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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 ^{incubated} 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. (Author)

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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 $\Lambda \subset \mathbb{Z}^d$ by the hamiltonian

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

where $\Lambda_e = \Lambda \cap \mathbb{Z}'$, $\Lambda_o = \Lambda \cap \mathbb{Z}''$ with \mathbb{Z}' (\mathbb{Z}'') being the sublattice of \mathbb{Z}^d 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

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 A .

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 $\vec{m} = (m_e, m_o)$,

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

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

and

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

We then compute:

$$\begin{aligned} c(\vec{E}) &= \frac{1}{2} \lim_{N \rightarrow \infty} \left\{ \frac{2}{N} \sum_{i \in A_0} \ln \cosh(t_0 \epsilon_i) + \frac{2}{N} \sum_{i \in A_1} \ln \cosh(t_0 \epsilon_i) \right\} \\ &= \frac{1}{2} \left\{ \overline{\ln \cosh(t_0 \epsilon)} + \overline{\ln \cosh(t_0 \epsilon)} \right\} \end{aligned} \quad (7)$$

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

$$c(\vec{E}) = \frac{p}{2} \ln \cosh t_0 + \frac{p}{2} \ln \cosh t_0 \quad (8)$$

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

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

with m determined by the equation

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

or equivalently

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

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:

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

and

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

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 $m=0$:

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

where

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

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

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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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