are reviewed by Johnson.⁶⁸ In decapod crustaceans the HPT is a defined organ made up of lobules enveloped by a thin casing of connective tissue, that in noncaridean pleocyemata covers the dorsal and lateral walls of the foregut (Fig. 1A). The HPT consists of lobules of highly active proliferating cells (shown in Figure 1B-C by 5'-bromo-2'-deoxyuridine (BrdU) incorporation and mitotic figures in hematoxylin stained tissue). In dendrobranchiata, such as the penaeid shrimps, hemocytes are produced in paired nodules on the dorsolateral surface of the foregut and in some species supplementary HPTs are localised at the base of

the maxillipeds, or adjacent to the antennal artery.⁶⁹⁻⁷¹ The lymphoid organ (LO) of penaeid shrimps consists of lobes of folded tubules located ventro-anterior to the hepatopancreas. This organ has been mistaken for a hematopoietic tissue, but according to detailed studies by Van de Braak et al⁷² it is clear that the LO mainly has a role in bacterial clearance and is homologous in function to the phagocytic organ described by Cuenot⁷³ and fixed phagocytes associated with the hepatic artery in decapod crustaceans.⁶⁸

The organisation of crustacean HPTs have been described in detail by electron microscopy studies in the crab *Carcinus maenas*,^{74,75} the shrimp *Sicyonia ingentis*,^{71,76} the lobster *Homarus americanus*⁷⁷ and in *P. leniusculus*.⁶⁰ These studies are in agreement with each other and we will in the further text adopt the nomenclature from Chaga et al⁶⁰ in classifying the HPT cells into five distinct categories based on morphological criteria. Type 1 cells have a large nuclei surrounded by small amount of cytoplasm which usually is characteristic of a stem cell while Type 2 also has large nuclei but larger cytoplasm containing cytoplasmic granules. Type 1 and 2 are the main proliferating cells in the HPT, whereas the other cell types in HPT can be categorised into precursors of granular hemocytes as Type 3 to 4, or as precursor of semigranular hemocytes as Type 5 (Fig. 1D).⁶⁴

The formation and development of mature hemocytes involve proliferation, commitment and differentiation from undifferentiated HPT cells. Several transcription factors have been characterised as lineage specific markers in *Drosophila* and are conserved across taxonomic groups from flies to mammals.⁷⁸ Also in *P. leniusculus* the importance of a GATA transcription factor as well as a Runx protein homologue, during hematopoiesis has been revealed.⁶¹ Apart from systematic detailed studies of hematopoietic transcription factors and signalling pathways associated with *Drosophila* hematopoiesis, little is known about the events regulating this process during development or an infection in insects.

In crustaceans, generally, hemocytes do not divide in the circulatory system and thus, new hemocytes need to be continuously and proportionally produced. Already experiments performed in the late 1800s revealed an increase in mitotic index in the HPT following experimental bleeding,⁶⁸ and since then several studies have confirmed that cell proliferation in the HPT can be influenced by moulting,⁶⁸ and different stress factors such as for example LPS injection,⁷² and Mn-exposure.⁷⁹ In addition, the number of blood cells can be experimentally decreased by injection of microbial polysaccharides and then rapid recovery is stimulated, mainly due to production and release of new cells⁶¹ form the HPT.

That new hemocytes are synthesised and partly differentiated in the HPT, but the final differentiation from stem cells not expressing proPO, into functional hemocytes expressing proPO is not completed until the hemocytes are released into the circulation are supported in several reports.^{61,68} A method was developed to isolate the HPT cells from *P. leniusculus* in order to study their proliferation and differentiation in vitro.⁶² The key to the successful stem cell culture was the isolation and characterisation of a new group of cytokines named astakines present in crayfish plasma. Two different astakines have been detected in crustaceans (Table 2) and astakine 1 is a small protein of 9 kDa containing a so-called prokineticin-domain present in several vertebrates. In *Penaeus monodon* astakine 2 contains a 13 amino acid insert as compared to astakine 1,^{62,80} and this protein is also found in *P. leniusculus*. The importance of astakine 1 and astakine 2 in HPT cell proliferation, differentiation and release from the HPT has been shown by injection of recombinant protein as well as by in vivo and in vitro gene silencing by RNAi.⁶² These experiments have revealed that astakine 1 supports differentiation of HPT cells into the SG cell lineage, since the addition of astakine 1 induces expression

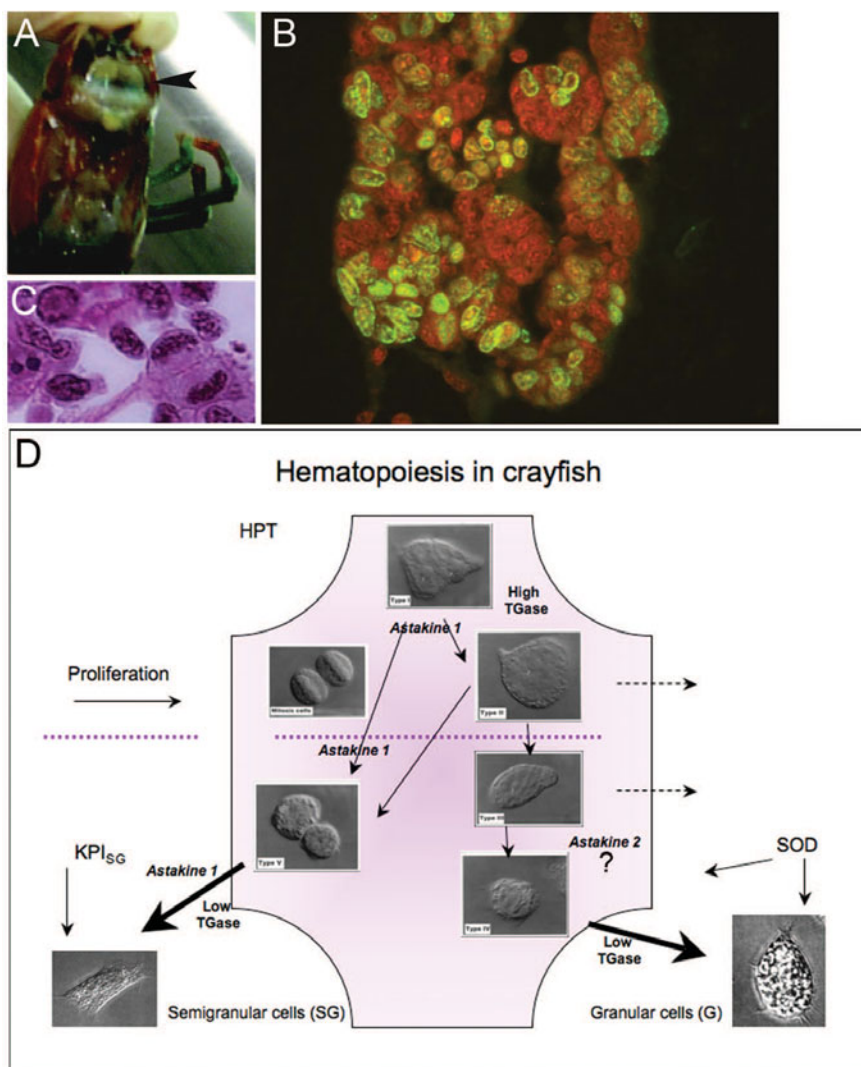


Figure 1. A) The hematopoietic tissue (HPT) of crayfish covers the dorsal and lateral walls of the foregut. B) Proliferating cells in HPT labelled by 5'-bromo-2'-deoxyuridine (BrdU) incorporation. C) Mitosis in HPT cells stained by hematoxylin. D) A hypothetical model for hemocyte development and release from the HPT in *P. leniusculus*. Based on data from references 9-12, 29. The cells in HPT follow two main cell lineages: one branch is from Type 1 via Type 5 cells to semigranular cells (SGC); the other is from Type 1 to Types 2, 3, 4 cells to granular cells (GC). Astakine 1 is involved in proliferation, differentiation through the SGC pathway and release of hemocytes into the circulation, while the definite role of astakine 2 still remains to be shown.

of the SGC specific Kazal type proteinase inhibitor.^{62,80,81} Astakine 2 has been shown to increase proliferation in HPT cells in penaeid shrimp, whereas its role in crayfish is not fully understood although it has some stimulatory effect on GC differentiation.⁸¹

Table 2. Crustacean astakines

	Species	Accession Number
Astakine 1	<i>Pacifastacus leniusculus</i>	AY787656
Astakine 2	<i>Pacifastacus leniusculus</i>	EF568370-EF568371
	<i>Penaeus monodon</i>	AY787657
	<i>Litopenaeus vannamei</i>	FE148214
	<i>Homarus americanus</i>	FE535609
	<i>Carcinus maenas</i>	DW585080
	<i>Daphnia pulex</i>	FE329237

Transglutaminases (TGases) form a family of Ca²⁺-dependent enzymes catalysing posttranslational remodelling of proteins by cross-linking and this enzyme acts as a clotting enzyme in crustacean hemolymph coagulation.¹ TGase is one of the most abundant proteins in crustacean HPTs^{63,82} where its mRNA expression as well as enzyme activity is very high. TGase has been shown to play an important role in keeping the HPT cells in an undifferentiated stage inside the hematopoietic tissue and if expression of TGase mRNA is blocked, the cells start to differentiate and migrate out into the circulating hemolymph.⁶³ Interestingly astakine 1 seems to play a role in this process, since astakine by some unknown mechanism decrease extracellular TGase activity and induce cell migration.

ANTIVIRAL REACTIONS

Viruses remain a major obstacle to crustacean aquaculture. Among the viruses, the most intensively studied have been characterised from cultured penaeids such as the white spot syndrome virus (WSSV), yellow head virus (YHV) and Taura syndrome virus (TSV). We recently published⁵ a review on antiviral reactions in crustaceans covering the literature up to 2008 so here mainly some recent developments are discussed. The molecular mechanisms that underlie the majority of crustacean antiviral immune responses are still unknown and are only starting to be addressed. Recently, high throughput identification of genes and proteins (e.g., EST, SSH, microarrays) have been taken into use in an attempt to solve this.⁸³

Antiviral substances have been isolated from several crustaceans although the mechanism of this inhibitory activity remains unclear.⁸⁴ A well known cationic protein, antilipopolysaccharide factor (ALF) originally isolated from horseshoe crab⁸⁵ has been studied in crustaceans for its antibacterial activity.^{86,87} Crayfish ALF was up-regulated by a WSSV challenge and was shown to be involved in antiviral response against WSSV. Silencing of ALF resulted in higher rates of WSSV propagation both in the animals and in an HPT cell culture.⁸⁸ In contrast, enhanced expression of ALF in the crayfish by the administration of UV-treated WSSV led to lower viral replication and a partial protection against a subsequent challenge with the active virus.⁸⁸ Silencing of *Lv*ALF1 resulted in a significant increase of mortality in *L. vannamei* challenged by *Vibrio penaeicida* and *Fusarium oxysporum* but no protection against WSSV.⁸⁶ However, a study on a *P. monodon* recombinant ALFPm3, showed that this protein affected viral

infection both in shrimp and in crayfish HPT cell cultures.⁸⁹ The mechanism for antiviral activity of crayfish ALF or shrimp ALF*Pm3* is still unknown.

Cytokine activation through JAK/STAT pathway of a number of genes has been suggested in counteracting viral infection in *Drosophila*.¹² Flies deficient in the JAK kinase Hopscotch show increased susceptibility to *Drosophila C* virus and contain a higher viral load. These data indicate that flies produce antiviral molecules in a JAK-STAT-dependent way.⁹⁰ However, the WSSV immediate early gene (*ie1*) was shown to employ a shrimp STAT as a transcription factor to enhance its expression.⁹¹ Additional studies showed that shrimp STAT was activated in response to WSSV infection and the WSSV does not disrupt JAK-STAT pathway but benefits from STAT activation in the shrimp.^{92,93} Also, some components of the Toll pathway (Toll and Dif) have been shown to be of importance for the resistance against *Drosophila X* virus.⁹⁴ However, in shrimp a recent study found that a Toll like receptor (IToll) was not involved in antiviral immunity.⁹⁵ Further work is needed to reveal if other Toll-like receptors are necessary for antiviral responses in crustaceans.

Apoptosis

Apoptosis is a critical cellular process for removing unnecessary or potentially harmful cells and possibly for limiting viral spread.⁹⁶ Caspases are central effectors in apoptosis and if the *M. japonicus Pj*caspase gene was silenced, the WSSV-induced apoptosis was significantly inhibited and the number of viral copies increased, indicating that apoptosis may play an antiviral role.⁹⁷ This proposal however needs to be ascertained by more experiments since knocking down caspase-3 reduces mortality in Pacific white shrimp challenged with a low dose of WSSV but not with a high-dose of WSSV. This suggests that apoptosis in some cases may increase rather than decrease mortality in WSSV-challenged shrimp.⁹⁸

Similarly, the wide spread apoptosis in *P. monodon* infected with YHV is a major cause of dysfunction and death of the host. The expression of ribophorin I, a protein involved in apoptosis, was up-regulated and remained high until the moribund stage in YHV infected shrimp⁹⁹ whereas the defender against apoptotic death 1, a negative regulator of apoptosis, decreased dramatically after YHV challenge.¹⁰⁰ It ought to be stressed though that apoptosis can be triggered by a multitude of signals and much more need to be known about apoptosis in crustaceans before the mentioned findings can be properly evaluated.

Antiviral Activity Induced by RNA Interference or Injection of dsRNA

Injection of dsRNA/siRNA specific to viral genes can block viral disease progression. For instance, viral replication was efficiently suppressed with injection of WSSV-specific dsRNA/siRNA targeting VP19, VP28, VP281, or WSSV protein kinase in penaeid shrimp.^{5,101}

Recent studies reveal the existence of both innate (nonsequence specific) and RNAi related (sequence specific) crustacean antiviral phenomena.¹⁰² However, the protection induced by the innate pathway could be overwhelmed by a higher dose (8-fold) of infectious virus, suggesting it mediates a low degree of resistance. Two components of RNA silencing, Argonaute and Dicer, have been characterised from *P. monodon*^{103,104} but the mechanism for this silencing is still not clear. Injection of

CpG oligodeoxynucleotides (CpG ODNs; these are actually typical for bacterial DNA) mediates a protection against WSSV propagation, possibly via Argonaute and Dicer.¹⁰⁵ Recently a protein homologous to HIV transactivating response RNA-binding protein was found to bind Dicer and to inhibit WSSV replication in *F. chinensis*,¹⁰⁶ These studies hint that the RNAi machinery may play an important role for antiviral activity in crustaceans, although much efforts remain to fully establish the mechanism(s) for this activity.

Studies on the mechanism of these antiviral responses have been hampered by absence of genome, tools for genetic manipulation and mutants and stable long-term cell lines for in vitro studies. We have succeeded in developing an HPT cell cultures from crayfish, which can be a useful tool for gene functional studies in crustaceans.¹⁰⁷ These HPT cultures can also be used to replicate WSSV and to study host-virus interactions.¹⁰⁸

CLOTTING, SYNTHESIS OF ANTIMICROBIAL PROTEINS AND MELANISATION

Clotting

Clotting is an important reaction aimed at preventing hemolymph loss and microbial spread at sites of injury. The reaction has been extensively studied in crustaceans, in particular freshwater crayfish,¹⁰⁹ from which the first crustacean clotting protein was cloned.¹¹⁰ For a review on the subject and a comparison with the corresponding reaction in other arthropods see ref 111. In the shrimp *Marsupenaeus japonicus* RNAi silencing of clotting protein and a transglutaminase resulted in a defect clotting system.¹¹² Interestingly, challenging such animals with *V. penaeida* or WSSV resulted in higher mortalities. Although the reason for this effect is unknown it could mean that initiation of clotting also triggers the onset of other immune reactions and/or that the clotting reaction itself interferes with the propagation of these pathogens by e.g., entangling them.

