

Ising model

①

Let us suppose a lattice composed of N spins with magnetic dipole moment μ strength $g\mu_B \sqrt{J(J+1)}$ and degeneracy $(2J+1)$. Each atom interacts with their q nearest neighbors and they are subject to the magnetic field \vec{H} .

The analysis of this system reveals that, depending on the dimensionality, a ferromagnetic-paramagnetic can emerge at the critical temperature T_c .

Most of systems have a curve $M(0, T)$ $M \propto T/T_c$ that are better fitted for $J = 1/2$. This reveals that the ferromagnetism is associated to the spin of electron and not with the orbital contribution. Next, I will present a better discussion about the interaction which is responsible for the ferromagnetism.

It is called "exchange interaction". When two electrons have equal spins, they should be "more separated" from each other (due to the Pauli-exclusion principle).

Conversely, when their spins are different, they can be closed to each other. Hence, different spatial separations give rise

to different electrostatic interactions and they are closely related to the relative orientations & their spins.

Now, we are going to present an argument/explanation, based on perturbation theory, about the "exchange interaction".

The wave-function of two electrons should be anti-symmetric. For instance, there are two possibilities:

$$\underbrace{\phi_s(\vec{r}_1, \vec{r}_2)}_{\begin{array}{l} \text{symmetric} \\ \text{spatial} \\ \text{wave-function} \end{array}} \times_A \underbrace{\chi_A(s_1, s_2)}_{\begin{array}{l} \text{anti-symmetric} \\ \text{spin} \\ \text{wave-function} \end{array}}$$

or

$$\underbrace{\phi_A(\vec{r}_1, \vec{r}_2)}_{\begin{array}{l} \text{anti-symmetric} \\ \text{spatial} \end{array}} \times \underbrace{\chi_s(s_1, s_2)}_{\begin{array}{l} \text{symmetric spin} \end{array}}$$

The first one have a spatial wave-function proportional to $\frac{1}{\sqrt{2}} [\phi_1(\vec{r}_1) \phi_2(\vec{r}_2) + \phi_2(\vec{r}_1) \phi_1(\vec{r}_2)]$

Whereas the latter one have a spatial wave function of type $\frac{1}{\sqrt{2}} [\phi_1(\vec{r}_1) \phi_2(\vec{r}_2) - \phi_2(\vec{r}_1) \phi_1(\vec{r}_2)]$

Regarding the spin wave-function, the first one is anti-symmetric and has total spin $s=0$ (singlet) and is of type $\frac{1}{\sqrt{2}} [|\Psi_L\rangle - |\Psi_R\rangle]$, whereas

the latter one is symmetric and accounts for three possibilities: $|111\rangle$, $|1\downarrow\downarrow\rangle$ or $\frac{1}{\sqrt{2}}(|1\uparrow\rangle + |1\downarrow\rangle)$

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An estimate of the difference energy ~~about~~
between above states is obtained by considering

$$\text{the quantity } \langle \Psi_1 | -\frac{e^2}{r_{12}} | \Psi_1 \rangle - \langle \Psi_2 | -\frac{e^2}{r_{12}} | \Psi_2 \rangle$$

where $|\Psi_1\rangle \rightarrow$ first state (Symmetric Spatial Wave function)

$|\Psi_2\rangle \rightarrow$ second state (Anti-Symmetric Spatial Wave function) $|\Psi_2\rangle$

Both $|\Psi_1\rangle$ and $|\Psi_2\rangle$ can be expressed as: $|\Psi_{1(2)}\rangle = \frac{1}{\sqrt{2}} [|\kappa_1^1\rangle + |\kappa_2^1\rangle + |\kappa_1^2\rangle + |\kappa_2^2\rangle]$

$\langle \Psi_1 | -\frac{e^2}{r_{12}} | \Psi_1 \rangle$ can be written as $\langle \Psi_2 | -\frac{e^2}{r_{12}} | \Psi_2 \rangle$

$$\int d^3\vec{r}_1 \int d^3\vec{r}_2 \langle \Psi_1 | \vec{r}_1, \vec{r}_2 \rangle \langle \vec{r}_1, \vec{r}_2 | -\frac{e^2}{r_{12}} | \Psi_1 \rangle =$$

$$= A + B + C + D \quad \text{where}$$

$$A = \int d^3\vec{r}_1 \int d^3\vec{r}_2 |\phi_1(\vec{r}_1)|^2 |\phi_2(\vec{r}_2)|^2 \left(-\frac{e^2}{r_{12}}\right)$$

