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ON THE EQUIVALENCE OF DILUTE ANTIFERROMAGNETS
AND FERROMAGNETS IN RANDOM EXTERNAL FIELDS:
CURIE-WEISS MODELS

by

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ON THE EQUIVALENCE OF DILUTE ANTIFERROMAGNETS AND FERROMAGNETS
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ABSTRACT

Using a method proposed by van Hemmen we compute the free energy of the Curie-Weiss version of the site-dilute antiferromagnetic Ising model in the presence of an uniform magnetic field. The solution displays an exact correspondence between this model and the Curie-Weiss version of the Ising model in the presence of a random magnetic field. The phase diagrams are discussed and a tricritical point is shown to exist.

Considerable attention has been devoted in the literature to the study of the Ising model in the presence of random magnetic fields (RMF) [1,6,7]. The critical behavior and the phase diagram of this model are believed to be related to those of a site dilute antiferromagnetic Ising model in the presence of an applied uniform magnetic field (DAF) [2,3]. This is specially relevant since site-dilute magnetic materials can be prepared and the magnetic-ordering transition as a function both of the external field and of the dilution of the magnetic species has been experimentally investigated [4].

In this letter we use an approach introduced by van Hemmen [5] to compute the free energy and the phase diagram of a Curie-Weiss mean field version of a DAF model. We then compare our results to those obtained by Salinas and Wrezinski [6] for the Curie-Weiss version of a RMF model and establish an exact correspondence between the parameters and phase diagrams of the two-models, as to show their complete equivalence.

Our mean field DAF model is described in a finite-volume $\Lambda \subset \mathbb{Z}^v$ by the hamiltonian

$$H_{DAF} = - \frac{J}{N} \sum_{\substack{i,j \in \Lambda_e \\ i \neq j}} \epsilon_i \epsilon_j \sigma_i \sigma_j - \frac{J}{N} \sum_{\substack{i,j \in \Lambda_o \\ i \neq j}} \epsilon_i \epsilon_j \sigma_i \sigma_j + \frac{J}{N} \sum_{\substack{i \in \Lambda_e \\ j \in \Lambda_o}} \epsilon_i \epsilon_j \sigma_i \sigma_j + H \sum_{i \in \Lambda} \epsilon_i \sigma_i \tag{1}$$

where $\Lambda_e = \Lambda \cap \mathbb{Z}_e^v$, $\Lambda_o = \Lambda \cap \mathbb{Z}_o^v$ with \mathbb{Z}_e^v (\mathbb{Z}_o^v) being the sublattice of \mathbb{Z}^v for which the sum of coordinates of each site are even (odd) integers. The interaction is antiferromagnetic ($J > 0$) between sites in different sublattices and ferromagnetic between sites in the same sublattice. The random

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variables $\epsilon_i \in \{0, 1\}$ describe the site dilution and they are taken to independent and identically distributed, with

$$\epsilon_i = \begin{cases} 1, & \text{probability } p \\ 0, & \text{probability } 1-p \end{cases}$$

The spin variables σ_i are, for simplicity, taken to be of Ising type: $\sigma_i = \pm 1$. The external magnetic field H is uniform and deterministic. N denotes the number of points in Λ .

Introducing the sub-lattice magnetizations:

$$m_e = \frac{1}{N} \sum_{i \in \Lambda_e} \sigma_i, \quad m_o = \frac{1}{N} \sum_{i \in \Lambda_o} \sigma_i$$

the hamiltonian (1) can be written in the form

$$H_{DAF} = -N \frac{J}{2} (m_e - m_o)^2 + NH (m_e + m_o) \quad (2)$$

As proved by van Hemmen [5] the quenched free energy density $f(\beta)$ of the system at inverse temperature β is given by:

$$-\beta f(\beta) = \max_{\vec{m} \in \mathbb{R}^2} \{Q(\vec{m}) + c^*(\vec{m})\} \quad (3)$$

where \vec{m} is the two-component vector $\vec{m} = (m_e, m_o)$,

$$Q(\vec{m}) = -\frac{\beta}{N} H_{DAF} \quad (4)$$

$$c^*(\vec{m}) = - \sup_{\vec{t} \in \mathbb{R}^2} (\vec{m} \cdot \vec{t} - c(\vec{t})) \quad (5)$$

and

$$c(\vec{t}) = \lim_{N \rightarrow \infty} \frac{1}{N} \ln \left[\frac{1}{2^N} \sum_{\{\sigma_i = \pm 1\}} \exp N \vec{t} \cdot \vec{m} \right] \quad (6)$$