Antimicrobial Proteins

Antimicrobial proteins are very important components of the immune system in many insects. They have received less attention in crustaceans, perhaps because their expression usually is not up-regulated as dramatically by the presence of microbial products as in holometabolous insects. In recent years a number of crustacean antimicrobial proteins (AMPs) have been purified and/or cloned. In some cases their effects on microbial growth in vitro have been investigated, but to what extent these activities reflect their importance in vivo is more difficult to assess. There are many kinds of crustacean AMPs that differ considerable in structure; two prominent groups with many members are the penaeidins and the crustins, thoroughly reviewed in Cuthbertson et al³ and in Smith et al⁴ respectively. Crustins have a wider occurrence among crustacean taxa than the penaeidins. However, a recently characterised spider crab AMP, hyastatin may judged from its Cys bond pattern be related to the penaeidins.¹¹³ The presence of a whey acidic protein (WAP) domain is a characteristic of crustins although WAP domains are present in many other types of proteins, e.g., proteinase inhibitors.¹¹⁴ A third group of AMPs are the antilipolysaccharide factors that also have

been implicated in antiviral reactions (see above). There are other potential sources of antimicrobial peptides and proteins. For example, some peptides released by limited proteolysis from hemocyanin exhibit antimicrobial properties.^{115,116}

In many insects AMP expression is governed by either the Toll or the imd pathways. As mentioned earlier, crustacean AMP expression tends to be more or less constitutive although several cases of inducible AMPs have been recorded.¹¹⁷ Some possible Toll pathway components were recently described in several shrimp species, e.g., Toll itself^{95,117-120} spätzle,¹²¹ relish,^{121,123} dorsal.¹²⁴ *L. vannamei* Relish and dorsal were shown to regulate the expression of penaeidin-4 in transfected insect cells. To what extent this reflects in vivo AMP expression in the shrimp remains to establish. In one study, the recombinant protein from the spätzle-like gene in *F. chinensis* was injected into the crayfish *Procambarus clarkii* (shrimps died if injected with this product), which resulted in an increase of transcript levels for crustin-2 but not for the other tested AMP genes.¹²¹ Also this finding needs confirmation by additional experiments. If transcription of one putative Toll-receptor was reduced by RNAi treatment in *L. vannamei* the animals became more susceptible to the bacterium *Vibrio harveyi* whereas the susceptibility to WSSV was unaffected.¹¹⁷ Whether this is due to any possible Toll effects on AMP production or other Toll-mediated effects is unknown and, furthermore, the number of putative Toll receptors in shrimp is not known. However, it should be noticed that at least crustin expression was unaffected by the presence of LvToll dsRNA in this case. In another study specific RNA interference of *L. vannamei* crustin caused in increased susceptibility to a related bacterium, *V. penaeicida*.¹²⁵ Also a putative imd homologue has been reported from *L. vannamei*¹²⁶ that requires further functional studies. Silencing of a Relish homologue in the shrimp *F. chinensis* resulted in a lowered penaeidin 5 transcription upon bacterial challenges, a result which could be interpreted as an imd pathway exists in shrimps. However, the extent (if any) to which AMP synthesis is regulated by Toll and imd homologues in crustaceans is still far from settled.

The Prophenoloxidase Activating System

The melanisation reaction is an important immune reaction and numerous studies in different types of animals have attested the crucial role of the phenoloxidase system in combating microbial infections. A large body of pioneering work (for reviews see refs. 127,128) on the proPO was carried out using freshwater crayfish as model. Recently several RNA interference studies aiming at the transcription of the proPO gene or genes whose products are involved in proPO activation have been carried out. In *P. leniusculus* reduced levels of proPO led to an increased susceptibility to the serious bacterial pathogen *Aeromonas hydrophila*.¹²⁹ Reducing the levels of pacifastin, a specific inhibitor of the proPO-activating proteinase, resulted into a higher melanisation capacity and increased survival to the pathogen. Also in vitro the products from an active crayfish PO are reducing the growth of several bacteria, both gram positive and gram negative species.¹³⁰ Two studies^{131,132} using RNAi depletion of proPO transcripts in *Penaeus monodon* have led to similar conclusions. Reduced transcription of the proPO genes or a gene for proPO-activating proteinase resulted into higher mortality upon challenge with *Vibrio harveyi*. An extensive study carried out with *M. japonicus* showed that after depletion of proPO, bacterial counts in hemolymph and other tissues increased.¹³³ Since these animals were not challenged either endogenous bacteria, or bacteria taken up from the rearing tank must be responsible for the increasing bacterial

loads. In contrast to animals injected with control dsRNA the proPO-depleted animals exhibited increased mortality and reduced hemocyte numbers. The increased mortality could to some degree, although not completely, be counteracted by the administration of antibiotics. Interestingly, an array with more than 2000 shrimp genes showed that 28 genes were up-regulated and 49 down-regulated on the third day after starting the proPO gene interference. Thus, a consequence of the lowering of the PO levels could be that other immune factors are produced at a reduced rate, decreasing the immune capacity of the animal. Among the genes down-regulated after proPO depletion were some AMPs (penaeidin and crustin) and two Kazal type inhibitors. Kazal type proteinase inhibitors have been shown to interfere with the growth of several bacteria. It is still not known to what extent the bacteriostatic activity is due to inhibition of microbial proteinases or to other effects (for a review see ref. 134). A single animal can produce a large number of different Kazal variants, a fact suggesting but not proving that these inhibitors is under selection pressure from various microbes.¹³⁵ Also other proteinase inhibitors, e.g., alpha-2-macroglobulin have been shown to be produced in a large number of sequence variants.¹³⁶ As is the case in the crustacean Kazal inhibitors the sequence variation is especially evident among those amino acid residues that takes part in the interaction with proteinases.

Once active PO has been produced, regulatory mechanisms to ensure that melanisation does not proceed uncontrolled for unlimited periods are likely to exist. Several such nonproteinaceous compounds and phenoloxidase inhibiting peptides are known from insects. One such control of excessive melanisation is likely to be carried out by the recently identified melanisation inhibition proteins (MIPs) from crayfish¹³⁷ and meal worms.¹³⁸ MIPs have been shown to prevent both proteolytic activation of proPO as well as to interfere with the melanin synthesis from quinones. The prevention of the melanin formation occurs at a late step(s) after the steps catalysed enzymatically by PO. Crayfish MIP is from an evolutionary viewpoint an interesting protein, since it is bearing significant sequence similarity with vertebrate ficolins, including a binding site for Ca²⁺. In vitro mutagenesis of this site has demonstrated its importance for full inhibitory activity.¹³⁷

CONCLUSION

Emerging evidence from other systems such as insects indicate that there is substantial cross-talk between different arms of the innate immune defence. For example, in several insect species some of the proteinases and regulatory serpins that are part of the Toll activation cascade and the proPO-activating system are shared.^{139,140} This makes sense since it is likely that achieving an efficient response towards many pathogens will require that several parts of the immune system cooperate. It will be interesting to see whether examples of such cross-talks within crustacean innate immunity will be discovered. To conclude, large progress in elucidating innate immune pathways have been made in recent years. In the near future genomic information will be available for additional crustacean species and the speed with which progress in this area is made will increase further. However, we are still far from efficient therapies for the diseases that are plaguing crustacean aquaculture.

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ECHINODERM IMMUNITY

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Abstract: A survey for immune genes in the genome for the purple sea urchin has shown that the immune system is complex and sophisticated. By inference, immune responses of all echinoderms may be similar. The immune system is mediated by several types of coelomocytes that are also useful as sensors of environmental stresses. There are a number of large gene families in the purple sea urchin genome that function in immunity and of which at least one appears to employ novel approaches for sequence diversification. Echinoderms have a simpler complement system, a large set of lectin genes and a number of antimicrobial peptides. Profiling the immune genes expressed by coelomocytes and the proteins in the coelomic fluid provide detailed information about immune functions in the sea urchin. The importance of echinoderms in maintaining marine ecosystem stability and the disastrous effects of their removal due to disease will require future collaborations between ecologists and immunologists working towards understanding and preserving marine habitats.

INTRODUCTION

Echinoderms are a phylum of ubiquitous benthic marine invertebrates, found in a continuous distribution from the intertidal zone to the deepest depths of the ocean explored to date. They are the sister group to the chordates and are the basal deuterostomes.¹ Approximately 7000 extant echinoderm species have been described, falling into five classes—echinoids (sea urchins and sand dollars), holothurians (sea cucumbers), asteroids (sea stars), crinoids (sea lilies and feather stars) and ophiuroids (brittle stars), as well as a number of extinct classes known from the fossil record. Many members of the phylum are large and long lived.²

Echinoderms are central players in many benthic ecosystems, variously as herbivores, carnivores and detritivores. They possess a simple gut which opens into an anus on the aboral surface, or, for crinoids, on the oral surface of the animal.¹ All echinoderms have basic pentameric radial symmetry in their adult forms and most metamorphose from bilateral larvae found in the plankton that function as the dispersal stage, feed and produce the adult rudiment. Adults lack obvious sensory organs or any kind of head and locomote largely by means of tube feet, which function by means of a water vascular system unique to the phylum. Activities are coordinated by a simple neural ring that communicates to five nerve bundles into the five ambulacral grooves that run longitudinally along each radial segment of the body. All echinoderms have some form of calcified skeleton composed of aragonite, which is more or less extensive in the various classes. At one extreme, echinoids are completely enclosed in a calcified test, with only a thin layer of muscular tissue lining the interior of the body cavity (or coelom) while holothurians possess a highly muscularized body wall in which the exoskeleton is reduced to microscopic ossicles dispersed throughout the dermis. The coelomic cavity contains the gonads and gut of the animal, bathed in coelomic fluid. This coelomic fluid is key to the immunological capabilities of echinoderms, being the medium in which the immunocytes or coelomocytes reside, and which also contains antimicrobial molecules.

A larval sea star was first used by Metchnikoff³ to demonstrate that invertebrate immune cells could recognize the presence of and respond to (encapsulate) a rose prickle. Subsequent characterization of the echinoderm immune system in adults relied on allograft rejection assays in sea cucumbers, sea stars,^{4,5} and sea urchins,⁶⁻⁸ and demonstrated the ability of echinoderms to recognize self from nonself. Later evaluation of the sea urchin graft rejection kinetics illustrated the innate characteristics of echinoderm immunity.⁹ Although the immune system is entirely innate, it has recently been found to be highly complex and sophisticated^{10,11} and is clearly effective given that echinoderms have survived as a phylum for 450-500 million years and that individual echinoderms can survive the constant assault by pathogens in the marine environment (reviewed by ref. 12) for upwards of 100 years.²

COELOMOCYTES, THE IMMUNE MEDIATORS IN ECHINODERMS

The body cavity of echinoderms is filled with coelomic fluid,¹³ which bathes the internal organs and forms the fluid medium in which the coelomocytes are suspended. The composition of coelomic fluid (CF) is similar to sea water in terms of minor dissolved salts and other minerals and contains proteins, the best-characterized of which are those involved in antipathogen responses. Indeed, whole CF (wCF) is a complex tissue that



Figure 1. Live coelomocytes from the sea urchin, *Paracentrotus lividus*. A) petaloid phagocyte. B) red spherule cell. C) colorless spherule cell. D) vibratile cell. Scale bar = 5 microns. Images taken by R. Bonaventura.

mediates responses to wounding and microbial infections by undergoing reactions such as opsonization, coagulation, encapsulation and phagocytosis. Initial publications on echinoderm coelomocytes appeared in the late 19th century (e.g., ref. 14) and since then many reports have appeared describing the morphologies and functions of the various coelomocytes^{9,15-17} (reviewed by refs. 18,19). Observations of live cells in CF from sea urchins suggest three basic categories of coelomocytes; phagocytes, spherule cells (also called amoebocytes,^{20,21} or morula cells) and vibratile cells (Fig. 1); however further analyses suggest additional categories (Table 1) (reviewed in ref. 22). Within the spherule cell category, there are red and colorless spherules within the cells (Fig. 1B,C) and there are three different categories of phagocytes depending on cytoskeletal morphology and size (Fig. 2). Additional cell types have been noted in nonechinoid species including crystal cells, fusiform cells and progenitor cells, about which very little is known. The descriptions below refer to sea urchin coelomocytes, which are the best studied.

Phagocytes

Phagocytes have been variously referred to as leukocytes, or as bladder, petaloid or filiform phagocytes in the literature and constitute the most abundant type of coelomocyte (Table 1) in the coelomic fluid. In suspension the cells appear in a petaloid form (Fig. 1A;

Table 1. Coelomocytes in sea urchins

Cell Type	% in Coelomic Fluid	Function
Phagocyte Types	Total phagocytes	Encapsulation, Opsonization,
Type 1—Discoidal cells	<i>Sp</i> * 40-80%	Graft rejection, Chemotaxis,
Type 2—Polygonal cells	<i>Sd</i> 67%	Phagocytosis, Antibacterial
Type 3—Small phagocytes	<i>Pl</i> 80%	activity, Cellular clotting
Red spherule cells	<i>Sp</i> 7-40%	Oxygen transport. Antibacterial
	<i>Sd</i> 8%	activity from echinochrome A.
	<i>Pl</i> 4.7%	
Colorless spherule cells	<i>Sp</i> 3.7-25%	Cytotoxicity, Clotting?
	<i>Sd</i> 6.5%	
	<i>Pl</i> 7.8%	
Vibratile cells	<i>Sp</i> 11.9-20%	Movement or agitation of
	<i>Sd</i> 18.5%	coelomic fluid? Associated
	<i>Pl</i> 7.5%	with clotting.

**Sp*, *Strongylocentrotus purpuratus*. *Sd*, *Strongylocentrotus droebachiensis*. *Pl*, *Paracentrotus lividus*
 Information collected from references 7, 16, 17, 19, 22, 24, 28, 33, 34, 38, 39, 50, 95, 222 and 223.

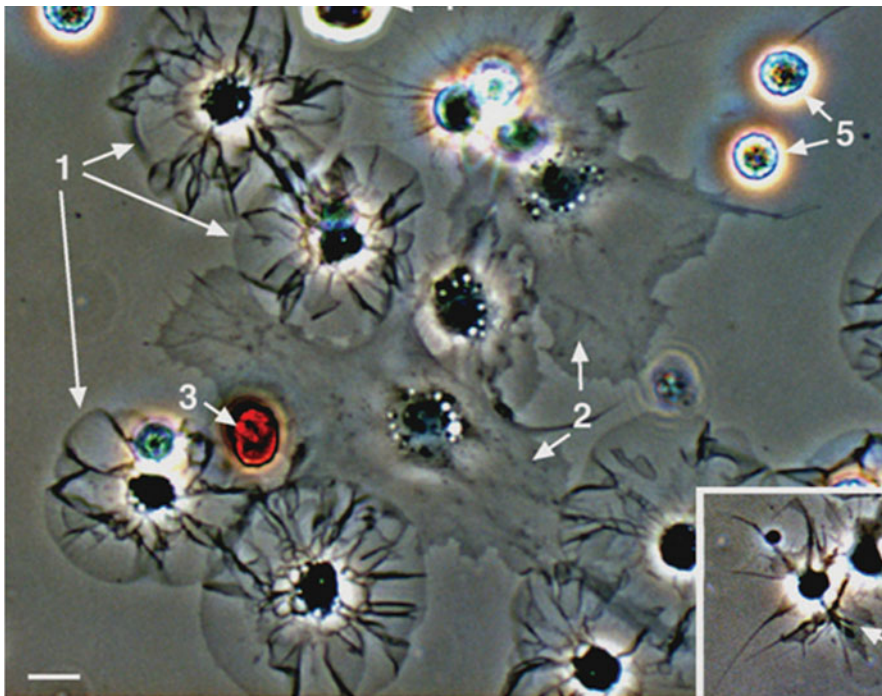


Figure 2. Coelomocytes from the sea urchin *S. droebachiensis*. Cells were withdrawn in anticoagulant and settled onto a glass coverslip. All cell types are shown and labeled with numbers. Large phagocytes; 1 = discoidal phagocyte; 2 = polygonal phagocyte; 3 = red spherule cell; 4 = colorless spherule cell; 5 = vibratile cell (the lower cell has lost the prominent flagellum seen in the upper cell). Inset; 6 = small phagocyte. Bar = 10 microns.

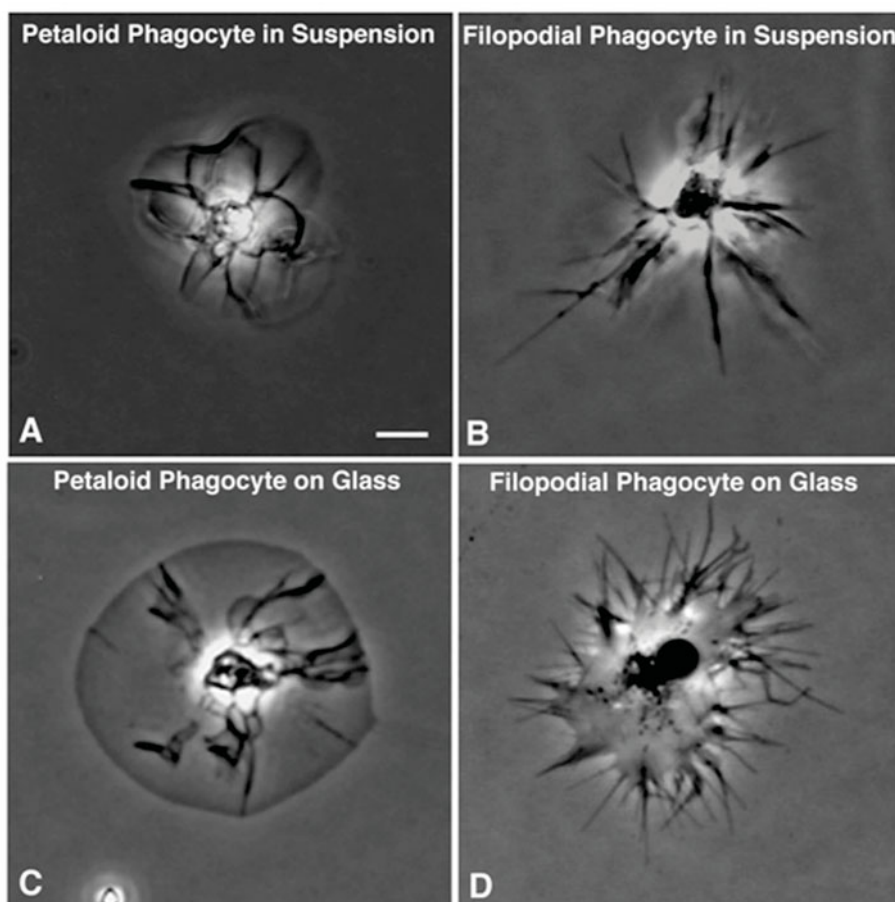


Figure 3. Phagocytes undergo an inducible petaloid/lamellipodial to filopodial/filiform shape change that is important in the coelomic fluid clotting process. This shape change occurs in cells in suspension; panel (A), petaloid cell; panel (B), filopodial cell. Shape change also occurs when cells are attached to glass substrate; panel (C), petaloid (lamellipodial) cell; panel (D), filopodial cell. Shape change was induced by hypotonic shock. Bar = 10 microns; magnifications of panels (A-D) are equivalent.

