

The *stuA* gene controls development, adaptation, stress tolerance, and virulence of the dermatophyte *Trichophyton rubrum*

Elza A.S.Lang, Tamires A.Bitencourt, Nalu T.A.Peres, LuciaLopes, Larissa G.Silva, Rodrigo A.Cazzaniga, AntonioRossi, Nilce M.Martinez-Rossi

<https://doi.org/10.1016/j.micres.2020.126592>

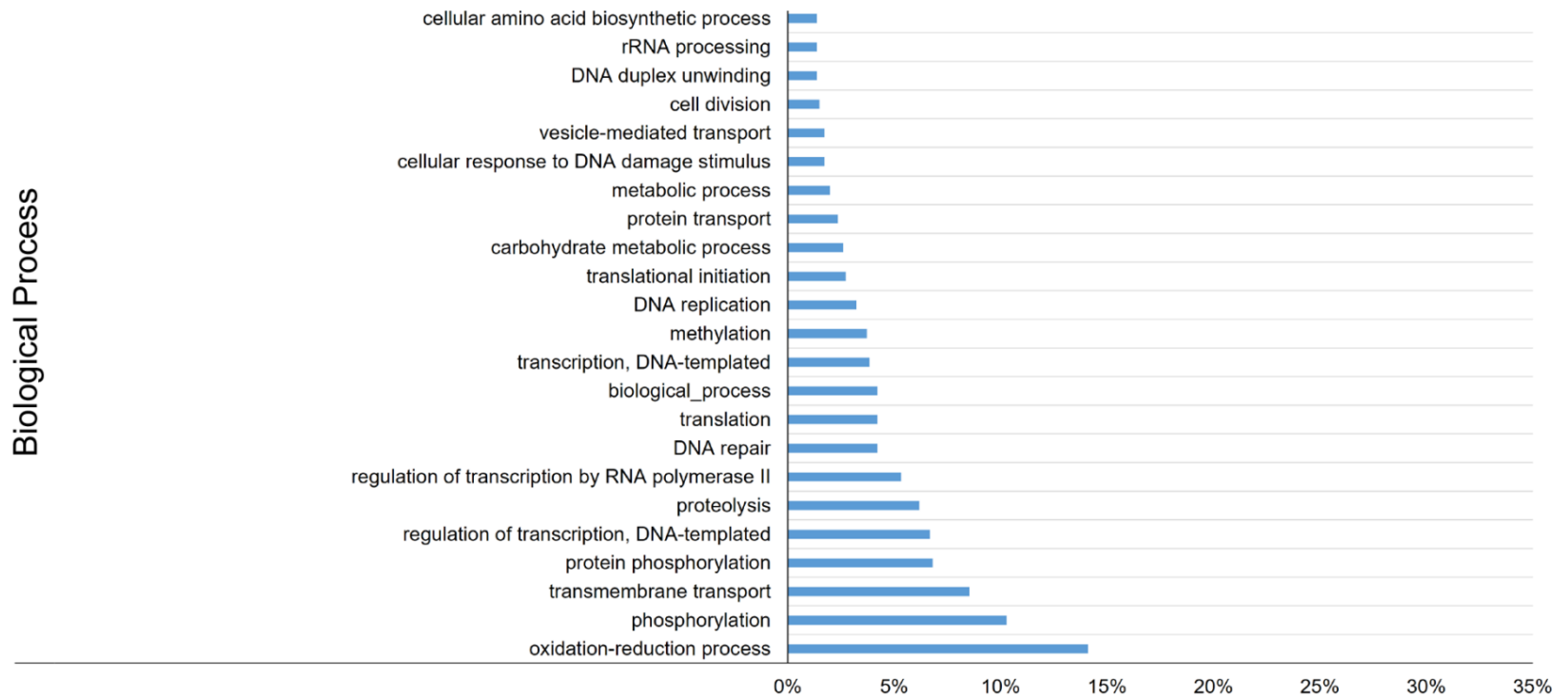
Supplementary Table 1. Primers used in this study

Primer name	Sequence (5'- 3')	Application
Primers used for gene targeting constructs		
M13_F	GTAAAACGACGGCCAGT	Amplification of <i>hph</i> cassette
M13_R	CAGGAAACAGCTATGAC	
P1	TGTCAGCGTCTCGTCTCTACC	Amplification of <i>stuA</i> 5' flanking sequence
P2	<u>ACTGGCCGTCGTTTTACC</u> ATCCATGTACGGCTGTGTC ^(a)	
P3	<u>GTCATAGCTGTTTCCTGAAACGCTCATCGACTCTGC</u> ^(b)	Amplification of <i>stuA</i> 3' flanking sequence
P4	TGCTTCTTTTCTGCACGTTG	
P5	ATAGTGATCAACCAGACCAGCCAATCTGC	Generation of 5' split-marker fragment
H1	GATGTTGGCGACCTCGTATT	
P6	TAACAGGCCTGGATGGCGTGGTGAGTATCT	Generation of 3' split-marker fragment
H2	CTGCCTGAAACCGAACTGC	
Primers used for mutant screening		
S1	AGGCAGCCTCTGCTTCCA	Mutant screening
S2	TTTGTACCGTTGATCATATGGTTGT	

P1	TGTCAGCGTCTCGTCTCTACC	Mutant screening
P4	TGCTTCTTTTCTGCACGTTG	
Primers used for qRT-PCR		
stuA_S1	AGGCAGCCTCTGCTTCCA	qRT-PCR
stuA_S2	TTTTGTACCGTTGATCATATGGTTGT	
rpb2_F	TGCAGGAGCTGGTGGGAAGA	qRT-PCR
rpb2_R	GCTGGGAGGTACTGTTTGATCAA	
hypA_F	TCCTGCTGCAACACTGAGAC	qRT-PCR
hypA_R	AACCCTTGAGGAGGGAGAAG	

^(a) The sequence with homology with M13_F sequence is underlined

^(b) The sequence with homology with M13_R sequence is underlined



Supplementary Figure S1. *In silico* analysis of StuA target genes in *T. rubrum* genome. Gene ontology-based functional categorization of genes in the *T. rubrum* genome with a recognition site for the motif consensus (A/TCGCGT/ANA/C) in their promoter regions.