Lecture 21 – Weiss model of ferromagnetism

PREVIOUSLY: We investigated a paramagnet with independent, non-interacting spins. Recall the simple 2-state paramagnet (e.g. spin ½ e⁻): in a magnetic field B pointing up, each spin can either be aligned or anti-aligned, with energy $\varepsilon = -\mu_B Bs$ and $s = \pm 1$ for spin up or spin down, respectively ($\mu_B \equiv e\hbar / 2m_e$ is the "Bohr magneton").

For a single spin:

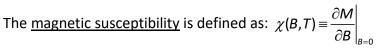
$$Z_{1} = e^{\mu_{B}B/k_{B}T} + e^{-\mu_{B}B/k_{B}T} = 2\cosh\frac{\mu B}{k_{B}T}$$

$$\langle s \rangle = \frac{1 \cdot e^{\mu_{B}B/k_{B}T} - 1 \cdot e^{-\mu_{B}B/k_{B}T}}{Z_{1}} = \tanh\frac{\mu_{B}B}{k_{B}T}$$

For a material with a density *n* of spins, the magnetization density is:

$$M(B,T) = n\mu_B \langle s \rangle = n\mu_B \tanh \frac{\mu_B B}{k_B T}$$

For small B, $\tanh x \approx x$ and $M(B,T) = n\mu_B^2 \frac{B}{k_B T}$



$$\chi(B,T) = \frac{n\mu_B^2}{k_B T} \propto \frac{1}{T}$$
 Curie law

TODAY: Ferromagnetism

Notice that for paramagnets, $M \rightarrow 0$ as $B \rightarrow 0$. Ferromagnets (e.g. iron) have a spontaneous magnetization $M \neq 0$, even when B = 0

The simple theory we used previously cannot explain this phenomenon. We need interactions between spins to get ferromagnetism. The "exchange" interaction leads to tendency for neighboring spins to align.

(Note: this interaction is purely quantum mechanical and comes from the symmetry properties of multi-particle wavefunctions under particle exchange).

For a pair of spins *i* and *j*, $\varepsilon_{exch} = -2J_{ij}s_is_j$, so

$$\varepsilon_{tot} = -\sum_{i=1}^{N} \mu_{B} B s_{i} - \frac{1}{2} \sum_{i=1}^{N} \sum_{\substack{j=1 \ i \neq j}}^{N} 2 J_{ij} s_{i} s_{j}$$

(Again, the factor of ½ is there to avoid double counting, and cancels the factor of 2 in the exchange term).

The exchange interaction is important only when wavefunctions of two particles overlap. It is therefore short-range, and we usually assume only nearest-neighbor spins interact, with a constant $J_{ii} = J$:

$$\varepsilon_{tot} = -\mu_{B}B\sum_{i} s_{i} - J\sum_{i,j}' s_{i}s_{j}$$

where $\sum_{i,j}$ signifies a sum over nearest neighbors only.

Note: this is called the Ising model, exactly solvable only under certain circumstances (1-D, 2-D with B=0)

KEY CONCEPT: Weiss molecular field

As for the van der Waals model, we will treat this problem approximately using a <u>mean field</u> model, replacing the interaction term with an <u>average over all spins</u> $j \neq i$.

For the *i*-th spin:

$$\varepsilon_{i} = -\mu_{B}Bs_{i} - Js_{i} \sum_{j \neq i} 's_{j} \approx -\mu_{B}Bs_{i} - Js_{i} \sum_{j \neq i} '\left\langle s_{j}\right\rangle$$

$$Js_{i} \left\langle s\right\rangle N_{n,n}$$

 $N_{n.n.}$ = number of nearest neighbors

Since $M = n\mu_B \langle s \rangle$, the averaged exchange term is ∞M and

$$\varepsilon_{i} \approx -\mu_{B}Bs_{i} - \mu_{B}\lambda Ms_{i} = -\mu_{B}(B + B_{eff})s_{i}$$

with $\lambda = J \ V \ / \mu_{\scriptscriptstyle R}^2$.

The average interaction term acts like an effective *B*-field $B_{eff} = \lambda M$ generated by neighboring spins, which tend to align *i*-th spin. This effective *B*-field is called the <u>Weiss molecular field</u>.

The rest is easy: take paramagnet solution and add Beff to the external B field!