Fig. 3A) in which numerous cytoplasmic lamellipodia give the impression of the petals of a flower.²³ More recent work indicates that the phagocyte population actually consists of several different cell types that can be distinguished based on cytoskeletal morphology and organization, actin-based motility patterns,²⁴⁻²⁷ and differential gene and protein expression.^{19,28,29} Phagocytes are most easily distinguished as three morphologies in spread cells; two types of large phagocytes²⁴⁻²⁷ and small phagocytes.^{28,29} The discoidal-shaped phagocytes (Fig. 2) are large phagocytes that display rapid, actin-based retrograde/centripetal flow over their entire diameter, and contain a dense cortical actin meshwork rich in the Arp2/3 complex (Fig. 4C), radial actin bundles that connect the cortex to the perinuclear region, and a centralized distribution of myosin II (Fig. 4B).^{25,27} The retrograde/centripetal flow activity in discoidal cells tends to restrict the distribution of microtubules (Fig. 4A), organelles (endoplasmic reticulum, mitochondria, lysosomes and granules) and

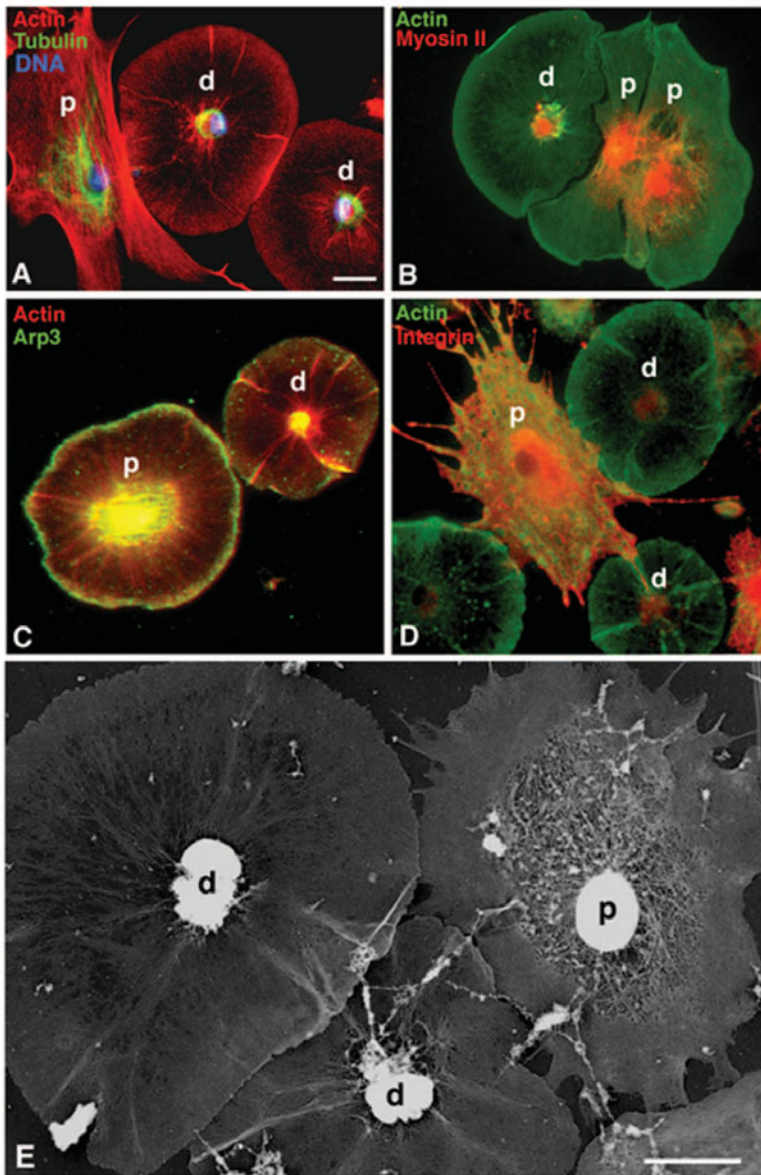


Figure 4. Large phagocytes of the discoidal (d) and polygonal (p) subpopulations can be differentiated based on cytoskeletal organization. Actin (panels A-D) in discoidal cells displays a broad cortical meshwork containing Arp3 (panel C) that feeds into a radial array of centralized bundles. The cortical actin and Arp3 meshwork is more limited in polygonal cells (panel C) and cells often contain prominent actin bundles oriented either radially in discoidal cells (panel C) or parallel to the long axis in polygonal cells (panels A,D). Both myosin II (panel B) and microtubules (panel A) are perinuclear in discoidal cells and more widespread in polygonal cells. β integrin expression is restricted to the polygonal cells (panel D). The significant difference between the structural organization of the cytoskeletons of the two large phagocytes is best appreciated in TEM images of critical point dried and rotary shadowed replicas of detergent extracted cells (panel E). Bar = 10 microns; magnifications of panels A-D are equivalent.

associated kinesin motor proteins to the perinuclear region.²⁶ The second type of large phagocyte is polygonal-shaped cells (Fig. 2, labeled 2), which display retrograde flow in a thin rim along the cell margin corresponding to more limited distribution of the Arp2/3 complex-containing actin meshwork (Fig. 4C). These large cells are integrin-positive (Fig. 4D) and the interior cytoplasm contains elongate actin bundles associated with myosin II (Fig. 4A-D),^{25,27} as well as a broad array of microtubules (Fig. 4A) and associated organelles and kinesin (Fig. 4E).²⁶ Furthermore, both cell types undergo significant shape changes from a lamellipodial/petaloid to a filopodial form (Fig. 3B; see also ref. 20), inducible by the calcium-dependent clotting process, treatment with hypotonic shock²³ or calcium ionophore.³⁰

Large phagocytes display phagocytic activity associated with clearance of bacteria, xenogeneic cells, foreign particles and proteins, both *in vivo* and *in vitro* (reviewed in ref. 18). Phagocytes have also been implicated in encapsulation and graft rejection responses, as well as cytolytic/cytotoxic reactions,^{8,31} (reviewed in refs. 18, 19). Phagocytes participate in the cellular clotting process, in which their filopodial form facilitates the cell-cell interactions leading to cellular clot formation and retraction.^{16,23} Based on their size, preponderance in the CF and their phagocytic and encapsulation functions, the large phagocytes appear to be the major mediators of cellular immunity in echinoderms.

The third and least common form of phagocyte is the small phagocyte (Fig. 2, labeled 6)^{28,29} of which some express the Sp185/333 family of highly diverse immune response-related proteins (see below). These cells are characterized by their small size relative to the large phagocytes, numerous small cytoplasmic granules and filopodial morphology (Fig. 5). The morphology of small phagocyte filopodia appears different from that of large phagocytes in that they are less numerous, thicker and often contain periodic knobs and microtubule bundles (Fig. 5D).²⁹

Spherule Cells

Red spherule cells (Fig. 1B; Fig. 2, labeled 3; Table 1) have been called amoebocytes, spherulocytes, morula cells, pigment cells, granulocytes and eleocytes in the literature, and are small cells filled with spherical red granules containing echinochrome A, a naphthoquinone pigment with antibacterial properties. Red spherule cells have been reported to move towards bacteria to initiate an encapsulation response and to degranulate in the presence of bacteria.³²⁻³⁵ They have been reported to migrate to and form a rim around the edge of wounds, infections and tissue grafts.^{8,36,37} When settled onto a substrate, red spherule cells exhibit dynamic actin-based, amoeboid mobility, which may account for their ability to surround wounds and infections. Colorless spherule cells (Fig. 1C; Fig. 2, labeled 4; Table 1) contain granules and have therefore been referred to as morula cells (from Latin for 'mulberry'). Their properties and functions are not well known, however, a recent study suggests that these cells have potent cytolytic activity that is augmented by the presence of phagocytes.³⁸

Vibratile Cells

Vibratile cells are round, highly motile, flagellated cells that contain large cytoplasmic granules (Fig. 1D; Fig. 2, labeled 5; Table 1). Exocytosis of these granules may trigger the initiation of the clotting reaction^{32,39} (Sacchi and Smith, unpublished), however this hypothesis will require additional testing.

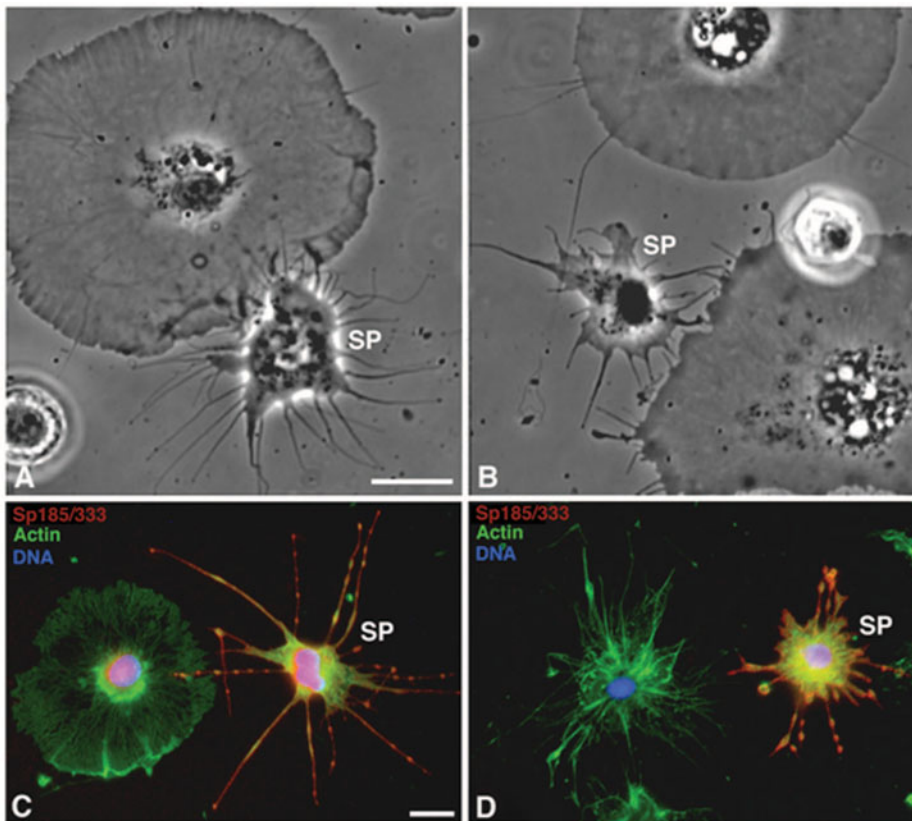


Figure 5. Small phagocytes in living preparations. Small phagocytes (SP) can be identified based on their relative small size, unusual filopodial morphology and granular cytoplasm (panels A,B). These cells are clearly labeled by antibodies against the Sp185/333 family of proteins (SP cell in panels C,D), whereas the petaloid and filopodial discoidal large phagocytes do not stain for Sp185/333. The Sp185/333 labeling of the small phagocytes highlights the atypical nature of their filopodia which are often thick and contain knobs, not common for filopodia of discoidal phagocytes (left-hand cell in panel D). Note that in some small phagocytes the filopodia become very elongate (panels A,C). Bar = 10 microns.

Coelomocytes in Other Echinoderms

Asteroids clear their coelomic cavities efficiently despite having only two major cell types, phagocytes and colorless spherule cells.^{22,40} Rapid activation and differentiation of the phagocytes into multiple functional cell types upon bacterial infection is likely responsible for efficient microbial clearance.⁴¹ The phagocytes in the sea star, *Asterias rubens*, are present in a continuum of small to large cells that change relative abundance upon immune stimulation. Larger cells become more numerous and exhibit strong phagocytic reactions to bacteria and the total number of phagocytes triple 24 hrs after challenge.⁴¹ Increased numbers of total coelomocytes have also been noted in response to wounding and hypoxic conditions.⁴² Swift responses, immune challenges, and the possible differentiation of phagocytes into cells with augmented phagocytic activity is an interesting approach to infection in sea stars that have fewer morphotypes of coelomocytes compared to echinoids.

Box 1. Larval immune cells.

While most investigations of echinoderm immunity have been carried out in the adult, the larvae of indirectly-developing species possess specialized mesodermal cells derived from embryonic blastocoelar cells that display a variety of immune behaviors. The immune activities of the larval immunocytes were recognized at the inception of cellular immunity by Metchnikoff and his infamous experiment of encapsulation of a rose prickle by blastodermal cells of a sea star larva.³ This was the first demonstration of phagocytosis and encapsulation and led to a Nobel prize. More recently, recognition and phagocytic behavior of larval immune cells responding to bacteria or yeast injected into the blastocoel has been demonstrated.^{218,219} The immune cells in the embryo and larvae are the blastocoelar cells located in the blastocoel²²⁰ and the pigment cells in the ectoderm¹⁹ that produce echinochrome A.²²¹ Embryonic and larval immune cells are also known to express immune factors, including the complement homologue SpC3⁹² and the immune response genes *Sp185/333*.^{10,70,92} Besides immune effector functions, these cells carry out a number of immune functions including surveillance-like behavior, wound healing and the expression of a complex suite of immune recognition, regulatory and effector genes¹⁰ (Rast, unpublished). As more data are accumulated about this simple stage of the sea urchin life cycle, postgastrula embryos and larvae are becoming an increasingly valuable model system in which to investigate immunity.

Holothurians have lymphocytes, morula cells, amoebocytes, crystal cells, fusiform cells and vibratile cells.^{22,40} In addition to phagocytosis, holothurian coelomocytes exhibit brown body formation in response to multicellular parasites. These are pigmented aggregates of phagocytes and spherule cells that encapsulate parasites that are too large to be phagocytized.⁴³ Brown bodies in the sea cucumbers, *Holothuria polii* and *H. tubulosa*, are tissue nodules containing entrapped parasites that are first surrounded by a fibrous, electron-dense, noncellular layer—probably melanin—and then covered by layers of specialized cells^{44,45} including subsets of elongated phagocytes⁴⁴ and spherule cells.⁴⁶ Spherule cells likely degranulate to chemically kill and/or degrade the invader,⁴⁶ while other cells within the brown bodies possess active phenoloxidase resulting in melanization.⁴⁴ Brown bodies are progressively eliminated from the animal through coelo-rectal canaliculi.^{44,45,47} Essentially, coelomocytes appear to have two means for clearing microbes and parasites; phagocytosis and encapsulation.

Crinoids have phagocytes, red spherule cells and colorless spherule cells (or morula cells), whereas ophiuroids have phagocytes, colorless spherule cells and possibly crystal and vibratile cells.^{22,40} Because little work has been done on these classes of echinoderms, little is known about the functions of coelomocytes in these groups.