$$B = \int d^3\vec{r}_1 \int d^3\vec{r}_2 |\phi_2(\vec{r}_1)|^2 |\phi_1(\vec{r}_2)|^2 \left(-\frac{e^2}{r_{12}}\right)$$

$$C = \int d^3\vec{r}_1 \int d^3\vec{r}_2 \phi_1^*(\vec{r}_1) \phi_2^*(\vec{r}_2) \left(-\frac{e^2}{r_{12}}\right) \phi_2(\vec{r}_1) \phi_1(\vec{r}_2)$$

$$D = \int d^3\vec{r}_1 \int d^3\vec{r}_2 \phi_1^*(\vec{r}_2) \phi_2^*(\vec{r}_1) \left(-\frac{e^2}{r_{12}}\right) \phi_1(\vec{r}_1) \phi_2(\vec{r}_2)$$

The quantity $\langle \psi_2 | -\frac{e^2}{r_{12}} | \psi_1 \rangle$ is analogous
and given by

$$(A + B) - (C + D)$$

Hence, the difference between the "singlet"
and "triplet" states is given by $2(C + D)$
and depends solely on the "exchange"
term. This energy is closely dependent
on the coulombian interaction.

Now, we are going to show that
a Hamiltonian of type $H = \vec{\nabla}_1 \cdot \vec{\nabla}_2 + D$
is able to reproduce the "separation"
of two states of electrons. More specifically,
we are going to evaluate $\langle s | \vec{\nabla}_1 \cdot \vec{\nabla}_2 | s \rangle -$
 $\langle t | \vec{\nabla}_1 \cdot \vec{\nabla}_2 | t \rangle$
 \uparrow triplet
 \downarrow singlet

$$\vec{\nabla}_1 \cdot \vec{\nabla}_2 = S_{1x} S_{2x} + S_{1y} S_{2y} + S_{1z} S_{2z},$$

where S_x, S_y, S_z are Pauli matrices

Given by $S_x = \frac{\hbar}{2} \sigma_x, S_y = \frac{\hbar}{2} \sigma_y, S_z = \frac{\hbar}{2} \sigma_z$

$$\text{and } \sigma_x = \begin{pmatrix} 0 & 1 \\ 1 & 0 \end{pmatrix}, \quad \sigma_y = \begin{pmatrix} 0 & -i \\ i & 0 \end{pmatrix} \quad (3)$$

$$\sigma_z = \begin{pmatrix} 1 & 0 \\ 0 & -1 \end{pmatrix}.$$

All of them are Hermitian matrices and have the same eigenvalues $\lambda = \pm 1$.

Their eigenvectors are given by

$$\sigma_x |\pm\rangle_x = \pm |\pm\rangle_x$$

$$\sigma_y |\pm\rangle_y = \pm |\pm\rangle_y$$

$$\sigma_z |\pm\rangle = \pm |\pm\rangle$$

$$\text{where } |\pm\rangle_x = \frac{1}{\sqrt{2}} [|+\rangle \pm |-\rangle]$$

$$|\pm\rangle_y = \frac{1}{\sqrt{2}} [|+\rangle \pm i |-\rangle]$$

It is easy to show that

$$\left\{ \begin{array}{l} S_{1x} S_{2x} |+-\rangle = \frac{\hbar^2}{4} |-+\rangle \\ S_{1x} S_{2x} |-+\rangle = \frac{\hbar^2}{4} |+-\rangle \\ S_{1y} S_{2y} |+-\rangle = \frac{\hbar^2}{4} |-+\rangle \\ S_{1y} S_{2y} |-+\rangle = \frac{\hbar^2}{4} |+-\rangle \\ S_{1z} S_{2z} |+-\rangle = -\frac{\hbar^2}{4} |-+\rangle \end{array} \right. \quad \left\{ \begin{array}{l} S_{1x} S_{2x} |++\rangle = 0 \\ S_{1y} S_{2y} |++\rangle = 0 \\ S_{1z} S_{2z} |++\rangle = \frac{\hbar^2}{4} |++\rangle \\ S_{1x} S_{2x} |--\rangle = 0 \\ S_{1y} S_{2y} |--\rangle = 0 \\ S_{1z} S_{2z} |--\rangle = \frac{\hbar^2}{4} |--\rangle \end{array} \right.$$

