

Magnetic properties of Free Electron Gas

In the presence of an applied magnetic field \vec{H} , the electron gas will develop a net magnetization via two effects

- 1) the intrinsic spins of the electrons anti-align with $\vec{H} \Rightarrow$ magnetic moments align with $\vec{H} \Rightarrow$ paramagnetic effect

Pauli Paramagnetism

- 2) the electrons move in closed orbits,
 \Rightarrow circulating currents \Rightarrow magnetic moments anti-aligned with $\vec{H} \Rightarrow$ diamagnetic effect.

Landau Diamagnetism

We consider first Pauli Paramagnetism
(A+M Chpt 31)

An electron with intrinsic spin \vec{s} ($s_z = \pm 1$) has intrinsic magnetic moment $\vec{\mu} = -\mu_0 \vec{s}$ where $\mu_0 = \frac{e\hbar}{2mc}$ is the Bohr magneton

the interaction energy of the spin with the applied magnetic field is

$$8J = -\vec{\mu} \cdot \vec{H} = \mu_0 \vec{s} \cdot \vec{H}$$

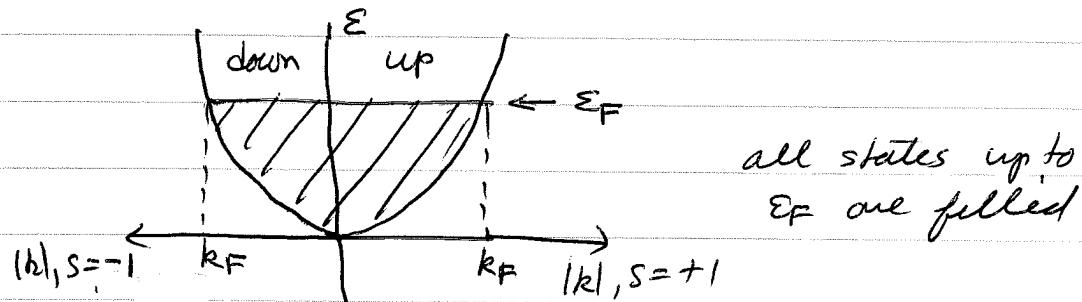
If we take $\vec{H} = H \hat{z}$, the above & FR then gives the single electron energy eigenvalues,

$$E_{ks} = \frac{\hbar^2 k^2}{2m} + s\mu_0 H \quad \text{with } s=\pm 1$$

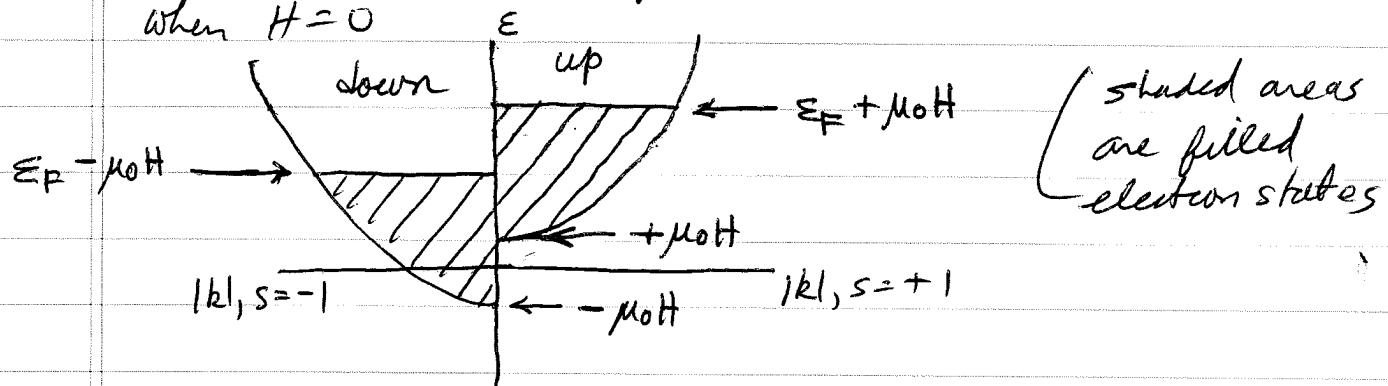
(parallel) spin up electrons, $s=+1$, increase their energy
 (antiparallel) spin down electrons, $s=-1$, decrease their energy

\Rightarrow In equilibrium, this shift of electron energy with s results in a net excess of $s=-1$ electrons

For $H=0$, the dispersion curves look like



Now imagine turning on a small \vec{H} , but keeping the $s=\pm 1$ populations of electrons the same as when $H=0$



We see that above cannot remain the ground

state, as the energy will be lowered by having ~~up~~ up electrons at $\epsilon_F + \mu_0 H$, $s = +1$ convert into down electrons and go into the empty states at $\epsilon_F - \mu_0 H$, $s = -1$.