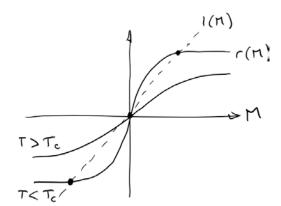
$$M(B,T) = n\mu_B \tanh \frac{\mu_B (B + B_{eff})}{k_B T}$$

First consider case where external *B* field = 0:

$$M = n\mu_B \tanh \frac{\mu_B \lambda M}{k_B T}$$
 a transcendental equation, since M is on both sides. Solve graphically

Define functions *r* and *l* for the right- and left-hand sides of the equation:

$$r(M) \equiv n\mu_{\rm B} \tanh \frac{\mu_{\rm B} \lambda M}{k_{\rm B} T}$$
, and $I(M) = M$



Intersection(s) are the solutions

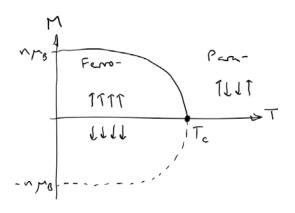
At high T, only one solution: M = 0

At low *T*, three solutions: $\underbrace{M > 0, M < 0}_{\text{stable}}$, $\underbrace{M = 0}_{\text{unstable}}$

Below some temperature T_C , system has $M \neq 0$ despite B = 0: spontaneous magnetization

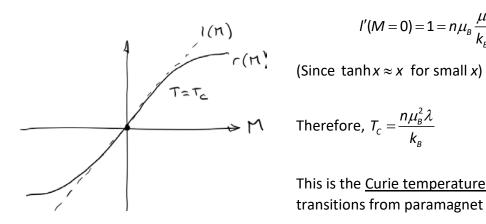
$$M > 0$$
 = spins up, $M < 0$ = spins down

In the absence of an external B field, either state $(\uparrow \uparrow \uparrow \uparrow)$ or $\downarrow \downarrow \downarrow \downarrow)$ is equally likely. We say symmetry is spontaneously broken.



Question 1: Derive an expression for T_c in terms of parameters of the system

This will occur when slopes of r(M) and l(M) at M=0 are equal:



$$I'(M=0) = 1 = n\mu_B \frac{\mu_B \lambda}{k_B T_C} = r'(M=0)$$

Therefore,
$$T_C = \frac{n\mu_B^2 \lambda}{k_B}$$

This is the Curie temperature – T at which system transitions from paramagnet to ferromagnet.

Some typical numbers: Ni has T_C = 631 K which gives $\lambda \sim 2000$

$$M(T = 0) \approx 0.17 \text{ T}$$
, so $B_{eff} = \lambda M \sim 10^{2-3} \text{ T}$

Note: this is a very large field, much larger than *B* field acting on 1 spin due to neighboring spins:

$$B_{actual} \sim \frac{\mu_B}{a^3} \sim 0.1 \text{ T} \ll B_{eff}$$
 (a = lattice spacing)

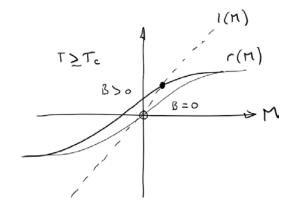
B_{eff} is not a classical dipole-dipole coupling interaction. Exchange interaction is purely quantum mechanical effect, much stronger than that expected classically.

Now consider case with external field B:

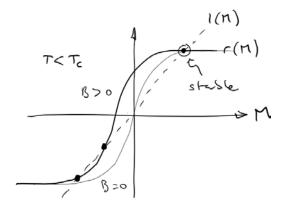
$$M(B,T) = n\mu_B \tanh \frac{\mu_B (B + \lambda M)}{k_B T}$$

Question 2: Plot M vs. B for $T > T_c$ and $T < T_c$ by solving the above equation graphically

For $T \ge T_C$

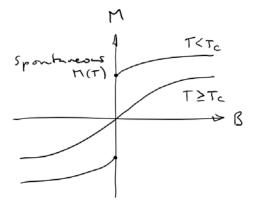


For $T < T_C$



Magnetization dependent on *B* (paramagnet)

Magnetization spontaneous, weakly dependent on *B* (ferromagnet)