CLOT FORMATION

Clotting is mediated by a variety of agglutination factors. It is an important response to injury in echinoderms functioning to block loss of CF resulting from wounds, and to sequester pathogens and prevent their invasion throughout the body. A 220 kDa agglutinating factor thought to be involved with coagulation is secreted from coelomocytes of the sea cucumber *Holothuria polii*.⁴⁸ Coelomocytes from the purple sea urchin express amassin-1, a 75 kDa, multidomain protein with an olfactomedin domain that is stabilized by multiple disulfide

Table 2. ESTs from the purple sea urchin and a sea cucumber

Category	Examples
Defense	Sp185/333, DD104, DD186, Complement C3, Complement factor B, Galectin, SpEchinoidin and other C-type lectins, Thrombospondin, Amassin, Integrin- β C, Ferritin, Melanotransferrin, Annexin, Serum amyloid A, Echinonectin, Fibrinogen-like protein, Ficolin-like protein, Kazal-type serine proteinase inhibitors
Cytoskeleton	Actin, Gelsolin, Cofilin, Thymosin β , Protein tyrosine kinase-9, Fascin, Tubulin, Microtubule-associated protein
Signal Transduction	SpTie1/2, Steroid hormone receptor, LPS-induced TNF α , Protein tyrosine phosphatase receptor, Adenylyl cyclase, Receptor for activated protein kinase C, Guanine nucleotide binding protein β , Rho-GTPase, Rho-GDP dissociation inhibitor, GTPase-activating protein, FK Binding Protein-12, GF14/14-3-3
Nuclear Activities, Splicing	RNA/DNA binding protein, DNA methyl transferase-associated protein, Nonhistone nucleic acid-binding protein, Splicing factor 30, Paraspeckle protein, ET putative translation product, Heterogeneous nuclear ribonucleo-protein R, PolyA binding protein
Transcription factors	Steroid hormone receptor, LPS-induced tumor necrosis factor α , Immediate early-response protein, SpRunt, SpNFkB
Protein Metabolism, Synthesis, Degradation	Translation elongation factor 1 α , 1 γ , 2, Translation initiation factor, Peptide chain release factor, Signal sequence receptor, Protein disulfide isomerase, Binding Protein, Heat shock protein, gp96, Presenilin, Proteosome subunit, Ubiquitin
Molecular Transport	Dynein, Kinesin, Vesicle trafficking protein, Coated vesicle membrane protein, ER transport protein, Vacuolar protein sorting protein, Rab7, Rab5-interacting protein, Sec22, vSNARE, Mannose-6 phosphate receptor
Endosomal System	Vacuolar H ⁺ -ATPase, Cathepsins, Lysozyme, Arylsulfatase
Proliferation, Apoptosis	Polo-like kinase, Bax inhibitor-1, Allograft inflammatory factor-1
Metabolism	ATP synthase, Cytochrome C oxidase subunit 1, 4, Cytochrome b, NADH dehydrogenase subunit 2, 4, 5, Fatty acid desaturase, ATP/ADP translocase, Vacuolar ATP synthase subunit, Na ⁺ /phosphate cotransporter, Na ⁺ K ⁺ -transporter, Citrate synthase, Malate dehydrogenase

From references 49, 52, 53, 88 and 165.

bonds (Table 2; Table 3;⁴⁹ Dheilly, Raftos and Nair, unpublished). Amasin-1 functions in cellular clot formation through homooligomerization and cross-linking coelomocytes.⁵⁰ A search of the sea urchin genome identified a putative receptor for amassin-1, colmedin, which also has an olfactomedin domain and is expressed in coelomocytes.⁵¹

Other molecules involved in clotting mechanisms, including transglutaminase homologues, were identified during an expressed sequence tag (EST) study from gut tissue of the sea cucumber *Holothuria glaberrima*.^{52,53} Many of the genes referenced from

Table 3. Proteomic analysis of whole coelomic fluid from *Strongylocentrotus purpuratus*

Classification	Examples
Immune Response	Complement components (SpC3, SpBf), Scavenger receptors cysteine-rich, Sp185/333
Pathogen Destruction	Arylsulfatase and other lysosomal enzymes, α -2-macroglobulin
Clotting	Amassin, Annexin V, Von Willebrand factor
Metal binding	Major yolk protein, Transferrin, Ferritin, Ceruloplasmin
Cytoskeleton	Actin, Profilin, Fascin, Cofilin, Gelsolin, Myosin, Microtubule-associated protein, Arp 2/3 complex proteins, Coronin, Tubulin, α -actinin, Tetraspannin, Talin, Vinculin, Rab
Cell Adhesion	Integrin, NCAM, Selectin, Cadherin, Fibronectin
Signalling	Ras
Cytoplasmic Enzymes	Oxidative enzymes

From Dheilly, Raftos and Nair, unpublished.

these EST studies were originally studied in the context of the intestinal regeneration that occurs after sea cucumbers expel their internal organs as part of an antipredation response, and then regenerate the entire digestive tract in 30 days. As such, evisceration and regeneration exposes the coelomic cavity to microorganisms from the surrounding seawater and the animal's own enteric microflora. Thus, genes related to immune function and tissue regeneration are upregulated in intestinal tissues undergoing regeneration and are likely essential for the animal to survive the process.⁵²⁻⁵⁵

KEY MOLECULES IN ECHINODERM IMMUNITY

Proteins with Leucine Rich Repeats

Toll-Like Receptors

The Toll-like receptors (TLRs) are best characterized in insect and mammalian systems, in which they form small gene families of ~10 genes.^{56,57} These receptors have an N-terminal solenoid-like leucine-rich repeat (LRR) ectodomain, a transmembrane region and a cytoplasmic Toll/interleukin 1 receptor (TIR) in the C-terminus.⁵⁸ The genes are often encoded in a single exon. There are more than 200 TLRs in the *Strongylocentrotus purpuratus* genome, far more than has been identified in any other species to date.¹⁰ These genes fall into three categories: a large family of more than 200 genes with vertebrate-like structure; a set of three genes with structure similar to *Drosophila* Toll (which differs from that of the vertebrate TLRs and *Drosophila* Toll-9⁵⁹) and a family of five genes with a short truncated ectodomain structure (Fig. 6). In vertebrates, the ectodomain consists of canonical leucine-rich repeats flanked by specialized, cysteine-rich domains (designated single cysteine cluster, sccTLR). In

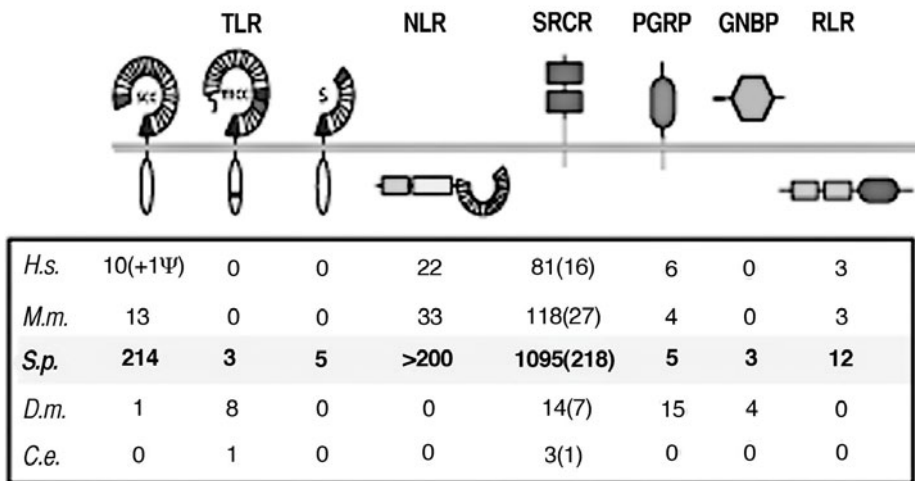


Figure 6. Pattern recognition receptor gene families are significantly expanded in the purple sea urchin with respect to other model organisms. The numbers of gene models encoding Toll-like receptors (TLR), Nod-like receptors (NLR), scavenger-receptor cysteine-rich (SRCR), peptidoglycan recognition proteins (PGRP), Gram negative binding proteins (GNBP) and RIG-I-like receptors (RLR) for human (*H.s.*), mouse (*M.m.*), purple sea urchin *S. purpuratus* (*S.p.*), fruit fly *Drosophila melanogaster* (*D.m.*) and nematode *Caenorhabditis elegans* (*C.e.*)²¹² are shown. Given the complexity associated with identifying SRCR gene models accurately, the number of SRCR domains is shown, with the number of gene models that contain multiple SRCR domains indicated parenthetically. In the electronic version of this chapter, colors are used to illustrate protein domains and to emphasize the sea urchin gene model numbers. A color version of this image is available at www.landesbioscience.com/curie.

contrast, the *Drosophila* Toll has specialized domains that are located in the center of the ectodomain (multiple cysteine cluster; mccTLR).⁵⁸ The mccTLRs may be the ancestral form,¹⁰ as mccTLR genes are present throughout eumetazoans, with the exception of the vertebrate lineage, in which it has apparently been lost. The sea urchin short TLRs have distant similarity within the TIR domain to the ancient Toll genes.⁶⁰

The large family of sea urchin TLRs is unusual in both its multiplicity and the apparent rapid diversification of some of its subfamilies. The ~210 genes in this family can be divided into seven subfamilies based on phylogenetic analysis of the TIR domain. Some of these subfamilies are composed of many members that differ primarily within their ectodomains while the TIR domains show greater conservation. Divergence within the leucine-rich repeats takes a number of forms, including point mutations, insertion-deletions between LRRs and insertion-deletions of whole LRR units.¹⁰ The extensive diversity of this class of sea urchin TLRs, along with a relatively large proportion of pseudogenes, differs from the more conserved picture seen in vertebrate TLR evolution⁶¹ and suggests that this complex family of receptors function fundamentally differently.⁶⁰

Many of the sea urchin TLR families are expressed most highly in coelomocytes in addition to gut tissue.¹⁰ Expression of sea urchin TLRs is not detected in the embryo but many families are expressed in the feeding larvae. Thus, expression patterns and diversity of sea urchin TLRs are consistent with an immune rather than developmental function. Consistent with this is the fact that extensive investigations of vertebrate TLRs only show immune functions.

NOD-Like Receptors

The second family of expanded immune receptors in the sea urchin is the NOD-like receptors (NLRs).¹⁰ NLRs are cytoplasmic pattern recognition receptors (PRRs) that recognize a variety of microbial signatures, including LPS, peptidoglycan, dsRNA and flagellin.⁶² Structurally, NLRs are composed of C-terminal LRRs, a central NACHT domain and one of several N-terminal domains that function in protein-protein interactions. NLRs in mammals function in the immune response by serving as scaffolding proteins to assemble protein complexes that lead to the activation of the NF κ B and MAPK signalling pathways. NLRs have also been shown to activate inflammatory caspases, including caspase-1, which is responsible for the processing of pro-interleukin-1 β .⁶³ There is a significant expansion of the NLR gene family in the *S. purpuratus* genome, which has over 200 NLR gene models compared to vertebrates that typically have ~20 NLR genes (Fig. 6).¹⁰ NLRs appear to be restricted to the deuterostome lineage and have not been identified in any of the sequenced protostome genomes, including *Drosophila* or *Caenorhabditis elegans*. As in vertebrates, the sea urchin NLR genes contain LRRs, a central NACHT domain and an N-terminal protein-protein interaction domain. However, unlike in the vertebrate system, this N-terminal domain is most commonly a DEATH domain, another member of the death-domain superfamily. A small number of sea urchin NLRs also contain CARD domains at the N-terminus, while the PYD domain is absent outside of the vertebrate lineage. Although many of the NLR gene models encode C-terminal LRR domains, these domains are lacking from some sea urchin gene models, which may be due to problems with accurate computational prediction of gene structure. The sea urchin NLR family appears to be the result of a sea urchin-specific gene expansion, which is seemingly more diverse than the similarly expanded sea urchin TLR family. The extent of this diversity is likely underrepresented given the incompleteness of the gene models, particularly within the LRR region. Although the function of the sea urchin NLRs is unknown, it is notable that they are most highly expressed in the gut,¹⁰ and therefore may be involved in managing gut microflora. This mimics the role of the NLR NOD2 protein in mammalian systems, which when mutated, results in inappropriate inflammatory reactions in the gut tissue and leads to Crohn's disease.⁶⁴ The LRR gene families in the sea urchin are greatly expanded compared to families in vertebrates and insects and it is noteworthy that the LRR families in amphioxus are also expanded,⁶⁵ suggesting that these receptors play an important role in the innate immune functions of these deuterostome invertebrates.

***Sp185/333*—A DIVERSE FAMILY OF GENES AND PROTEINS EXPRESSED IN RESPONSE TO IMMUNE CHALLENGE**

The *Sp185/333* Gene Family

The initial discovery of the *Sp185/333* family was the result of an EST analysis of transcripts that are upregulated in response to LPS challenge (see Box 2).⁴⁹ The diversity observed among the *Sp185/333* transcripts is intriguing for a putative immune response repertoire and resulted, in part, from an extraordinarily diverse gene family. The *Sp185/333* genes are small with two exons, of which the first is short and encodes a hydrophobic leader sequence, while the second encodes the remainder of the highly variable protein. The *Sp185/333* genes are atypical for *S. purpuratus* in three respects: (1) the second exon

ranges in size from 771-1431 base pairs (bp), which is notably larger than the average exon length (100-115 bp) as characterized from all gene models in the genome; (2) the intron is smaller than average (~400 bp, compared to an average intron size of ~750 bp); and (3) the average *S. purpuratus* gene has 8.3 exons.^{11,66} The most interesting aspect of the *Sp185/333* genes is the structure of the second exon. It is composed of contiguous blocks of sequence called *elements* that are defined from sequence alignments that require the insertions of large gaps (Fig. 7).^{49,66,67} Elements are variably present or absent in different genes (and transcripts, see Box 2) in recognizable mosaic combinations that have been called *element patterns*. Each element is actually a set of sequences that are similar but not necessarily identical—elements in the genes and transcripts differ by single nucleotide polymorphisms (SNPs) and small insertions or deletions (indels). Much of the sequence diversity among the gene results from the element patterns in the second exon. Although there are only six *Sp185/333* genes assembled in the v. 2.1 build of the *S. purpuratus* genome, three independent lines of evidence suggest that the gene family is composed of 40 to 60 paralogous loci: quantitative PCR (qPCR) of genomic DNA,⁶⁸ statistical estimates based on the frequency with which unique genes were cloned,⁶⁹ and an estimate based on results from screening two BAC libraries for *Sp185/333* genes combined with assumptions about gene linkage.⁷⁰

The *Sp185/333* genes have six types of repeats (Types 1-6)⁶⁹ in addition to elements (Fig. 7). Individual genes contain between two and four Type 1 repeats, which are located tandemly at the 5' end of the second exon. In contrast, repeat types 2-6 are arranged as mixed, interspersed groups in the 3' half of the second exon. The complexity of these repeats facilitates multiple sequence alignments, two of which have been analyzed in detail.⁶⁶ The “cDNA-based alignment” (Fig. 7A) was generated using the location of gaps in the *Sp185/333* transcript sequences as a guide.⁴⁹ Alternatively, because of the repeats within the sequences, genes (and transcripts) can also be aligned according to the boundaries of the repeats, which results in the “repeat-based alignment” (Fig. 7B). Regardless of the criteria used to align the gene sequences, they are characterized by a similar diversity in element pattern variation as is observed among the transcripts (see Box 2). To date, 171 genes have been isolated from four animals, of which 121 have unique coding sequences and have 33 different element patterns.⁶⁶ Furthermore, identical sequences have not been isolated from more than one individual, suggesting a level of diversity that exceeds the estimated 4-5% sequence divergence between *S. purpuratus* individuals.⁷¹ Large, diverse gene families are a common theme in purple sea urchin immunity (see above) and understanding the mechanisms by which this diversity is generated and maintained is an important aspect of investigations of sea urchin immunity.

Gene-Level Diversification through Recombination

Despite the striking number of unique genes that have been isolated to date, the *Sp185/333* sequences are surprisingly similar. Overall, the genes share >88% pairwise identity. The number of different versions of each element is relatively low.⁷² The complexity of the *Sp185/333* gene family, therefore, is the result of a mosaic pattern of these few element sequences distributed among the genes. Analysis of the evolutionary histories of five of the elements that were present in all of the genes revealed that each element evolved independently. This level of incongruence suggests that the extant *Sp185/333* gene family is the result of rapid and recent diversification events.⁶⁹ It is peculiar, given this high rate of diversification that pseudogenes have yet to be identified.

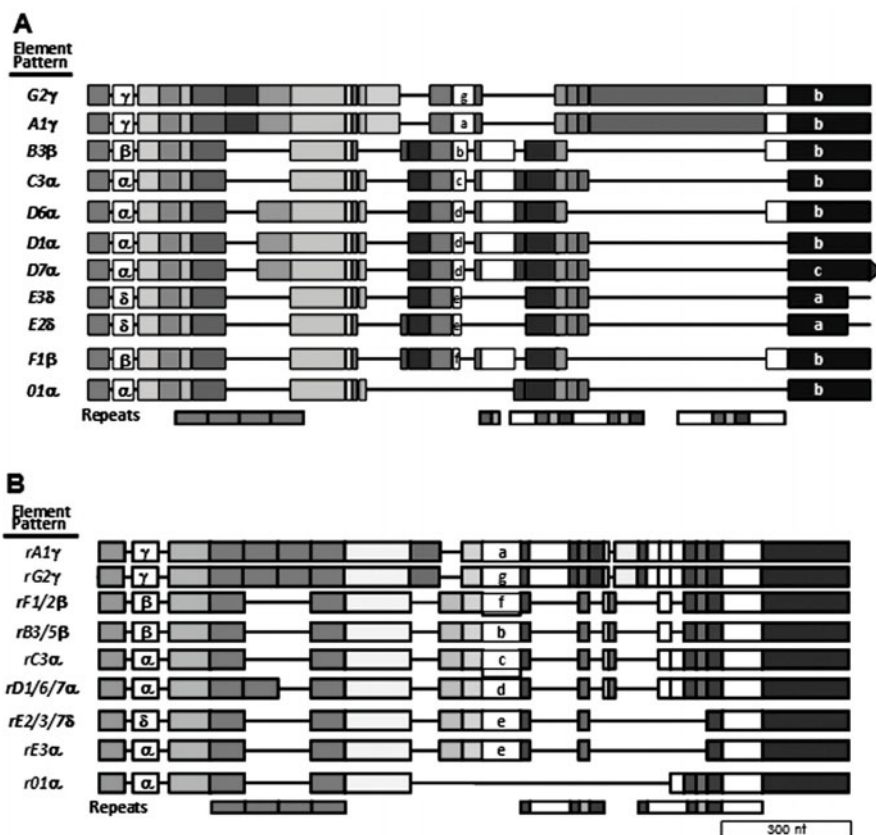


Figure 7. Two optimal alignments of the *Sp185/333* genes. The sequence complexity and repeats within the *Sp185/333* genes facilitate multiple alignments, two of which have been analyzed in detail.⁶⁶ The alignments differ in the pattern of elements in the second exon; the first exon and intron sequence do not have elements. Phylogenetic analysis of the intron sequences define five major types⁶⁶ (designated α - ϵ). The type of intron associated with each element pattern is indicated. A) The “cDNA-based alignment” results from the gaps used to optimize alignments of the *Sp185/333* ESTs and cDNAs.^{49,68} This alignment divides the second exon into 25 elements. The terminal element is differentiated (designated a-c) based on the position of the first of three possible stop codons. The locations of each of the repeat types is indicated by the colored boxes under the alignments. B) An alternative alignment, the “repeat-based alignment”, results from inserting gaps so that the repeats and elements correspond as much as possible.⁶⁶ This alignment divides the second exon into 27 elements based on the locations of gaps, as well as the locations of the repeats. The elements in this alignment that correspond to the repeats are color coded accordingly. The electronic version of this chapter shows the elements and repeats in color. A color version of this image is available at www.landesbioscience.com/curie.