The ground state will instead look like



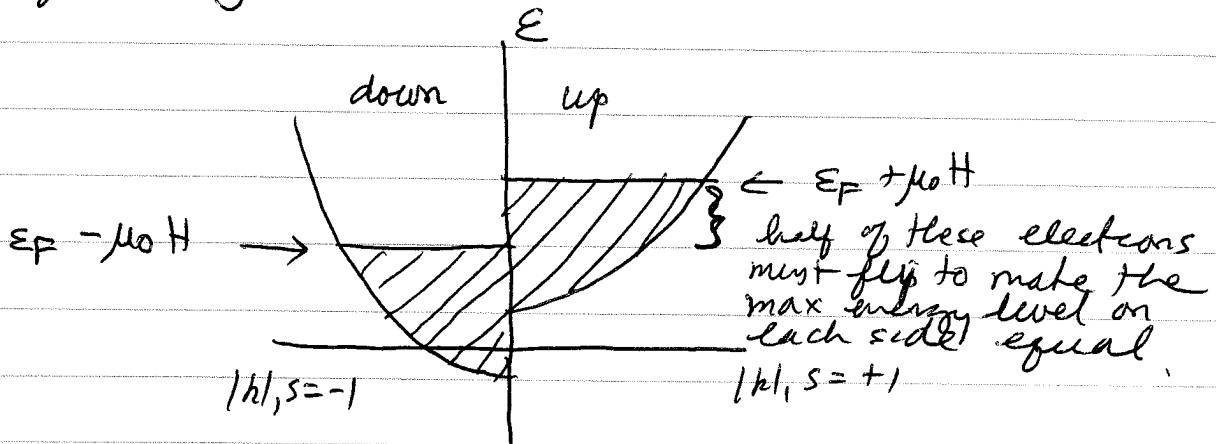
and we thus see that there will be more down than up electrons in the ground state.

Since $s = \pm 1$ electrons have magnetic moment $\mu = -\mu_0 s$, the system has a net positive magnetization M aligned with H .

To see how big $\propto M$ we need to compute the number of up electrons that flip into down electrons when H is turned on.

We will assume that H is small enough that $\mu_0 H \ll \epsilon_F$. When this is so, we can ignore the fact that the density of states has a slight variation with energy ϵ over the range $\epsilon_F + \mu_0 H$ to $\epsilon_F - \mu_0 H$ and assume it to be roughly constant $f(\epsilon_F)$.

The number of up electrons that flip is then easily computed from the following sketch



The number of up electrons that must flip is therefore

$$g_+(\epsilon_F) \Delta E$$

where $g_+(\epsilon_F) = \frac{1}{2} g(\epsilon_F)$ is the density of states of up electrons at ϵ_F , which is half the total density of states at ϵ_F , and $\Delta E = \frac{1}{2} [\mu_0 H - (-\mu_0 H)] = \mu_0 H$ is the energy interval that must flip. The number that flip is therefore

$$\Delta M = \frac{1}{2} g(\epsilon_F) \mu_0 H.$$

In the new ground state, the number of down electrons is now $n_0 + \Delta M$, and the number of up electrons is $n_0 - \Delta M$, where $n_0 = \frac{1}{2} M$ is the ~~not~~ number when $H=0$.

so the net magnetization density (at $T=0$) is then

$$\begin{aligned} M &= \mu_0 (m_- - m_+) \\ &= \mu_0 (m_0 + \Delta m - (m_0 - \Delta m)) \\ &= 2\mu_0 \Delta m \\ &= g(\epsilon_F) \mu_0^2 H \end{aligned}$$

and the Pauli paramagnetic susceptibility is

$$\chi_P = \frac{\partial M}{\partial H} = g(\epsilon_F) \mu_0^2$$

proportional to
density of states
at Fermi energy

For the free electron gas we have

$$g(\epsilon_F) = \frac{3}{2} \frac{m}{\epsilon_F}$$

$$\Rightarrow \boxed{\chi_P = \frac{3}{2} \frac{m}{\epsilon_F} \mu_0^2}$$

$$m \sim k_F^3, \epsilon_F \sim k_F^2, \text{ so } \chi_P \sim k_F \sim \frac{1}{(r_s/a_0)}$$

$$\chi_P = \frac{2.59}{(r_s/a_0)} \times 10^{-6}$$

corrections to above result at finite T of order

$\left(\frac{T}{T_F}\right)^2$ so above is very good ~~not~~ at all $T \ll T_F$
and so good at room temperature.