Hence, a singlet state $|s\rangle = \frac{1}{\sqrt{2}} [|+\rightarrow\rangle - |-\rightarrow\rangle]$

and a triplet state $|t\rangle = \frac{1}{\sqrt{2}} [|+\rightarrow\rangle + |-\rightarrow\rangle]$

have difference energies given by

$$\langle s | \vec{s}_1 \cdot \vec{s}_2 | s \rangle = -\frac{3}{4} \hbar^2 \tilde{c} \quad \rangle = -\hbar^2 \tilde{c}$$

$$\langle t | \vec{s}_1 \cdot \vec{s}_2 | t \rangle = \frac{\hbar^2}{4} \tilde{c}$$

which is qualitatively similar to the previous argument, allowing us to identify the coefficient \tilde{c} with the scalar product between spin vectors and the exchange term. When $\tilde{c} > 0$ the ferromagnetism is favored, whereas $\tilde{c} < 0$ favors the anti-ferromagnetism.

Once presented physical arguments for the Hamiltonian $\tilde{H} = -J \sum_{\langle i,j \rangle} \vec{s}_i \cdot \vec{s}_j + H \sum_i \vec{s}_i$,

We are going to study their main features in the realm of mean-field theory and beyond the mean-field. For simplicity

We are going to consider the simplified case $\vec{s}_i \cdot \vec{s}_j = s_{iz} s_{jz}$ which is

the Ising model and quantum treatment is not required.

Bragg - Williams

mean - field - approximation

This first kind of mean - field approximation is called Bragg - Williams and consists of neglecting correlation among spin sites

$$\langle \sigma_i \sigma_j \rangle \approx \langle \sigma_i \rangle \langle \sigma_j \rangle$$

From this approximation, the ensemble average of above Hamiltonian $\left(H = -J \sum_{\langle ij \rangle} \sigma_i \sigma_j - h \sum_i \sigma_i \right)$ reads.

$$\text{with } \sigma_i = \pm 1$$

$$U = \langle H \rangle$$

$$= -JdN \langle \sigma_i \rangle \langle \sigma_j \rangle - hN \langle \sigma_i \rangle,$$

where \underline{d} is the system dimensionality.

Since we are dealing with the simplest case of $S_i = \pm \frac{1}{2}$ (or $\sigma_i = \pm 1$)

we have that the magnetization per site is given by

$$m = \langle \sigma_i \rangle = \frac{N_+ - N_-}{N} \quad \begin{matrix} \uparrow \\ \text{number of} \\ \text{spins "up"} \end{matrix} \quad \begin{matrix} \rightarrow \\ \text{number of} \\ \text{spins "down"} \end{matrix}$$

magnetization per site

and \underline{U} then reads

$$U = -JdN m^2 - hNm.$$

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The entropy system is expressed in terms of N_+ and N_- and given by

$$S = K_B \ln \frac{N!}{N_+! N_-!} \quad \text{where } N_+ - N_- = Nm \\ N_+ + N_- = N$$

$$= K_B \ln \frac{N!}{\left(\frac{N+m}{2}\right)! \left(\frac{N-m}{2}\right)!}$$

By using the Stirling approximation, S is given by

$$S = -K_B \left[\left(\frac{N+m}{2}\right) \ln \left(\frac{1+m}{2}\right) - \left(\frac{N-m}{2}\right) \ln \left(\frac{1-m}{2}\right) \right]$$

$$\text{or } \nu = \frac{S}{N} = -K_B \left[\left(\frac{1+m}{2}\right) \ln \left(\frac{1+m}{2}\right) + \left(\frac{1-m}{2}\right) \ln \left(\frac{1-m}{2}\right) \right]$$

The functional $g(T, H; m)$ is given by

$$g(T, H; m) = u(T, H; m) - T S(m)$$

$$\text{or } g(T, H) = \min_m \{ g(T, H; m) \}$$

$$\text{where } g(T, H; m) = -Jdm^2 - hm + K_B T \left[\left(\frac{1+m}{2}\right) \ln \left(\frac{1+m}{2}\right) + \left(\frac{1-m}{2}\right) \ln \left(\frac{1-m}{2}\right) \right]$$

The minimization of above functional leads to the following equation:

$$\left(\frac{\partial g}{\partial m} \right) = 0 \Leftrightarrow -2Jdm - h + K_B T \left[\left(\frac{1+m}{2} \right)^{-\frac{1}{2}} + \frac{1}{2} \ln \left(\frac{1+m}{2} \right) + \left(\frac{1-m}{2} \right)^{-\frac{1}{2}} - \frac{1}{2} \ln \left(\frac{1-m}{2} \right) \right]$$

$$\Rightarrow h = -2Jdm + \frac{K_B T}{2} \ln \frac{1+m}{1-m} \quad \text{or}$$

even $m = \tanh \left(\frac{1}{2} \beta J dm + \beta H \right)$,

which is the previous Curie-Weiss with ($\beta \lambda = 2Jd$).