With one exception, all but one of the cloned genes are predicted to encode intact open reading frames.⁶⁶ This may reflect a bias, however, in the primers that were designed in the untranslated regions of expressed genes,⁴⁹ so results do not confirm with certainty to a true lack of pseudogenes within the genome. It should be noted that, extensive computational analysis of elements and regions of the intron suggest that element boundaries do not necessarily serve as “recombination hotspots”, but rather, that recombination appears to occur throughout the length of the genes.

The mechanisms that promote this frequent recombination are unknown. However, there are a few characteristics of the *Sp185/333* gene family that may contribute to its rapid diversification. Amplification of the regions between genes and preliminary analysis of a sequenced BAC insert with six *Sp185/333* genes show that many genes are closely linked (~3 kb). In addition to the six types of repeats found within the coding regions, the genes are flanked on either side by stretches of di- and trinucleotide microsatellites⁶⁶ (Miller, Buckley and Smith, unpublished). These repeats are closely associated with the boundaries of two types of large segmental duplications that include the *Sp185/333* genes (Miller, Buckley, Easley and Smith, unpublished). Microsatellites have been associated with genomic instability and increased recombination frequency^{73,74} and have been implicated in mediating recombination of the variable surface glycoprotein (VSG) genes from *Trypanosoma brucei*⁷⁵ and plant *R* genes.⁷⁶ The genomic organization, high sequence similarity and repeats within and surrounding the genes likely promote the diversification of the *Sp185/333* gene family through frequent recombination and thereby contributing to the complex protein repertoire.

***Sp185/333* Transcript Editing**

Given the diversity of the *Sp185/333* gene family, comparison of the gene and message sequences from individual animals yielded the surprising result that the two sets of sequences are very different.⁷² Specifically, 148 messages and 53 genes were isolated from a single animal, of which only five of the sequences matched identically to another. Similarly, there was little or no overlap in gene and message sequence in two other animals. Furthermore, the gene and message repertoires were generally characterized by different element patterns, such that the predominantly expressed element pattern following immune challenge was *E2*, whereas the most common gene element pattern was *D1*. Although about half of the messages isolated from immunoquiescent animals had a truncated *E2* element pattern, called *E2.1*, the SNP that introduced an early stop codon was never found among the *Sp185/333* genes. In fact, no genes were identified with premature stop codons or indels resulting in frame shifts, which were both common features of *Sp185/333* transcripts.⁶⁷ When genes and message from individual animals were compared, the large majority of messages expressed both before and after immune challenge were the likely product of a few genes.⁷² Conversely, most of the genes that comprise the large *Sp185/333* gene family were not transcribed. Notably, the pattern of nucleotide substitutions between the messages and the genes from which they were most likely transcribed indicated a bias towards transitions, specifically a uridine in the message at a position in which the gene contained a cytidine. This pattern of nucleotide substitution is consistent with cytidine deaminase activity. A number of cytidine deaminase-like molecules have been annotated within the sea urchin genome, but phylogenetic analysis of these sequences fails to identify homologues of activation-induced cytidine deaminase (AID).¹⁰ AID, which has only been identified in vertebrates, is involved in class switch recombination and somatic hypermutation of immunoglobulins in B cells.⁷⁷ Alternatively, it is possible that the observed differences between *Sp185/333* gene and message sequences result from low-fidelity polymerase activity and the sea urchin genome does contain a homologue of terminal deoxytransferase and polymerase μ (Tdt/Pol μ).^{10,78} In higher vertebrates, this enzyme is also involved in immunoglobulin diversification, as well as low-fidelity DNA replication.^{79,80}

Two Levels of Diversity

The *Sp185/333* gene family is an intriguingly diverse facet of the sea urchin immune response.⁶⁶ In response to immune challenge, this gene family is highly expressed and produces a diverse message repertoire.^{49,67,68} The genes are believed to diversify through frequent recombination that does not appear to be limited to element boundaries and may be mediated by repeats within and flanking the coding sequences.⁶⁹ Given the diversity within the gene family, it is surprising that a second tier of diversification appears to affect the message sequences. That the majority of the messages appear to be derived from a few genes suggests that many of the *Sp185/333* genes may be nonexpressed pseudogenes and serve as a source of sequence diversification to the expressed genes. Alternatively, it may be that these genes are simply not expressed under the limited immunological challenges with which the animals have been presented.^{67,68} Thus, the complex *Sp185/333* gene family represents a novel form of invertebrate immunological diversification both at the genomic and, also, possibly, at the posttranscriptional level.

In addition to diversity that appears to be generated by gene recombination, duplication, deletion, conversion,⁶⁹ and mRNA editing,⁷² the array of *Sp185/333* proteins show unexpectedly greater structural complexity than predicted from the genes and messages.⁸¹ Not only are the arrays of *Sp185/333* proteins different among different individual sea urchins, but the majority of sizes are at least twice as large as predicted and up to ≥ 200 kDa (Fig. 8).²⁹ The pI values range from 3-10 although the majority of isoforms have a pI more acidic than predicted. There are up to 260 discrete isoforms in

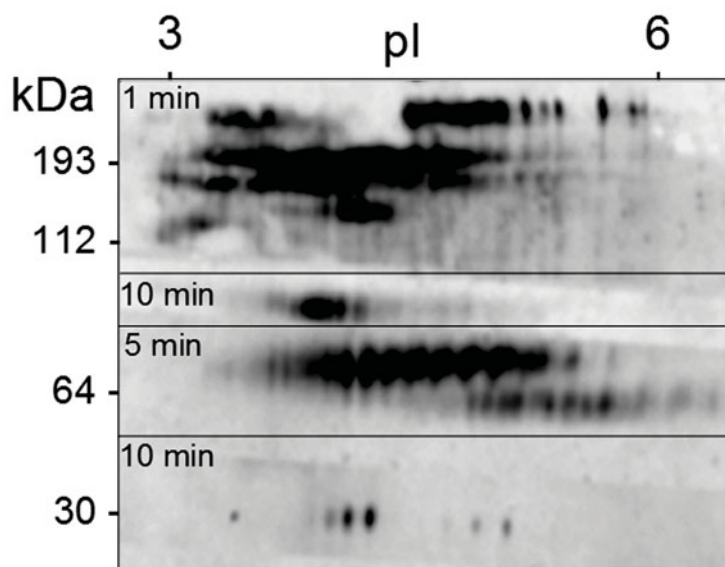


Figure 8. Over 260 spots for *Sp185/333* proteins are present in CF from a single sea urchin. Total proteins from CF were separated by two dimensional gel electrophoresis and analyzed by Western blot with anti-185 antisera. The image is a composite of different regions of the blot that received different exposure times to optimize the spot intensities. pI units are shown at the top and molecular masses (kDa) are shown to the left. Reproduced from reference 81 with permission, ©2009, The American Association of Immunologists, Inc.

individual sea urchin coelomocytes and many of the expressed proteins are truncated, likely a result of mRNA editing.^{67,81} Finally, the arrays of Sp185/333 proteins change in response to different pathogen-associated molecular patterns (PAMPs), which may result from a combination of variations in gene expression, mRNA editing and posttranslational processing of the proteins. How these several levels of diversification are coordinated and perhaps directed towards different types or species of pathogens will be the focus of future research.

COMPLEMENT IN ECHINODERMS

The complement system is a system of central importance in immunity for vertebrates and comprises over 30 known humoral and cell surface proteins.⁸² Complement activation occurs by three major pathways: the classical pathway that is activated by antigen-antibody interaction, the lectin pathway that is activated by mannose binding lectin (MBL) or ficolins and the alternative pathway that is initiated through C3 autohydrolysis (reviewed in ref. 83). C3, a thioester containing protein, is the central component of the cascade and is activated by all three initiation pathways. It acts to coat the surfaces of pathogens, functioning as the initiator of the terminal pathway, an adjuvant for activating the adaptive immune response⁸⁴ and as an opsonin for direct pathogen recognition, phagocytosis and killing.

Investigations of opsonization and phagocytosis by coelomocytes from the green sea urchin *Strongylocentrotus droebachiensis* show that opsonizing target cells with mammalian C3 enhances the coelomocyte response,^{39,85-87} suggesting that coelomocytes have receptors for mammalian C3 and consequently, sea urchins themselves may express C3-like proteins. The identification of two expressed sequence tags (ESTs) from the purple sea urchin, *S. purpuratus* with sequence homology to C3 and to factor B (Bf) called SpC3 and SpBf⁸⁸⁻⁹⁰ was the first evidence that invertebrates have a complement system. Gene models annotated from the sea urchin genome reports several gene models encoding thioester proteins, including a second C3 homologue, Sp-C3-2, several factor B genes, MBL and ficolins (Table 4) (Rast, unpublished).¹⁰

SpC3

SpC3 has several conserved functional regions that are also present in other members of the thioester protein family, including a conserved thioester site, a histidine that regulates binding specificity, a putative C3-convertase site, a two-chain structure putatively disulfide bonded via cysteines in conserved positions, putative factor I cleavage sites and other conserved regions.⁹⁰ Immunoquiescent sea urchins responding to LPS generally show dramatic increases in the amounts of SpC3 in the CF⁹¹ with localization to small transport vesicles in subpopulations of discoidal phagocytes and polygonal phagocytes.²⁸ Expression of *Sp064*, the gene which encodes SpC3, responds to immune challenge in adult animals and also in embryos cultured in the presence of heat-killed bacteria.⁹² The function of SpC3 could be predicted from the deduced amino acid sequence and through comparison to mammalian C3 functions. The characteristics of the thioester, which forms covalent bonds with target molecules that lack protection against complement attack (i.e., microbial surfaces), suggests that SpC3 functions as an opsonin. Classic assays demonstrate that SpC3 binds methylamine, a small nucleophile that interacts covalently with thioester.⁹³ Furthermore, not only does SpC3 undergo autolysis, a reaction that, under appropriate

Table 4. Complement proteins in the sea urchin, *Strongylocentrotus purpuratus*

Gene Model or cDNA	Encoded Protein	Predicted Pathway
<i>Sp-064</i>	SpC3	Alternative
<i>Sp-C3-2</i>	SpC3-2	Alternative?
<i>Sp-TCP1, Sp-TCP2</i>	Thioester containing proteins	?
<i>Sp-thioester containing protein-1, -2, -3/4</i>	Thioester containing proteins	?
<i>Sp-factor B</i>	SpBf	Alternative
<i>Sp-factor B-2, -3</i>	SpBf-2, SpBf-3	Alternative?
<i>SpSM30-F</i>	Mannose-binding protein	Lectin
<i>Sp-C1q-like (4*)</i>	SpC1q	Lectin
<i>Sp-MACPF (21*)</i>	Perforin-like proteins	?
<i>Sp-CD59, Sca2-likel, 2</i>	CD59	Regulatory
<i>Sp5</i>	SpCRL	Regulatory?
<i>Sp5013</i>	SpCRS	Regulatory?

*Numbers of gene models.
From references 10 and 124.

conditions, results in the cleavage of the peptide bond between glutamic acid and glutamine within the thioester (reviewed in ref. 94), but methylamine binding blocks SpC3 autolysis.⁹³ When CF containing SpC3 is incubated with yeast, SpC3 can be detected on the yeast surface and augmented phagocytosis of the yeast by coelomocytes is inhibited by addition of anti-SpC3 antibody.⁹⁵

SpBf

The deduced amino acid sequence and domain structure of SpBf show significant similarity to the vertebrate Bf/C2 family of proteins.⁸⁹ SpBf is a mosaic protein with short consensus repeat (SCR) domains, a Von Willebrand Factor (vWF) domain and a serine protease domain. SpBf has a conserved cleavage site for a putative factor D that is conserved compared to cleavage sites in other Bf/C2 proteins. Members of the Bf/C2 family are mosaic proteins and most have three SCRs, although some have more than three⁹⁶ including SpBf, which has five.⁸⁹ Sequence analysis of these small domains shows that the first two may be the result of a duplication event and that SCR4 may be the result of a recombination between SCR3 and SCR5.⁸⁹ Furthermore, alternative splicing produces some mRNAs with three or four SCRs.⁹⁷ The results suggest redundancies of SCRs in the SpBf protein and that the predicted function may be conserved in SpBf, even though additional SCRs are present, as the relative order of the SCRs in SpBf is maintained.⁸⁹ The gene encoding SpBf, *Sp152*, is expressed in the phagocyte fraction of coelomocytes with low levels detected in ovary, testes, gut and esophagus,⁹⁷ although it is not clear whether expression in tissues is actually due to coelomocytes present in the tissues rather than the tissue cells themselves. Unlike *Sp064* expression, *Sp152* expression is not induced by LPS and appears to be constitutive.

Complement Phylogeny and Evolution

The identification of a C3 homologue in the purple sea urchin⁸⁸ inspired searches for complement components in a wide range of invertebrates. Phylogenetic analysis of thioester proteins including homologues of C3 with α -2-macroglobulin homologues as the outgroup shows that vertebrate C3, C4 and C5 cluster into three well supported clades (Fig. 9). The invertebrate C3 homologues cluster in three paraphyletic clades at the base of the vertebrate complement clades. SpC3 clusters with other invertebrate C3 sequences, while the tunicate sequences form a chordate cluster with the vertebrate sequences (see also ref. 98). The structure of the tree suggests that the thioester complement proteins diverged after the separation of the major phyla. Phylogenetic analyses of members of the Bf/C2 family show that the sea urchin homologue, SpBf, is positioned near the base of the tree⁸⁹ with cnidarian Bf/C2 being more ancient.⁹⁸ A number of analyses of the complement family of proteins have suggested that they evolved from a restricted set of primordial genes^{99,100} (reviewed in refs. 101, 102). Conserved sequence motifs, such as the thioester site (GCGEQ) and similar organization of domains suggest that the ancestral complement system may have included a thioester protein, a Bf/C2 protein and a mannose binding lectin. These are all present in a number of invertebrates including the purple sea urchin (Table 4).

LECTINS

Lectins are a large and heterogeneous group of proteins and glycoproteins present in plants, microorganisms and animals, that function to bind mono- and disaccharides.^{103,104} Lectins can be soluble proteins or integral membrane proteins that often exist as oligomers, contain at least two carbohydrate recognition domains (CRDs) and are capable of agglutinating cells and/or precipitating glycoconjugates.¹⁰⁵ In immunity, lectins are key molecules that function in cell-cell interactions, self/nonself discrimination and interactions between cells and the extracellular matrix (ECM), among other functions.¹⁰⁶ Lectins in invertebrates fall into four major groups based on similarity of structure to vertebrate lectins. C-type lectins are the most common type in various invertebrates¹⁰⁷⁻¹¹⁰ and consist of both soluble and integral membrane proteins that require divalent cations (Ca^{2+}) to maintain the CRD structure to bind carbohydrates. A second major group of invertebrate lectins are S-type lectins that specifically bind β -galactosyl residues,^{111,112} S-type lectins are predominantly intracellular and employ free thiols for binding carbohydrates. A third major group of invertebrate lectins resemble vertebrate pentraxins and share properties with C-reactive protein and serum amyloid protein.¹¹³⁻¹¹⁵ Finally, a fourth group of invertebrate lectins includes all those that cannot be otherwise categorized based on lack of information on their primary structure.