We then compute:

$$\begin{aligned} c(\vec{t}) &= \frac{1}{2} \lim_{N \rightarrow \infty} \left\{ \frac{2}{N} \sum_{i \in \Lambda_e} \ln \cosh(t_e \epsilon_i) + \frac{2}{N} \sum_{i \in \Lambda_o} \ln \cosh(t_o \epsilon_i) \right\} \\ &= \frac{1}{2} \left\{ \overline{\ln \cosh(t_e \epsilon)} + \overline{\ln \cosh(t_o \epsilon)} \right\} \end{aligned} \quad (7)$$

where \bar{A} denotes the average of the quantity $A(\epsilon)$ with respect to the probability distribution of the random variable ϵ . The last equality in (7) is a consequence of the strong law of large numbers, which is a key ingredient in the technique of van Hemmen. Therefore

$$c(\vec{t}) = \frac{p}{2} \ln \cosh t_e + \frac{p}{2} \ln \cosh t_o \quad (8)$$

If we now introduce the order parameter $m = m_e - m_o$, the free energy is given by

$$\begin{aligned} f_{DAF}(\beta, J, p, H) &= \frac{1}{2} J m^2 - \frac{\beta}{2p} \ln [\cosh \beta (Jm - H)] \\ &\quad - \frac{\beta}{2p} \ln [\cosh \beta (Jm + H)] \end{aligned} \quad (9)$$

with m determined by the equation

$$\frac{2m}{p} = \tanh[\beta(Jm - H)] + \tanh[\beta(Jm + H)] \quad (10)$$

or equivalently

$$f_{\text{DAM}}(\beta, J, p, H) = \inf_m g_{\text{DAM}}(\beta, J, p, H, m) \quad (11)$$

where g_{DAM} , the free energy density in the magnetization ensemble, is defined by the r.h.s. of (9).

Let us now consider the Curie-Weiss RMF hamiltonian

$$H_{\text{RMF}} = - \frac{J}{N} \sum_{i, j \in \Lambda} \sigma_i \sigma_j + \sum_{i \in \Lambda} h_i \sigma_i \quad (12)$$

where the random variables $h_i, i \in \Lambda$ are independent and identically distributed, being equal to $\pm h$ ($h > 0$) with probability $\frac{1}{2}$. Applying the method of van Hemmen to H_{RMF} , the free energy density can be computed [6]:

$$f_{\text{RMF}}(\beta, J, h) = \frac{1}{2} J m^2 - \frac{1}{2\beta} \ln [\cosh \beta (Jm - h)] - \frac{1}{2\beta} \ln [\cosh \beta (Jm + h)] \quad (13)$$

with the order parameter m determined by

$$2m = \tanh [\beta (Jm - h)] + \tanh [\beta (Jm + h)] \quad (14)$$

or equivalently

$$f_{\text{RMF}}(\beta, J, h) = \inf_m g_{\text{RMF}}(\beta, J, h, m) \quad (15)$$

with g_{RMF} defined as the r.h.s. of (13).

Comparing expressions (9) and (13) we see that:

$$f_{\text{DAF}}(\beta, J, p, H) = p f_{\text{RMF}}(p, \beta J, H) \quad (16)$$

and

$$g_{\text{DAF}}(\beta, J, p, H, m) = p g_{\text{RMF}}(\beta, \beta J, H, \frac{m}{p}) \quad (17)$$

which establishes the correspondence between the two models.

As in [6] the phase diagram of the DAF model can be analysed by expanding g_{DAM} in powers of m around $m=0$:

$$g_{\text{DAF}}(m) = A + Bm^2 + Cm^4 + Dm^6 + \dots \quad (18)$$

where

$$\begin{aligned} A &= - \frac{p}{\beta} \ln \cosh \beta H \\ B &= \frac{J}{2} - \frac{p}{2\beta} (\beta J)^2 \operatorname{sech}^2 \beta H \\ C &= \frac{p}{2\beta} (\beta J)^4 (\operatorname{sech}^2 \beta H) (1 - 3 \operatorname{tgh}^2 \beta H) \\ D &= - \frac{p}{6\beta} (\beta J)^6 (\operatorname{sech}^2 \beta H) (\operatorname{tgh}^4 \beta H - \operatorname{tgh}^2 \beta H + \frac{2}{15}) \end{aligned} \quad (19)$$

There is a critical line determined by the condition $B=0$ i.e.

$$p\beta J = \cosh^2 \beta H \quad (20)$$

On this critical line, there is a tri-critical point determined by the conditions $B=C=0, D>0$:

$$p\beta J = \frac{3}{2}, \quad \operatorname{tgh}^2 \beta H = \frac{1}{3} \quad (21)$$

The critical line represents a first order phase transition

for $p\beta J > \frac{3}{2}$ (since $c < 0$) and a second order phase transition
 for $p\beta J < \frac{3}{2}$ (since $c > 0$).

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