The critical point T_C is then given by $\frac{2Jd}{K_B T_C} = 1$ or $T_C = \frac{2Jd}{K_B}$.

Therefore, such Mean-field treatment Ising model predicts a ferromagnetic-paramagnetic phase transition, even for $d=1$ (this is actually incorrect),

whose equation of state is the same as before (but proposed ^{under a} phenomenological way)

Curie-Weiss Model

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Instead of neglecting correlations, as performed previously, one can modify the Ising model allowing it exactly solvable. Such a model modification is given by

$$H \rightarrow -\frac{J}{2N} \sum_{i=1}^N \sum_{j=1}^N \sigma_i \sigma_j - H \sum_i \sigma_i$$

interaction among all spins (instead of only among nearest-neighbor ones), but they decay with the distance as $\frac{1}{N}$, in order to ensure the "thermodynamic limit". The partition function then reads

$$\begin{aligned} Z &= \sum_{\{\sigma\}} e^{\frac{\beta J}{2N} \sum_{i=1}^N \sum_{j=1}^N \sigma_i \sigma_j + \beta H \sum_i \sigma_i} \\ &= \sum_{\{\sigma\}} e^{\frac{\beta J}{2N} \left(\sum_{i=1}^N \sigma_i \right)^2 + \beta H \sum_i \sigma_i} \end{aligned}$$

By using the Gaussian identity

$$\int_{-\infty}^{\infty} e^{-x^2 + 2ax} dx = \sqrt{\pi} a^2, \text{ the partition function then reads}$$

$$\frac{\beta J}{2N} \left(\sum_{i=1}^N \sigma_i \right)^2 = \frac{1}{\sqrt{\pi}} \int_{-\infty}^{\infty} e^{-x^2 + 2x \sqrt{\frac{\beta J}{2N}} \sum_{i=1}^N \sigma_i} dx$$

and then we have that

$$\begin{aligned} Z &= \frac{1}{\sqrt{\pi}} \int_{-\infty}^{\infty} e^{-x^2 + 2 \sqrt{\frac{\beta J}{2N}} x \sum_{i=1}^N \sigma_i + \beta H} \frac{dx}{\sum_{i=1}^N \sigma_i} \\ &= \frac{1}{\sqrt{\pi}} \int_{-\infty}^{\alpha} e^{-x^2} dx \sum_{i=1}^N e^{-\sum_{i=1}^N \sigma_i} \\ &= \frac{1}{\sqrt{\pi}} \int_{-\infty}^{\alpha} e^{-x^2} dx \left(e^{\alpha} + e^{-\alpha} \right)^N \\ &= \frac{1}{\sqrt{\pi}} \int_{-\infty}^{\alpha} e^{-x^2} dx - (x^2 - N \ln(2 \cosh \alpha)) \end{aligned}$$

$$\text{where } \alpha = 2 \sqrt{\frac{2 \beta J}{N}} x + \beta H$$

By introducing the new variables

$$\sqrt{\frac{2 \beta J}{N}} x = \beta J m, \quad \text{above partition}$$

$$\begin{aligned} \text{function becomes } & - \left[\frac{N}{2 \beta J} \beta^2 J^2 m^2 - N \ln \left[2 \cosh \left(\beta J m + \beta H \right) \right] \right] \\ Z = \sqrt{\frac{N \beta J}{2 \pi}} & \int_{-\infty}^{\infty} dm e^{-\left[\frac{N}{2 \beta J} \beta^2 J^2 m^2 - N \ln \left[2 \cosh \left(\beta J m + \beta H \right) \right] \right]} \end{aligned}$$

$$Z = \sqrt{\frac{NPJ}{2\pi}} \int_{-\infty}^{\infty} dm e^{-N\beta g(T, H; m)} \quad (7)$$