Innate immune functions performed by lectins include recognition and specific binding of microbial surface carbohydrates through the CRDs,¹¹⁶ which exhibit seven different structural folding patterns for binding to different carbohydrate motifs.¹¹⁷ Both mannose-binding lectin (MBL) and ficolins have been identified in the sea urchin genome¹⁰ and a homologue of MBL has been characterized in a sea cucumber, *Apostichopus japonicus*.¹¹⁸ It is speculated that these homologues may initiate the lectin pathway of complement in echinoderms (Table 4). MBLs and ficolins selectively bind mannose,

Figure 9, viewed on previous page. The phylogenetic relationships among members of the thioester protein family. Amino acid sequences for the thioester family were obtained from GenBank. The alignment was done in T-coffee²¹³ and manual editing was done with Mesquite.²¹⁴ A consensus tree was constructed in PAUP*²¹⁵ using maximum parsimony. Bootstrap support was generated with 10,000 iterations. Similar cladogram results were obtained by the maximum parsimony method in PAUP*, Neighbour-joining distance method in PAUP* and the Bayesian method in Mr. Bayes.²¹⁶ The Bayesian method was used with default priors and the GTR+G+I as the nucleotide substitution model suggested by jModelTest.²¹⁷ The α 2macroglobulin sequence from horseshoe crab (*Limulus polyphemus*) was chosen as the outgroup. Sp-C3, *Strongylocentrotus purpuratus*, NP_999686; Hs-C3, *Homo sapiens*, AAR89906; Rn-C3, *Rattus norvegicus*, NP_058690.2; Me-C3, *Macropus eugenii*, AAW69835; Gg-C3, *Gallus gallus*, NP_990736; Xl-C3, *Xenopus laevis*, AAB60608; Lj-C3, *Lethenteron japonicum*, AAR13241; Ej-C3, *Ectosphepus japonicus*, Q00685; Bb-C3, *Branchiostoma belcheri*, BAB47146; Cr-C3, *Carcinoscorpius rotundicauda*, AAQ08323; Se-C3, *Swiftia exserta*, AAN86548; Hr-C3, *Halocynthia roretzi*, BAA75069; Ci-C3, *Ciona intestinalis*, Q8WPD8; Eb-C3, *Eptatretus burgeri*, CAA77677; Om-C3, *Oncorhynchus mykiss*, I51339; Cc-C3, *Cyprinus carpio*, BAA36618; Es-C3, *Euprymna scolopes*, ACF04700; Vd-C3, *Venerupis decussates*, ACN37845; Gg-C4, *Gallus gallus*, T28153; Xl-C4, *Xenopus laevis*, BAA11188, Hs-C4, *Homo sapiens*, AAB67980; Cc-C4, *Cyprinus carpio*, BAB03284; Mm-C4, *Mus musculus*, CAA28936; Ts-C4, *Triakis scyllium*, BAC82347; Hs-C5, *Homo sapiens*, AAI13739; Mm-C5, *Mus musculus*, P06684; Gg-C5, *Gallus gallus*, XP_415405; Om-C5, *Oncorhynchus mykiss*, AAK82852; Lp-A2M, *Limulus polyphemus*, BAA19844; Lj-A2M, *Lethenteron japonicum*, BAA02762; Pt-A2M, *Pan troglodytes*, XP_001139559; Xl-A2M, *Xenopus laevis*, AAY98517; Cc-A2M, *Cyprinus carpio*, BAA85038; Cf- α 2M, *Chlamydia farreri*, AAR39412; Ci-A2M, *Ciona intestinalis*, NP_001027688; Hs- α 2M, *Homo sapiens*, P01023.

fucose and other specific carbohydrates present exclusively on the surface of pathogens resulting in the direct activation of complement.¹¹⁹⁻¹²²

Lectins were first demonstrated in echinoderms in the early 1980s.¹²³ Lectins have been identified in asteroids, echinoids and holothurians and characterized essentially from a functional and biochemical point of view. With improved molecular methods greater numbers of lectins and proteins with putative lectin function have been identified. Preliminary analysis of the sea urchin genome shows gene models encoding more than 100 small C-type lectins, over 400 mosaic proteins with lectin domains, 34 galectins, in addition to a few pentraxins and fucolectins¹²⁴ (Cohen and Smith, unpublished). This indicates that lectins in general are likely to have a variety of important functions in echinoderms, including recognition of foreign cells. Many, but not all of the lectins that have been identified in echinoderms have been C-type lectins (Table 5). Many show opsonin and agglutinin functions with the capability of binding carbohydrates on the surface of pathogens. It is generally accepted that lectins in echinoderms play an important role in the immune system, functioning as key molecules in immune responsiveness and to augment coelomocyte functions in host defense.

SCAVENGER RECEPTORS

Another expanded gene family in the sea urchin genome encodes a large repertoire of scavenger receptors containing multiple scavenger-receptor cysteine-rich (SRCR) domains that are both membrane-bound and secreted proteins.^{125,126} Receptors of this structure are found throughout the animal kingdom but the sea urchin genome encodes more than 1000 SRCR domains in ~180 gene models, which greatly exceeds the multiplicity of these genes in other characterized species (Fig. 6).¹⁰ These proteins are known to act as phagocytic receptors and some family members in mammals have been shown to bind bacteria.¹²⁷ These receptors are highly polymorphic in the population of purple sea urchins and show

Table 5. Echinoderm lectins

Class	Species	Name	Structure	Specificity	Agglutination	Reference
Asteroidea	<i>Asterina pectinifera</i>	(-)	C-type lectin	α -N-acetyl-galactosamine	yes	224
	<i>Oreaster reticulatus</i>	(-)	C-type lectin	galactosyl	(-)	225
Echinoidea	<i>Strongylocentrotus purpuratus</i>	SpEchinoidin	C-type lectin	galactose and derivatives	(-)	88,124
	<i>Anthocidaris crassispina</i>	Echinoidin SUEL	C-type lectin unique	N-acetyl-galactosamine D-galactoside	yes (-)	108 123
	<i>Lytechinus variegatus</i>	Echinonectin	(-)	galactoside	(-)	226
	<i>Paracentrotus lividus</i>	(-) P/SL	C-type lectin (-)	(-) D-glucose, L-rhamnose D-arabinose, L-fucose N-acetyl-D-glucosamine	yes yes	146 227
	<i>Toxopneustes pileolus</i>	SUL-I SUL-II	(-) (-)	D-galactose D-fucose D-galactose	(-) (-)	228 228
	<i>Tripneustes gratilla</i>	TGL-I	C-type lectin	(-)	(-)	228

continued on next page

Table 5. Continued

Class	Species	Name	Structure	Specificity	Agglutination	Reference
Holothuroidea	<i>Apostichopus japonicus</i>	SJL-I	(-)	N-acetyl-D-galactosamine	yes	229
		SJL-II	(-)	simple carbohydrates	yes	229
		SPL-I	C-type lectin	D-glucuronic acid, D-galacturonic acid	(-)	230
		SPL-II	C-type lectin	D-galactosamine, D-galactose	(-)	230
		MLB-AJ	C-type lectin	α -D-mannans	yes	118
<i>Cucumaria echinata</i>		CEL-I	C-type lectin	N-acetyl-galactosamine	yes	231
		CEL-II	C-type lectin	(-)	no	232
		CEL-III	C-type lectin	(-)	yes	231
		CEL-IV	C-type lectin	N-acetyl-galactosamine, α -galactose	yes	232
<i>Cucumaria japonica</i>		(-)	(-)	branched α -D-mannans	yes	233
	<i>Holothuria scabra</i>	HSL	(-)	galactose derivatives, T-antigen	yes	234

(-), not known, not done, not named.

pronounced variability in coelomocyte expression among individual animals suggesting a complex expression control system.¹²⁵ It is notable that representatives of all three of the most expanded families of sea urchin receptors (TLR, NLR and SRCR receptors) form a coregulated immune circuit that, in mammals, functions in gut immunity.¹²⁸

ANTIMICROBIAL PEPTIDES

Antimicrobial peptides (AMPs) have been identified in a wide variety of species including bacteria, fungi, plants, insects, tunicates, amphibians, birds, fish and mammals.¹²⁹⁻¹³² Since the discovery of cecropins in insects¹³³ and defensins in mammals,¹³⁴ more than 1,200 different eukaryotic AMPs have been characterized.¹³⁵ In eukaryotes they form the first line of host defense against pathogenic infections and are a key component of the innate immune system. AMPs have an enormous variety of sequences and structures, but certain features are common. Most have a net positive charge and are 12-50 amino acids long, of which approximately half are hydrophobic.^{136,137} However, a few peptides of up to 100 residues are also recognized as AMPs.¹³⁸

Early work to document antimicrobial activities of crude extracts from echinoderms showed a wide range of activities against bacterial and fungal isolates^{45,139} of which some functioned as antifoulants to deter the settlement of barnacle and bryozoan larvae in addition to bacterial colonization.¹⁴⁰ More recently, a variety of molecules with antimicrobial properties have been isolated from echinoderms, including steroidal glycosides,¹⁴¹⁻¹⁴³ polyhydroxylated sterols,¹⁴¹ naphthoquinone pigments such as echinochrome A,^{33,144} and complement homologues.^{89,90} Lysozymes with antibacterial activity have also been detected,¹⁴⁵⁻¹⁴⁷ and concentrations up to 15 µg/ml have been found in red spherule cells of the sea urchin *Paracentrotus lividus*.³⁴ Lysozyme and other antibacterial factors may act synergistically to provide effective defense against bacterial infections. In the Strongylocentrotids, the antimicrobial pigment echinochrome A is present in vesicles of red spherule cells and is bound to uncharacterized coelomocyte proteins.¹⁴⁸ A semi-purified coelomocyte fraction from *Paracentrotus lividus*, included fragments of beta-thymosin that were proposed to have antibacterial activity.¹⁴⁹ In extracts from coelomocytes of the sea star *Asterias rubens*, a number of partial peptide sequences were obtained and identified as fragments of actin, histone H2A and filamin A.^{150,151} Antibacterial activity was detected in extracts of several tissues from the green sea urchin *Strongylocentrotus droebachiensis*, the common sea star *Asterias rubens*, and the sea cucumber *Cucumaria frondosa*,¹⁵² with most activity in coelomocytes and body wall.

Strongylocins

Scans of the gene models in the purple sea urchin genome did not detect any sequences encoding recognizable AMPs.¹⁰ This may be due to short exons that are difficult to recognize computationally, but may also be due to the divergent nature of these small proteins. However, two cysteine-rich AMPs, called strongylocins, have recently been isolated and characterized from the green sea urchin, *S. droebachiensis*.¹⁵³ Homologues are also present in the sister species *S. purpuratus*, called SpStrongylocins.¹⁵⁴ The strongylocin peptides are members of the cysteine-rich AMP family, which have six cysteines with three disulfide bonds involved in peptide conformation, stabilization

and resistance to proteases,¹⁵⁵ and which are crucial for the antimicrobial activity.^{156,157} The strongylocins have a novel cysteine pattern (Table 6) suggesting a different conformation than the other members of the group, perhaps to resist proteolysis within the coelomocytes and in the CF. Both the native and recombinant peptides show antibacterial activity against both Gram-positive and Gram-negative bacteria (Table 7). In addition, other peptides have been isolated and characterized from *S. droebachiensis* that appear as heterodimers and have strong activity against Gram-positive and Gram-negative bacteria (Stensvåg, unpublished). The heavy chain from one of the heterodimeric peptides also has strong activity against fungi and yeast.

The strongylocins are composed of three regions: a signal peptide, a prosequence and the mature peptide, and the strongylocin 1 peptides show high amino acid sequence similarity throughout (Fig. 10). However, SpStrongylocin 2 shares an identical signal peptide with strongylocin 1, instead of strongylocin 2. The prosequences are negatively charged, which may act to neutralize and stabilize the positive charge of the mature peptide^{153,154} and presumably function as an intracellular steric chaperone during folding.¹⁵⁸⁻¹⁶⁰ The peptides become active after the prosequences are cleaved during maturation. The first amino acid in the mature peptide of strongylocin 2 from *S. droebachiensis* is a tryptophan which is likely brominated. Although the recombinant SpStrongylocin 2 is not brominated, it shows equivalent antimicrobial activity to native peptides. Therefore, the posttranslational modification of tryptophan may affect the properties of the peptides by enhancing stability rather than mediating antimicrobial activity. The site of strongylocin activity is likely to be intracellular based on membrane integrity assays.¹⁵⁴

Nonechinoids also synthesize a range of AMPs, of which many have been characterized. For example, the antibacterial activity in the CF of the orange-footed sea cucumber, *Cucumaria frondosa*, has been traced to small peptides (≤ 6 kDa) that appear to be active at low pH (5.0-6.5) and which may be similar to the clavanins found in solitary tunicates.¹⁶¹ Other immune-active chemical compounds with roles in maintaining antiseptic environments in nonechinoids include saponins and saponin-like compounds in sea stars and brittle stars, which are active against some Gram-positive bacteria.¹⁴¹ Sea stars and brittle stars in particular express steroidal glycosides that exhibit antifungal activity and toxicity against brine shrimp.¹⁶²⁻¹⁶⁴ It is likely that multitudes of molecules

Table 6. Cysteine patterns in AMPs containing six cysteines

Peptide Family	Cysteine Pattern ^a	Group of Organism
Strongylocins	C - C - C - CC - C	Echinoderms
Beta-defensins	C - C - C - C - CC	Mammals, birds
Alpha-defensins	C - C - C - C - CC	Mammals
Tachystatins	C - C - CC - C - C	Horseshoe crab
Knottin-type AMPs	C - C - CC - C - C	Plants
Thionins Type III and IV AMPs	CC - C - C - C - C	Plants
Insect defensins	C - C - C - C - C - C	Insects
Mytilus defensin	C - C - C - C - C - C	Molluscs

^aAdjacent double cysteine residues are highlighted. Information regarding cysteine arrangements in the different peptides was obtained from the *Antimicrobial Peptide Database*.¹³⁵

Table 7. Susceptibility of bacteria to strongylocins, recombinant SpStrongylocins and dimeric peptides from *Strongylocentrotus droebachiensis* and *S. purpuratus*

Peptide	Minimal Inhibitory Concentration (μ M)						
	<i>Listonella anguillarum</i>	<i>Escherichia coli</i>	<i>Corynebacterium glutamicum</i>	<i>Staphylococcus aureus</i>	<i>Penicillium roqueforti</i>	<i>Candida albicans</i>	
Strongylocin 1 ^a	2.5	5.0	2.5	2.5	Nt ^c	Nt	
Strongylocin 2 ^a	1.3	5.0	2.5	2.5	Nt	Nt	
Recombinant SpStrongylocin 1 ^b	15.0	7.5	7.5	15.0	Nt	Nt	
Recombinant SpStrongylocin 2 ^b	15.0	7.5	3.8	15.0	Nt	Nt	
Dimeric peptide 1 ^a	2.5	1.3	1.3	2.5	6.25 ^d	6.25 ^d	
Dimeric peptide 2 ^a	2.5	2.5	1.3	5.0	Nt	Nt	

^aMinimal inhibitory concentration was determined as the lowest concentration of peptide causing an optical density less than 50% of the growth control without any peptide present.

^bMinimal inhibitory concentration was determined as the lowest concentration of peptide causing 100% growth inhibition of the test organism compared to the growth control.

^cNt = Not tested.

^dThe peptide is the heavy chain of dimeric peptide 1. Growth inhibition was determined microscopically after 48 hrs of incubation.

with activity against all types of pathogens function efficiently in echinoderms and have central importance for immune functions in this group.

GENE EXPRESSION IN SEA URCHIN COELOMOCYTES

Before it was possible to scan for gene models encoding proteins with putative immune function in the sea urchin genome, the only feasible approach for understanding the echinoderm molecular immunology was through EST profiling of messages in coelomocytes under challenged vs nonchallenged conditions.^{49,88,165} ESTs encoding proteins with putative immune function have also been reported for the sea cucumber *Holothuria glaberrima*, undergoing gut regeneration after evisceration.^{52,53} A variety of categories of proteins are expressed in coelomocytes that illustrate the activities of these cells in an assortment of functions (Table 2). The main focus of these studies was to identify immune response genes (see Box 2), which included protein matches suggesting several mechanisms for opsonization, clotting, iron sequestration plus the activity of a variety of signaling pathways. Matches were found to proteins with functions in the endosomal system (lysosomes) that may be involved in killing phagocytosed pathogens. Coelomocytes express a number of genes encoding proteins involved in splicing transcripts and translating proteins. They process, package, transport and secrete proteins as suggested from the transcripts encoding proteins involved in trafficking transport vesicles. Phagocytes have extensive cytoskeletons (as illustrated in Figs. 1-5), which is borne out by the number of ESTs matching proteins that function as cytoskeletal elements and others that act to modulate the cytoskeleton. One of the benefits of generating

Box 2. EST studies identify immune-related transcripts; discovery of the *Sp185/333* sequences.

Differential characterization of ESTs of coelomocytes from *Strongylocentrotus purpuratus* before and after immune challenge provide a picture of global changes in transcriptional activity. Genes encoding complement homologues,⁸⁸⁻⁹⁰ transcription factors,¹²⁶ and a lectin,^{67,88} among many others are induced by immune challenge (Table 6). Significantly upregulated transcripts include a large and diverse set of novel transcripts designated *Sp185/333* that were first discovered with the use of differential display¹⁶⁵ and subtracted probes followed by EST analysis.⁴⁹ Probes representing transcripts from LPS-activated coelomocytes were used to screen a high-density arrayed, conventional cDNA library made from bacterially-activated coelomocytes. About 4.5% of the clones in the library were positive and about 60% of clones selected for EST analysis showed significant sequence similarity to two previously uncharacterized cDNAs: *DD185* and *EST333*.^{70,88,126} Given the considerable number of related *Sp185/333* sequences, the coding regions of the ESTs could be aligned. However, optimization of the alignment required the insertion of multiple, large artificial gaps⁴⁹ that defined 25 to 27 blocks of shared sequence called *elements*, which are variably present or absent in *element patterns* (Fig. 7).^{66,67} The extraordinary diversity of the *Sp185/333* transcripts is based on the mosaic element patterns in addition to extensive single nucleotide polymorphisms (SNPs). The induction of *Sp185/333* gene expression in response to LPS and other pathogen-associated molecular patterns (PAMPs)^{67,165} plus the diversity of the sequences suggest immune-related functions for the encoded proteins in *S.purpuratus*.

ESTs is that the level of expression rather than the level of match to known sequences can sometimes provide clues to putative immune function for unknown sequences. This has been the case for the *Sp185/333* sequences that are not known outside of the echinoids, but for which there is strong evidence of immune function.⁷⁰

PROTEIN EXPRESSION PROFILE OF SEA URCHIN COELOMOCYTES

In addition to genome scans and EST analyses, high throughput methods in proteomics are being applied to analyzing proteins in echinoderms. In response to LPS challenge, 319 proteins were identified in the wCF of the purple sea urchin, *Strongylocentrotus purpuratus*, of which 284 were encoded by gene models in the genome and 48 were encoded by hypothetical open reading frames (Table 3) (Dheilly, Raftos and Nair, unpublished). The majority of the proteins were involved in modulating the cytoskeleton and linkage between the cytoskeleton and cell adhesion molecules, all of which are essential for intracellular transport and behaviors such as locomotion and phagocytosis. Cell adhesion molecules and intracellular signaling proteins were also identified, suggesting that coelomocytes respond to LPS with large-scale alterations to the cytoskeleton. Furthermore, the response to immune challenge implies the secretion of cellular products, as well as the endocytosis of extracellular fluids and phagocytosis of pathogens. Proteins involved in clotting and coagulation suggest another mechanism to sequester pathogens and to clear them from the CF, in addition to the prevention of CF loss after trauma. A number of proteins involved in opsonization are present in wCF including complement homologues. Metal-binding proteins are present, including ferritin, which has been identified in several echinoderm species under various conditions and using different analytical approaches.^{49,52,88,166} Consequently, sequestration of iron must be an important mechanism for controlling the proliferation of invading microbes.