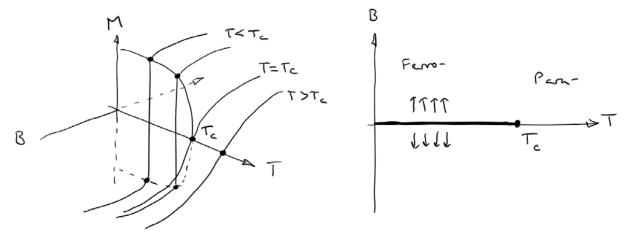
Plot M vs. B for different T

Note that the susceptibility

$$\chi(B,T) \equiv \frac{\partial M}{\partial B}\Big|_{B=0}$$
 = slope of M vs. B at origin

For
$$T \ge T_C$$
, $\chi > 0$; for $T < T_C$, $\chi \to \infty$

In 3-D, M(T,B) vs. T and B:



Looking down along M axis, we get phase diagram T_C is a critical point

This transition is second-order – no latent heat. Entropy changes continuously across transition, unlike gas-liquid transition (except at critical point).

KEY CONCEPT: Scaling laws near the critical point $(T \approx T_C, B \approx 0)$ Systems near a critical point exhibit <u>universal behavior</u>

Look at M(T) and $\chi(T)$ near T_C :

$$M = n\mu_B \tanh \frac{\mu_B \lambda M}{k_B T}$$
, $T_C = \frac{n\mu_B^2 \lambda}{k_B}$

Let's define a normalized magnetization $m = \frac{M}{n\mu_B}$, so we can write $m = \tanh \frac{mT_C}{T}$

Question 3: Show that near T_c , when m is small, the magnetization scales as $(T_c - T)^{\beta}$

When *m* is small, we can expand tanh to lowest order: $\tanh x \approx x - \frac{1}{3}x^3 + \cdots$ for x << 1

$$mathref{m} \approx m \frac{T_c}{T} - \frac{1}{3} m^{\frac{2}{3}} \left(\frac{T_c}{T} \right)^3 + \cdots \quad \text{or} \quad 1 \approx \frac{T_c}{T} \left(1 - \frac{1}{3} \left(m \frac{T_c}{T} \right)^2 \right)$$

Solving for m, we get: $m \approx \sqrt{3} \frac{T}{T_c} \left(1 - \frac{T}{T_c} \right)^{1/2}$

$$M \sim (T_c - T)^{\beta}$$
 $\beta = \frac{1}{2}$ is called a critical exponent

Now look at susceptibility χ for a small eternal field B

$$M = n\mu_{\rm B} \tanh\left(\frac{\mu_{\rm B}B}{k_{\rm B}T} + \frac{\mu_{\rm B}\lambda M}{k_{\rm B}T}\right)$$
 Define $b \equiv \frac{\mu_{\rm B}B}{k_{\rm B}}$, so that we can write $m = \tanh\left(\frac{b}{T} + m\frac{T_{\rm C}}{T}\right)$

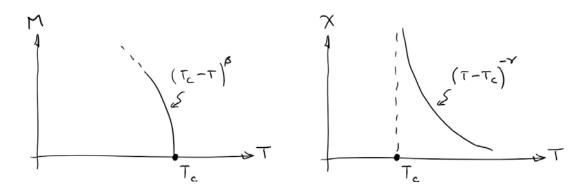
Here we only need to expand the tanh to linear order in *m*:

$$m = \frac{b}{T} + m\frac{T_c}{T} + \cdots$$

$$m\left(1 - \frac{T_c}{T}\right) = \frac{b}{T} \quad \text{and} \quad m = \frac{b}{T - T_c}$$

$$M \sim \frac{B}{T - T_c} \quad \text{and} \quad \chi \sim \left(T - T_c\right)^{-\gamma} \quad \gamma = 1$$

So χ diverges as $T \rightarrow T_c$. This is called the <u>Curie-Weiss law</u>



Amazingly, phase transitions for systems that appear unrelated (e.g. gas-liquid, ferromagnet-paramagnet) exhibit the same scaling laws near the critical point!

All mean field models can be shown to predict $\beta=\frac{1}{2}$ and $\gamma=1$ (there are other critical exponents, named α , δ , η , v, related to other physical quantities). Actual measurements give $\beta\approx 0.33$ and $\gamma\approx 1.2$. More sophisticated field theoretical techniques are required to get better agreement with experiments.

Critical	Uniaxial	Liquid-vapor	Fluid mixture	Mean field	Perturbative
exponent	magnet (exp)	(exp)	(exp)	theory	field theory
β	0.325	0.324	0.327	1/2	0.326
γ	1.240	1.241	1.235	1	1.239