Where

$$g(T, H; m) = \frac{\beta m^2}{2} - \frac{1}{\beta} \ln \left[2 \cosh (\beta Jm + \beta H) \right]. \quad (*)$$

above integral can be estimated through the Laplace method:

$$\int_a^b e^{Mf(x)} dx \approx e^{Mf(x_0)} \int_a^b e^{-M \left[f''(x_0) \left(\frac{x-x_0}{2} \right)^2 \right]} dx$$

Minimizing the above expression for (*) we have that

$$\frac{\partial g}{\partial m} = 0 \Leftrightarrow \beta m - \frac{1}{\beta} \frac{\sinh(\beta Jm + \beta H)}{\cosh(\beta Jm + \beta H)} \quad (P)$$

We obtain that

$$m = \tanh(\beta Jm + \beta H),$$

which is the same equation of state as before.

Mean-field approximation - ~~using~~ Bogoliubov inequality

This approach consists of "replacing" the expressions for the system we are interested (in which analytic solutions are generally not available) for an exactly solvable "upper ~~lower~~ bound".

More specifically, let us consider the Hamiltonian H (we are not able to solve it exactly) and the H_0 (exactly solvable) and ΔH_1 ^{being} the difference between H and H_0 .

We then have that

$$H(\lambda) = H_0 + \lambda H_1$$

$\lambda=0 \rightarrow$ one recovers the exactly solvable case H_0

$\lambda=1 \rightarrow$ we recover the Hamiltonian H we want to study.

The Bogoliubov inequality consists of "replacing / approaching" the Helmholtz / Gibbs free energy for the following expression

$$F(\lambda) \leq F(0) + \lambda \langle H_1 \rangle_0,$$

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where, Γ_0 is the free energy of the Hamiltonian (non perturbed) H_0 and $\langle \mathcal{H}_1 \rangle_0$ is the ensemble average of the Hamiltonian \mathcal{H}_1 , evaluated over the space of configurations respecting the Hamiltonian H_0 .

Let us prove the above relation.

For instance, we have that the free energy (Gibbs) $F(\lambda)$ is given by

$$-\beta F(\lambda) = \ln \left(\underbrace{\sum_{\sigma} e^{-\beta \mathcal{H}(\lambda)} }_{\text{configurations}} \right)$$

$\underbrace{\phantom{\sum_{\sigma} e^{-\beta \mathcal{H}(\lambda)}}}_{\text{partition function } Z(\lambda)}$

By deriving above expression with respect to λ we have that

$$-\beta \frac{dF(\lambda)}{d\lambda} = -\beta \frac{\sum_{\sigma} (-\beta \mathcal{H}_1) e^{-\beta(H_0 + \lambda \mathcal{H}_1)}}{\sum_{\sigma} e^{-\beta(H_0 + \lambda \mathcal{H}_1)}}$$

$= \langle \mathcal{H}_1 \rangle$

By proceeding analogously with $\frac{d^2 F}{d \lambda^2}$ we (10)

have that

$$\frac{d^2 F}{d \lambda^2} = \frac{d}{d \lambda} \left[\frac{\sum_{\sigma} H_{\sigma} e^{-\beta(H_0 + \lambda H_1)}}{\sum_{\sigma} e^{-\beta(H_0 + \lambda H_1)}} \right]$$

$= -\beta(A - B)$, where

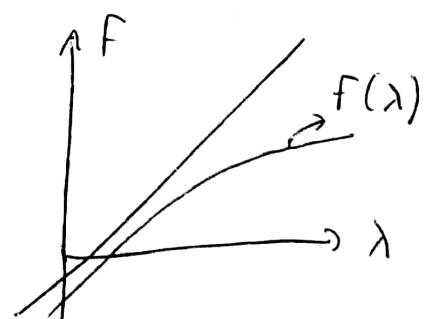
$$A = \frac{\sum_{\sigma} H_{\sigma}^2 e^{-\beta(H_0 + \lambda H_1)}}{\sum_{\sigma} e^{-\beta(H_0 + \lambda H_1)}} \quad 1$$

$$B = \left(\frac{\sum_{\sigma} H_{\sigma} e^{-\beta(H_0 + \lambda H_1)}}{\sum_{\sigma} e^{-\beta(H_0 + \lambda H_1)}} \right)^2 \quad 2$$

since $A - B \geq 0$ (always) the

derivative $\frac{d^2 F}{d \lambda^2} \leq 0$ and hence

the Gibbs free energy is concave with respect to λ and then it must to have the following shape



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Which justifies the following upper bound

$$F(\lambda) \leq F(0) + \lambda \left(\frac{dF}{d\lambda} \right) \Big|_{\lambda=0}$$



straight line whose linear coefficient
taken at $\lambda = 0$.