Proteomic analyses of coelomocytes from a second sea urchin species, *Heliocidaris erythrogramma*, demonstrated different expression profiles over time in response to sterile injections compared to injection of microbes (Dheilly, Raftos and Nair, unpublished). Results from shotgun proteomics showed two different profiles of proteomic changes. The proteins identified in the response to bacterial challenge were different from those identified in the controls. The proteomic profile changed in the coelomocytes from bacterially challenged animals within the first 6 hrs, largely due to the increased abundance of some proteins involved in cytoskeletal dynamics, while others decreased such as F-actin capping protein, advillin and α -actinin. Other cytoskeletal proteins that were absent in the controls, such as actins and annexin A7, were present in the challenged coelomocytes. Other proteins that appeared in the challenged coelomocytes included the complement homologue, C3 and SRCRs. The protein expression profile in sea urchin coelomocytes indicates that these cells show dynamic responses to wounding and immune challenge. Temporal analysis of proteomic changes in coelomocytes indicates that cellular responses to wounding and infection are biphasic. The initial phase, occurring within the first 24 hrs after treatment, appears to be a generalized response common to both types of insult. This phase involves reactions including CF coagulation and coelomocyte cytoskeletal remodelling, which resolves by 48 hrs to control levels. The second phase, which peaks at 48 hrs after injection, appears to be specific to microbial infections and the proteins expressed in this phase

are involved in pathogen recognition and opsonization as well as the destruction of the invading microbes. A number of attendant sub-cellular pathways involved in signal transduction, endocytosis and exocytosis are also enhanced during this phase. These two phases thereby function in an integrated manner to repair wounds and to neutralize microbial infections.

REGENERATION

Many classes of echinoderms are remarkably plastic in their abilities for repair and regeneration resulting from both proliferation and transdifferentiation of circulating cells of mesenchymal origin (see *Microscopy Research and Technique* vol 55 no. 6, 2001). New evidence suggests that cells required for regeneration originate from coelomocytes.¹⁶⁷⁻¹⁶⁹ Unfortunately, sea urchins have limited regenerative capabilities¹⁷⁰ compared to sea stars and crinoids, which have received more attention.¹⁶⁸ Sea stars are generally known for arm regeneration and levels of heat shock protein 70 (Hsp70) increase in the common sea star *Asterias rubens*, following arm tip amputation.¹⁶⁹ Furthermore, manganese treatment induces proliferation of coelomic epithelial cells that is coupled to traumatic stress responses including increased expression of Hsp70.¹⁷¹ Red spherule cells (or amoebocytes) and polygonal phagocytes appear to be involved in regeneration.¹⁶⁹ These findings suggest a role for increased coelomocyte numbers and the expression of classic stress molecules in the early repair phase of tissue damage and regeneration.

Toposome, which is also referred to as the major yolk protein (MYP), is the most abundant protein in the CF (Table 3).¹⁷² A monoclonal antibody to embryonic toposome also recognizes sea star coelomocytes and the coelomic epithelium.^{169,173,174} Although previous efforts in cloning the MYP gene in *Strongylocentrotus purpuratus* suggested its relationship to vitellogenin,¹⁷⁵ protein fragments were too short for unambiguous identification. The cDNA sequences encoding MYP from the sea urchins, *Pseudocentrotus depressus*, *Paracentrotus lividus* and *Tripneustus gratilla*, show that they are members of the transferrin family, lack iron-binding sites, and are not homologous to vertebrate vitellogenins.^{173,176} The toposome precursor, which has been postulated to serve multiple functions, is synthesized exclusively in the gut of the adult animal as a 180-190 kDa glycoprotein and the mature protein is found in the CF. The amount of toposome protein increases in response to traumatic stresses in agreement with increased gene expression as deduced from ESTs matching toposome in sea cucumbers regenerating gut tissue.¹⁷⁷ Toposome is expressed with other genes encoding proteins that function in wound healing, cell proliferation, differentiation, morphological plasticity, cell survival, stress response, immune challenge and neoplastic transformation.⁵⁵ It is emerging that circulating coelomocytes from some echinoderms originate from coelomic epithelia and are able to differentiate into a few tissue types, including nerve and muscle cells.¹⁷⁸ Accordingly, research on stem cells in marine organisms is becoming important for both comparative studies and for future applications.

ECOTOXICOLOGY AND THE ECHINODERM IMMUNE SYSTEM

Because coelomocytes are sensitive, stress-activated effectors of the echinoderm immune response,^{19,179} they are good candidate biosensors for monitoring environmental stress in an environmental management context. Several examples of using coelomocytes as indicators employ both analyses of specific proteins and of cell function. Unfractionated coelomocytes from the sea urchin *Paracentrotus lividus* cultured at stress-inducing temperatures, (4°C or 35°C), or exposed to acidic pH, or high levels of cadmium, express high levels of Hsp70,^{180,181} a well-recognized stress marker induced in response to a wide range of biological and physicochemical stresses.^{106,182} In addition to serving as molecular chaperones, secreted and membrane-bound heat shock proteins (particularly Hsp60, Hsp70, Hsp90 and gp96) are potent activators of the innate immune system capable of inducing the production of proinflammatory cytokines by the monocyte-macrophage system.¹⁸³ Sp-gp96, which is expressed in sea urchin coelomocytes⁸⁸ is present on the surface of coelomocytes.¹⁸⁴ High levels of Hsp70 are also observed in coelomocytes obtained from sea urchins collected from waters heavily polluted with urban run-off and industrial waste¹⁸¹ and, more recently, with the explosive 2,4,6-trinitrotoluene (TNT) from conventional weapons dumped at sea at the end of World War II.¹⁸⁵ Increased levels of Hsp70 levels in coelomocytes from specimens of the sea star *Asterias rubens*, collected along a transect from inland waters to the open sea along the Norwegian fiords correlate with a natural concentration gradient of heavy metals.¹⁸⁶ Increased production of reactive oxygen species (ROS) by coelomocytes from *A. rubens* is also observed after cadmium exposure in a dose-dependent manner.¹⁸⁷⁻¹⁸⁹ Cadmium, on the other hand, leads to reduced phagocytic activity by coelomocytes.¹⁹⁰ Other heavy metals, including cadmium, as well as UV light can also result in single strand DNA breaks in coelomocytes.^{20,191,192} Exposure to lead increases the phagocytic activity of sea star coelomocytes, though another pollutant, polychlorinated biphenyls (PCBs), do not have this effect.¹⁹⁰ Responses to temperature stress levels can also be detected in coelomocytes as increases in acetylcholine esterase (AChE) activity.¹⁹³ Overall, there are several approaches for monitoring protein levels and enzyme activities that can be employed as signs of environmental stress both in field and in laboratory studies.

Another approach for employing echinoderms as biosensors has been to evaluate the numbers of coelomocytes in the CF, which can be altered by environmental stressors. For example, the percentage of red spherule cells increase from 5% to 40% of total coelomocytes in animals collected from polluted seawaters or that are subjected to accidental injury.^{181,185} Similarly, the numbers of total coelomocytes increase in immunoquiescent sea urchins after simple immune challenge in the lab.²⁹ Therefore, levels of red spherule cells and perhaps total coelomocytes may be used as a practical marker of environmental stress in animals collected in costal surveys for marine management.

CONCLUSION

Echinoderms are important members of marine ecosystems and are required for the stable maintenance of habitats. This has been illustrated by the aftermath of the disappearance of the long-spined black sea urchin, *Diadema antillarum*, the top herbivore

on Caribbean coral reefs. The population crash in 1983-1984, with repeat crashes in 1985 and 1991-1992, reduced the population in the Caribbean and Western Pacific by 95-99%.^{194,195,196} General destabilization of the coral reef ecology resulting from herbivore release that was exacerbated by hurricane damage, lead to a change in the trophic cascade resulting in a swift and sustained switch to an alternative ecological state; reef cover of mostly coral switched to mostly soft algae.¹⁹⁷⁻²⁰⁰ The effect of the disappearance of *D. antillarum* was predicted by Sammarco²⁰¹ and modeling these population changes show the same outcome.²⁰²

The population crash of *D. antillarum* progressed from west to east in the Caribbean and was speculated to have been due to a pathogen¹⁹⁸ that may also have been a commensal.²⁰³ Disease outbreaks and mass mortalities have also been noted in the green sea urchin, *S. droebachiensis*, along the Atlantic coast of Nova Scotia.²⁰⁴ Little is known of the bacterial pathogens of sea urchins²⁰⁵⁻²⁰⁹ and much less is known of the fungal and viral pathogens of any class of echinoderm. This general lack of knowledge regarding pathogenesis in echinoderms is an important problem that will require coordinated efforts of both ecologists and immunologists. Understanding the population dynamics in a complex ecosystem such as coral reefs should include investigations of the host-pathogen interactions that are involved in the stability of the ecosystem or lack thereof.

Until recently, workers interested in the immune system of echinoderms were hobbled by identifying and analyzing one gene or protein at a time. Cross-phylum searches for genes encoding proteins involved in microbial recognition and immune effector functions were complicated by the rapid pace of immune gene evolution. However, the availability of the purple sea urchin genome¹¹ has greatly increased the sensitivity with which immune gene homologues can be identified, in addition to the efficient characterization of complex multigene families. Immune transcription regulators are generally very well conserved and BLAST type sequence identity searches are typically adequate to identify these factors. In the case of many immune receptors and effectors, primary sequence is poorly conserved but domain structure can be used as a unique identifier and combinatorial domain profile searches can be useful. A final class of immune mediators is encoded by genes that have novel structure relative to other phyla. These are typically identified in experimental surveys, but in some cases multiplicity of domains that are common to proteins with immune functions can be identified in purely bioinformatic surveys for candidates of novel immune mediators.

Analysis of the purple sea urchin genome has revealed a complex repertoire of immune receptors, regulators and effectors unlike those known in other phyla.^{10,60,78,210} In addition to the LRR-containing proteins and the *Sp185/333* gene family described above, the genome encodes a virtually complete set of homologues of (i) vertebrate hematopoietic and immune transcription factors, (ii) candidate effector proteins and (iii) genes with distant homology to key adaptive immune mediators of the jawed vertebrates.¹⁰ Transcription regulators of haematopoiesis include a nearly complete set with respect to the vertebrate homologues and include representatives of some subfamilies such as the PU.1/SpiB/SpiC Ets factors that are important regulators of vertebrate myeloid and lymphoid immunocyte development not found outside of deuterostomes.²¹¹ Transcription factors involved in the regulation immune response genes include GATA1/2/3, SCL and NFkB have also been identified.

Future work on echinoderm immunity will focus on the purple sea urchin, but additional echinoderm genomes are needed. Preliminary 1X sequence coverage of the genomes for

S. franciscanus and *Allocentrotus fragilis*, sister species of *S. purpuratus* are available ([http://www.hgsc.bcm.tmc.edu/project-species-o-Strongylocentrotus%20purpuratus.hgsc?pageLocation = Strongylocentrotus purpuratus](http://www.hgsc.bcm.tmc.edu/project-species-o-Strongylocentrotus%20purpuratus.hgsc?pageLocation=Strongylocentrotus%20purpuratus)) and preliminary analyses suggest a similar repertoire of immune genes. Additional genomes from species from other classes are needed and will not only promote phylogenetic and comparative evolutionary genomics, but will enable the characterization of the immune gene repertoire for a different echinoderm species. This will be of particular interest given that the immune genes and the mechanisms of immune gene diversification are dictated by the life history, pathogens and habitat of individual species.

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UROCHORDATE IMMUNITY

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Abstract: This chapter provides a short review of the immune system of urochordates, the closest living relative of vertebrates. Since adaptive immunity is a unique property of vertebrates, urochordates rely exclusively on innate immunity to recognize and eliminate pathogens. Here we discuss three immune systems of urochordates which show different evolutionary relationship with the vertebrate immune system. Urochordate Toll-like receptors (TLR) show a clear orthologous relationship with vertebrate counterparts, although they show unique characteristics most likely gained in the urochordate lineage. The urochordate complement system also shows orthologous relationship with the vertebrate complement system. From the structural and functional viewpoints, it seems to represent a more primitive state of the vertebrate complement system without any major deviation. In contrast, the allorecognition systems of urochordates show no evolutionary relationship with any invertebrate or vertebrate systems, suggesting that they were invented in the urochordate lineage.

INTRODUCTION

Vertebrates developed two types of the adaptive immune system, a well-defined one based on conventional lymphocytes and the major histocompatibility complex (MHC) of jawed vertebrates and an emerging one based on novel lymphocytes of jawless fish.¹ No evidence for the presence of adaptive immunity has been reported from invertebrates thus far, whereas various types of innate immunity, some common with vertebrates and others specific to certain phylogenetic groups, have been reported. To understand the origin and evolution of vertebrate immune system, it is essential to analyze the immune system of the closest relatives of vertebrates. Vertebrates, urochordates and cephalochordates constitute the phylum Chordata. Traditionally, cephalochordates are considered as the closest living

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relatives of vertebrates, with urochordates representing the earliest chordate lineage. This view is mainly justified by overall morphological similarities and an apparently increased complexity in cephalochordates and vertebrates relative to urochordates. However, recent molecular phylogenetic analyses provided compelling evidence that urochordates and not cephalochordates, represent the closest living relatives of vertebrates.² Comprehensive search for immune-related genes from draft genome information of one urochordate species, *Ciona intestinalis*, indicated that the pivotal genes for adaptive immunity, such as the MHC class I and II genes, T-cell receptors, or dimeric immunoglobulin molecules, are missing from the *Ciona* genome.³ In contrast many genes possibly involved in innate immunity are identified. Among them, here we discuss three well-characterized systems, TLR (Toll-like receptor), complement system and allorecognition. The former two are evolutionary related to their vertebrate counterparts, whereas the last one is unique to urochordate lacking the vertebrate counterpart.

TOLL-LIKE RECEPTORS (TLR)

What Are Toll-Like Receptors?

Toll-like receptors (TLRs) play pivotal roles in host defenses via the innate immune system. All TLRs are Type I transmembrane proteins which harbor an intracellular Toll/Interleukin-1 receptor (TIR) domain and extracellular leucine rich repeat (LRR) motifs.^{4,5} LRRs exhibit specific pathogenic ligand recognition and TIR participates in the activation of downstream signaling pathways. Nine functional human TLRs (hTLRs) have been identified. As summarized in Table 1, each hTLR directly recognizes their specific ligands (or pathogen-associated molecular patterns, PAMPs). Molecular diversity in the number and organization of LRRs enables the specific and sensitive recognition of PAMPs by respective TLRs. TLRs are expressed not only in immune cells such as lymphocytes, macrophages and dendritic cells but also nonimmune tissues including lung, small intestine, stomach and testis. TLR1, TLR2, TLR4, TLR5 and TLR6 recognize extracellular microbial pathogenic components on plasma membranes, whereas TLR3, TLR7, TLR8 and TLR9 respond to viral DNA or RNA on endosomes. In addition, TLR4 requires an extracellular associated protein, MD2, to recognize lipopolysaccharide (LPS).^{4,6} Interaction of TLRs with specific PAMPs triggers signal transduction pathways via adaptor proteins (MyD88, TIRAP, TRIF and TRAM) followed by activation of a wide range of inducible transcriptional factors such as NF- κ B, AP-1 and IRF, leading to production of an inflammatory cytokine TNF α , chemokines and/or Type I interferon.^{4,5} TLRs or their related genes have also been detected in fish,⁷⁻⁹ cyclostomes,¹⁰ amphioxus,¹¹ sea urchin,¹² annelid¹³ and cnidarian,¹⁴ although their functions, except for several fish TLRs, have yet to be elucidated.

TLRs of *C. intestinalis*

A *Ciona* genome survey and molecular cloning revealed the presence of two TLRs in *C. intestinalis*, namely, Ci-TLR1 and -2.^{3,15} Ci-TLR1 and Ci-TLR2 are composed of a TIR, transmembrane and LRR domain, which is typical of TLRs. Moreover, 7 and 13 LRRs are found in Ci-TLR1 and Ci-TLR2, respectively. Ci-TLR1 and -2 were most homologous to hTLR7 (26%) and hTLR8 (26%), respectively.¹⁵ However, the sequence homology is inconsistent with PAMP recognition and intracellular localization of Ci-TLRs (Table 1).

Table 1. Ligands of human and *Ciona* TLRs

TLR	Ligands	Intracellular Localization
TLR1/2	triacylated lipoprotein	PM
TLR2	Zymosan (yeast cell wall) 1, 3- β -glucan Lipoarabinomannan Heat-killed <i>Legionella pneumophila</i> (HKLP, Gram-negative) Heat-killed <i>Staphylococcus aureus</i> (HKSA, Gram-positive) Glycosylphosphatidylinositol (GPI)-anchored glycoprotein	PM
TLR3	poly(I:C) (double-stranded RNA)	ES
TLR4 (with MD2)	LPS (lipopolysaccharide from Gram-negative bacteria) Lipid A (lipid component of LPS)	PM
TLR5	Flagellin (bacterial flagellar filament)	PM
TLR6	MALP-2 (mycoplasma-derived macrophage-activating lipopeptide) FSL1 (micoplasma-derived lipoprotein)	PM
TLR7	Imidazoquimod (imidazoquinolone amino acid analog), single-stranded RNA	ES
TLR8	Single-stranded RNA	ES
TLR9	Unmethylated CpG DNA	ES
Ci-TLR1	Zymosan (yeast cell wall)	PM and ES
Ci-TLR2	Heat-killed <i>Legionella pneumophila</i> (HKLP, Gram-negative) poly(I:C) (double-stranded-RNA) Flagellin (bacterial flagellar filament)	

PM, plasma membrane; ES, endosome.

The Ci-TLR1 and Ci-TLR2 genes were expressed intensively in the stomach, intestine and numerous hemocytes and, to a lesser degree, the central nervous system.¹⁵ These findings indicate that Ci-TLRs function mainly in the alimentary tracts and hemocytes. Intriguingly, both of Ci-TLRs, unlike any vertebrate TLRs, were present on both the plasma membrane and a number of late endosomes.¹⁵ Moreover, Ci-TLR1 and Ci-TLR2 activated NF- κ B in response to multiple TLR ligands (Table 1), which are recognized by different mammalian TLRs. Zymosan (*Saccharomyces cerevisiae* cell wall) for hTLR2, heat-killed *Legionella pneumophila* (HKLP, a Gram-negative bacterium) for hTLR2, double-stranded RNA, poly(I:C) for hTLR3, *Salmonella typhimurium* Flagellin (the major component of the bacterial flagellar filament) for hTLR5 elicited a dose-dependent transactivation of NF- κ B in the *ci-tlr1*- or *ci-tlr2*-expressing cells.¹⁵ Poly(I:C) also elicited approximately 4-fold and 10-fold Ci-TNF α expression in the anterior and middle intestine, respectively.¹⁵ Likewise, induction of 4-fold and 10-fold Ci-TNF α expression by Flagellin was observed in the stomach and middle intestine, respectively.¹⁵ In contrast, no Ci-TNF α induction was detected in the posterior intestine.¹⁵ These profiles of the Ci-TNF α induction are compatible with the tissue-distribution of *ci-tlr* expression; *ci-tlr1* and *ci-tlr2* are abundantly expressed in the stomach, anterior and middle intestine, but not in the posterior intestine.¹⁵ These data lead to two important conclusions. Firstly, Ci-TLRs, like vertebrate TLRs, directly recognize their PAMPs and trigger the transactivation of NF- κ B. Secondly, Ci-TLRs are 'functionally hybrid TLRs' of vertebrate cell-surface TLRs and endosome TLRs:

poly(I:C) is recognized by hTLR3 on endosomes TLR, whereas hTLR2 and hTLR5 respond to Zymosan, HKLP and Flagellin on the cell surface, respectively (Table 1). In addition, the PAMPs of Ci-TLRs are in good agreement with their cellular localization to both the plasma membrane and endosomes.

C. intestinalis possesses only two TLRs,¹⁵ whereas other deuterostome invertebrates, amphioxus and sea urchin, were found to possess a great number of TLRs or their related genes: 72 genes in amphioxus¹¹ and 222 genes in sea urchin.¹² Furthermore, the molecular phylogenetic analyses demonstrated that most of these genes were generated via species-specific gene duplication, suggesting that these deuterostome invertebrates expand TLRs or their related genes in unique lineages of innate immunity, if most of the genes are functional. These findings lead to two scenarios of evolution of TLR or their related genes. First, only a few TLR or their related genes might have existed in a common deuterostome antecedent and *C. intestinalis* conserves the ancestral characteristics. Alternatively, a common deuterostome antecedent might have numerous TLR family genes. If this is true, *C. intestinalis* should have lost a large part of ancestral TLR family genes. Instead of such a gene deletion, Ci-TLRs are highly likely to have acquired multiple PAMP recognition and intracellular localization as mentioned above. Unfortunately, Ci-TLRs are at present the only invertebrate TLRs of which intracellular localization, PAMP recognition and signaling have been investigated. Elucidation of PAMPs and intracellular localization of sea urchin, amphioxus and cyclostome TLRs is expected to contribute not only to understanding of their biological roles but also to the investigation of molecular and functional divergence of the invertebrate TLR family.

COMPLEMENT SYSTEM

The mammalian complement system is a powerful defense mechanism consisting of more than 30 plasma and cell-surface proteins interacting in the recognition and elimination of pathogens.¹⁶ Three major physiological functions of the mammalian complement system are; opsonization of the foreign particles, induction of the inflammatory reactions and cytolysis. Evolutionary studies revealed that the origin of the multi-component complement system consisting of C3, Bf (factor B) and MASP (mannan-binding lectin associated serine protease) is traced back to the common ancestor of Eumetazoa.^{17,18} In addition, marked development of the complement system by gene duplication of the key components and subsequent functional differentiation likely occurred at the early stage of vertebrate evolution.¹⁸ Thus, the urochordate complement system represents the evolutionary stage just before this development and accumulating analyses made it the best-analyzed invertebrate complement system. Several complement genes have been identified mainly from two species, *Halocynthia roretzi* and *Ciona intestinalis*. Those genes are; C3,^{19,20} Bf,^{3,21} MASPs,^{3,22} mannan-binding lectin (MBL),²³ ficolin²⁴ and CR3 alpha²⁵ and beta.²⁶ In addition, a glucose binding lectin (GBL) lacking the collagen domain was reported from *H. roretzi* as a possible functional substitute for MBL.²⁷ For the functional aspect, *H. roretzi* C3, ficolin and GBL proteins were isolated from the body fluid and were shown to act as a component of the opsonic complement system. Moreover, the C3a fragment of *C. intestinalis* C3 was shown to have a chemotactic activity,²⁸ indicating that the role of the complement system in inflammation is also conserved between mammals and urochordates. In contrast, the third activity of the mammalian complement system, cytolytic activity, has not been recognized in the urochordate complement system.

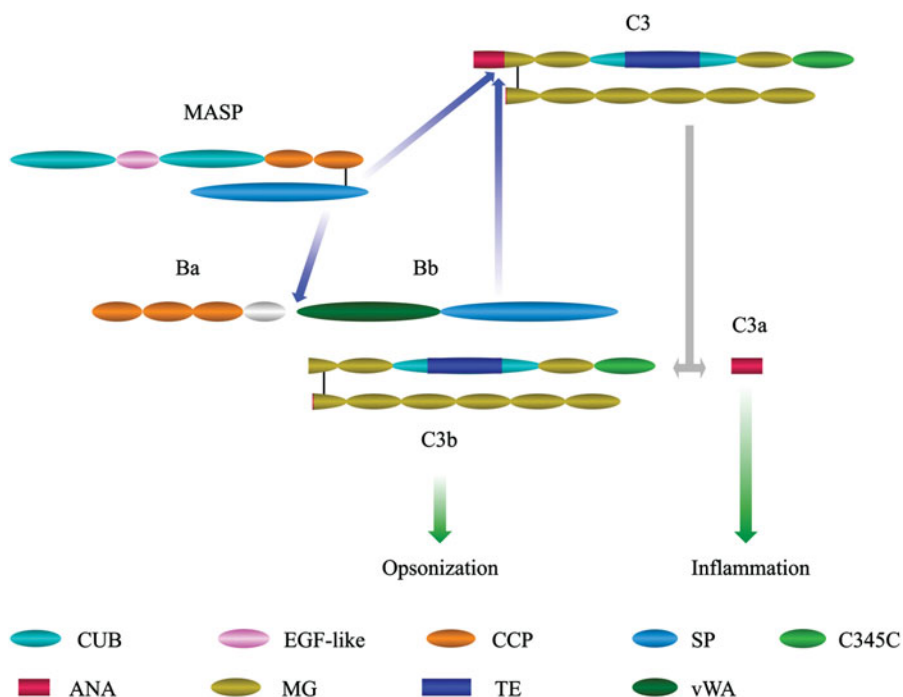


Figure 1. Schematic view of the core part of the urochordate complement system comprising C3, Bf and MASP. Conservation of the domain structure and functionally important residues of these components between mammals and urochordates suggests that the basic activation mechanism is also conserved. However, there is still no direct experimental evidence for proteolytic activation processes shown in blue arrows. The grey arrow indicates that C3 is cleaved by the C3 convertase (C3bBb) into two fragments, C3a and C3b and green arrows show biological functions of C3a and C3b. Abbreviations of domain names are: CUB, C1r, C1s, uEGF and bone morphogenetic protein; EGF-like, epidermal growth factor-like; CCP, complement control protein; SP, serine protease; C345C, C-terminal of C3, C4 and C5; ANA, anaphylatoxin; MG, macroglobulin; TE, thioester; vWA, von Willebrand factor Type A.

Although there are several C6-like genes with the membrane attack complex/perforin (MACP) domain in the *C. intestinalis* genome,³ all of them lack the C-terminal short consensus repeat (SCR) and factor I/membrane attack complex (FIM) domains reported to be essential for interaction with other complement components. Thus, it is unlikely that these C6-like molecules are integrated in the urochordate complement system. All these results indicate that the urochordate complement system represents the primitive evolutionary stage just before the development occurred in the common ancestor of vertebrates. It lacks some components and functions of the mammalian complement system and shows no sign of acquisition of unique function. Figure 1 shows a schematic representation of the putative activation mechanism of the core part of the urochordate complement system comprising C3, Bf and MASP. Conservation of most structural motifs involved in proteolytic activation and C3 convertase formation of these complement components between urochordates and mammals strongly suggests that the activation mechanism of the urochordate complement system is the same as that of the mammalian complement system, although direct experimental evidence is still missing.

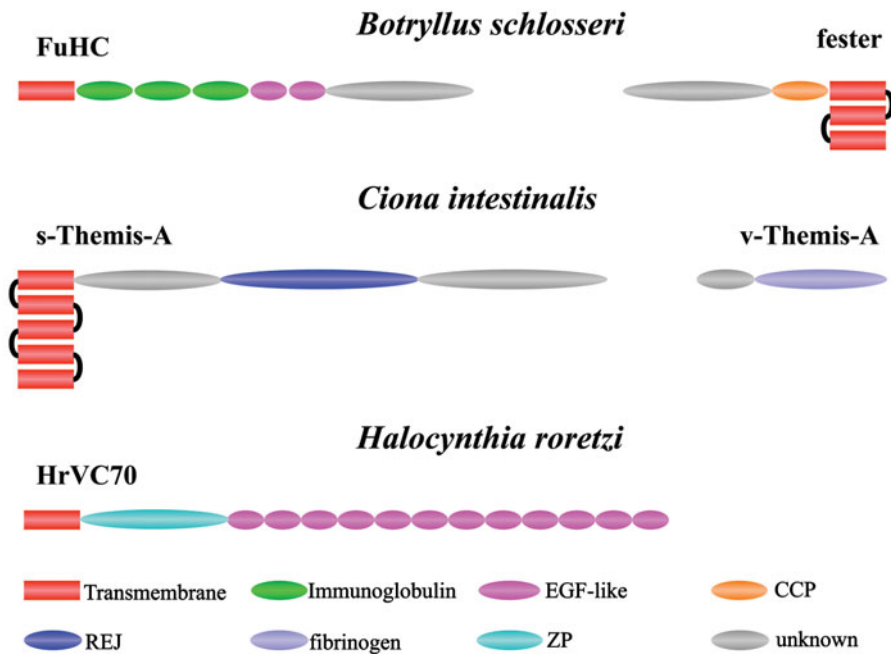


Figure 2. Domain structure of putative allorecognition molecules of three ascidian species. As shown here there is no orthologous relationship among these putative allorecognition molecules. Mutual interaction between FuHC and fester or between S-Themis and v-Themis is postulated, although it is still to be demonstrated directly. The *C. intestinalis* genome contains another set of the s-Themis and v-Themis genes containing similar domain structure as shown in this figure. Abbreviations of domain names not described in the legend to Figure 1 are: REJ, receptor for egg jelly; ZP, zona pellucida.

ALLORECOGNITION

Allorecognition is well known in vertebrates in the context of tissue transplantation where self or isogenic grafts are accepted whereas allogenic grafts are rejected. Although multiple genetic loci are involved in vertebrate allorecognition, by far the most important locus is the MHC. The evolutionary analyses indicated that the MHC was established in the common ancestor of jawed vertebrates¹ and urochordates completely lack the MHC. However, urochordates have two famous allorecognition systems working at colony fusion of the colonial ascidian²⁹ and fertilization of the solitary ascidian.³⁰ In both systems, the candidate genes for the key recognition molecules have been identified recently.

It has been known for 50 years that when two individuals of *Botryllus schlosseri*, a colonial ascidian, come into contact, they show histocompatibility reaction based on their genetic background.²⁹ If they share one or both alleles at a single histocompatibility locus, they will fuse. If they share no alleles, the colonies will reject each other. The candidate histocompatibility gene was isolated recently by positional cloning as described below and was termed *FuHC*.³¹ About 1 Mb region identified by segregation analysis was sequenced to identify a candidate gene which showed polymorphism correlating with

defined histocompatibility alleles in a fusion assay. The predicted open reading frame of this gene encoded a Type I transmembrane protein of 1007 amino acids in length. The amino terminus begins with a signal sequence, followed by an extracellular epidermal growth factor (EGF) repeats, two tandem immunoglobulin domains and the transmembrane domain and intracellular tail (Fig. 2). The high degree of polymorphism was demonstrated by identifying 18 alleles from 10 wild individuals. Most of allelic differences are single amino acid substitutions spread throughout the extracellular domains, with no obvious highly variable regions. Expression pattern analyzed by RT-PCR and in situ hybridization indicated that strong expression is observed in epithelia of ampullae and in a subset of blood cells, intimately associated with histocompatibility.

Another candidate for possible histocompatibility components is *fester* encoded near the *FuHC* locus.³² The *fester* locus is highly polymorphic although this polymorphism does not contribute to histocompatibility, since it is not correlated with defined histocompatibility alleles. The *fester* is a Type I membrane protein having a signal peptide and several extracellular domains containing a single SCR domain. Alternative splicing generates several forms both membrane bound and secreted, all expressed in tissues intimately associated with histocompatibility. SiRNA-mediated knockdown of *fester* resulted in no histocompatibility reaction in both compatible and incompatible pairs, suggesting that *fester* is a receptor involved in histocompatibility. These data suggest that *FuHC* and *fester* are involved in allorecognition of *B. schlosseri*, although the underlying molecular mechanism including the possibility that *FuHC* and *fester* bind to each other is still to be clarified.

Ascidians are hermaphroditic and exhibit self-incompatibility (SI) at fertilization, self-sterility. Two species, *Ciona intestinalis* and *Halocynthia roretzi*, have been studied in detail for their SI system.³⁰ The SI system of *C. intestinalis* is genetically determined by multiple loci and takes place in the interaction between sperm and vitelline coat (VC), since removal of VC by acid treatment results in the loss of self-sterility. Recently, positional cloning of the SI loci was carried out using acid-induced self-fertilized siblings, the draft genome sequences and the detailed physical map.³³ Two loci A and B were identified in chromosome 2q and 7q, respectively. At both loci, a pair of genes termed s-Themis and v-Themis are present with a curious configuration that the v-Themis gene is located in the first intron of the s-Themis gene in opposite transcriptional direction. s-Themis is a polycystin-I receptors and is expressed in testis. On the other hand, v-Themis is fibrinogen-like molecule and is a component of VC. Both s-Themis and V-Themis are highly polymorphic and autologous interaction between them is believed to reduce the binding ability of sperms. *H. roretzi* has much more strict SI system than *C. intestinalis*. In addition, *H. roretzi* has another allorecognition system termed "contact reaction", in which allogenic hemocytes show cytotoxic reactions. Although the common underlying mechanism is suggested for the SI system and contact reaction, their molecular basis is still to be clarified. Recently, a candidate for the VC molecule responsible for the SI system was reported.³⁴ A 70 kDa VC protein termed HrVC70 consists of 12 EGF-like repeats and show a high degree of polymorphism. HrVC70-agarose beads binds more nonself-sperms than self-sperms and pretreatment of sperm with nonself-HrVC70 more strongly inhibited fertilization than the pretreatment of sperm with self-HrVC70. These results suggest that HrVC70 is involved in the SI system of *H. roretzi*. Therefore, at least two different SI systems seem to be present in ascidians, although detailed molecular mechanism and evolution are still to be clarified.

CONCLUSION

Two major arms of innate immunity, TLRs and complement, show clear orthologous relationship between urochordates and vertebrates. Although functional information is still missing, Cnidaria also has the orthologous genes of vertebrate TLRs and complement, suggesting that their evolutionary origin can be traced back to the common ancestor of eumetazoa. In contrast, allorecognition systems of ascidians seem to be innovated in the urochordate lineage and use totally different genes from vertebrate MHC. Although vertebrate MHC has evolved as the antigen presentation system of adaptive immunity and its involvement in allorecognition is an accidental side effect, allorecognition seems to be the original purpose of the urochordate allorecognition systems.

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