Since $\frac{dF}{d\lambda} = \langle \gamma_1 \rangle$ we arrive
at the following relation

$$F(\lambda) \leq F(0) + \lambda \langle \gamma_1 \rangle_0 ,$$

where

$$\langle \gamma_1 \rangle_0 = \sum_{\sigma} \gamma_1 e^{-\beta \gamma_0}$$

and

$$\sum_{\sigma} e^{-\beta \gamma_0}$$

$$f(0) = -K_B T \ln Z_0, \text{ with } Z_0 = \sum_{\sigma} e^{-\beta \gamma_0}.$$

Next, we shall exemplify for the

$$\text{Let } H = -J \sum_{\langle i,j \rangle} \sigma_i \sigma_j - H \sum_i \sigma_i \rightarrow \text{the Hamiltonian}$$

in which we don't know the exact solution and $H_0 = -\eta \sum_i \sigma_i$ the Hamiltonian which is exactly solvable

$$H_1 = +H - H_0 = -J \sum_{\langle i,j \rangle} \sigma_i \sigma_j - (H - \eta) \sum_i \sigma_i$$

$$\langle H_1 \rangle = -J \left\langle \sum_{\langle i,j \rangle} \sigma_i \sigma_j \right\rangle_0 - (H - \eta) \left\langle \sum_i \sigma_i \right\rangle_0$$

$$Z_0 = \sum_{\{\sigma\}} e^{+\beta \eta \sum_i \sigma_i} = (2 \cosh \beta \eta)^N$$

$$\text{and then } F(0) = -N k_B T \ln(2 \cosh \beta \eta)$$

$$\begin{aligned} \left\langle \sum_{\langle i,j \rangle} \sigma_i \sigma_j \right\rangle_0 &= N d \left\langle \sigma_i \sigma_j \right\rangle_0 \\ &= N d \left\langle \sigma_i \right\rangle_0 \left\langle \sigma_j \right\rangle_0 \\ &\quad \uparrow \text{the Hamiltonian } H_0 \end{aligned}$$

does not presenting coupling between sites \underline{i} and \underline{j} .

$$\begin{aligned} \text{Hence } \langle \sigma_i \rangle_0 &= \frac{1}{N} \frac{\sum_{\{\sigma\}} (\sum_i \sigma_i) e^{-\beta H_0}}{\sum_{\{\sigma\}} e^{-\beta H_0}} \\ &= \tanh \beta \eta \end{aligned}$$

Hence

$$\frac{\langle \tau_1 \rangle_0}{N} = - J d (\tanh \beta \eta)^2 - (H - \eta) \tanh \beta \eta$$

and the functional free energy

$$\begin{aligned} F(T, H; \eta) &= - K_B T \ln (2 \cosh \beta \eta) + \\ &\quad - J d (\tanh \beta \eta)^2 - (H - \eta) \tanh \beta \eta \end{aligned}$$

Since above expression for $F(T, H; \eta)$ is an upper bound, by maximizing it with respect to η , we have that

$$\begin{aligned} \frac{\partial F}{\partial \eta} &= - K_B T (\beta \tanh \beta \eta) - 2 J d \tanh \beta \eta \times \\ &\quad \times \left(- \frac{\beta \eta}{\cosh^2 \beta \eta} \right) + \tanh \beta \eta + \\ &\quad - (H - \eta) \frac{(-\beta)}{\cosh^2 \beta \eta} = 0. \end{aligned}$$

$$= - \frac{\beta}{\cosh^2 \beta \eta} [- 2 J d \tanh \beta \eta - H + \eta] = 0.$$

Since $\cosh^2 \beta \eta \neq 0$ we have that

$$H = \eta - 2 J d \tanh \beta \eta$$

By noting that $\langle \sigma_i \rangle_0 = \tanh \beta \eta = m$.
 (magnetization per site) one finally arrives at

$$H + 2 J d m = \eta$$

$$H + 2 J d m = \frac{1}{\beta} \tanh^{-1} m \quad \text{or}$$

$$m = \tanh(\beta H + 2 \beta J d m),$$

which correspond to the Curie-Weiss equation obtained previously under three different ways.