Compare to experiment

<u>metal</u>	<u>r_s/a_0</u>	<u>χ_p^{theory}</u>	<u>χ_p^{ext}</u>	$\times 10^{-6}$
Li	3.25	0.80	2.0	
Na	3.93	0.66	1.1	
K	4.86	0.53	0.8	
Rb	5.20	0.50	0.8	
Cs	5.62	0.46	0.8	

turns out that the discrepancy between theory and expt is mainly due to having neglected electron-electron interactions!

Note that χ_p above is very different from what one gets with classical statistics.

$$\text{Classically } M \sim \left[\frac{(+)e^{-\mu_0 H/k_B T} + (-)e^{+\mu_0 H/k_B T}}{e^{-\mu_0 H/k_B T} + e^{+\mu_0 H/k_B T}} \right] (-\mu_0) m$$

$$= \left[\frac{e^{\mu_0 H/k_B T} - e^{-\mu_0 H/k_B T}}{e^{\mu_0 H/k_B T} + e^{-\mu_0 H/k_B T}} \right] \mu_0 m$$

$$\sim \frac{\mu_0^2 H M}{k_B T} \quad \text{when } \mu_0 H \ll k_B T$$

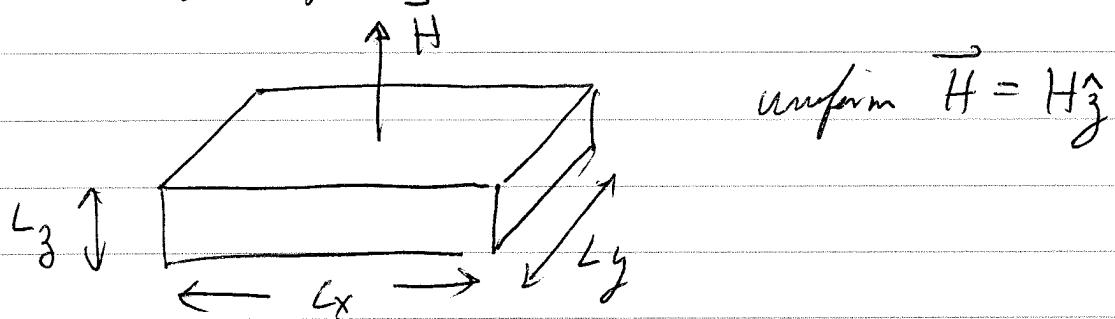
$$\chi_{\text{classical}} = \frac{dM}{dH} \sim \frac{\mu_0 m}{k_B T} \quad \text{Curie law } \sim \frac{1}{T}$$

$$\text{so } \frac{\chi_p}{\chi_{\text{classical}}} \sim \left(\frac{T}{T_F} \right) \ll 1$$

Landau Diamagnetism - Landau Levels

Here we wish to consider the effect of the magnetic field on the orbital motion of the conduction electrons. To do so we must solve the quantum mechanical problem of a charged particle moving in a uniform magnetic field.

The geometry we consider is



For a particle of charge q in a static uniform magnetic field, the Hamiltonian is

$$\mathcal{H} = \frac{1}{2m} \left(\frac{\hbar^2}{c} \vec{\nabla} - \frac{e}{c} \vec{A} \right)^2$$

where \vec{A} is the vector potential, $\vec{H} = \vec{\nabla} \times \vec{A}$
 $q = -e$ is the charge of the electron

for $\vec{H} = H\hat{z}$ we will use $\vec{A} = -\frac{q}{2} H \hat{x}$

Substitute these into \mathcal{H} to get:

$$\mathcal{H} = \frac{1}{2m} \left(\frac{\hbar}{i} \vec{\nabla} - \frac{e}{c} \vec{y} \vec{H} \right)^2$$

$$= \frac{1}{2m} \left[-\hbar^2 \frac{\partial^2}{\partial z^2} - \hbar^2 \frac{\partial^2}{\partial y^2} + \left(\frac{\hbar}{i} \frac{\partial}{\partial x} - \frac{e}{c} H_y \right)^2 \right]$$

We want to find the eigenstates ψ that solve

$$\mathcal{H}\psi = \varepsilon\psi$$

ε is eigenvalue of energy

try solution of the form

$$\psi(x, y, z) = e^{ikx} e^{ikz} \phi(y)$$

This form is suggested as \mathcal{H} is translationally invariant in x and z , but not in y (due to our particular choice for \vec{A})

Substitute this ψ into above Schrödinger Equation to get

$$\frac{1}{2m} \left[\hbar^2 k_z^2 - \hbar^2 \frac{\partial^2}{\partial y^2} + \left(\hbar k_x - \frac{e}{c} H_y \right)^2 \right] \phi(y) = \varepsilon \phi(y)$$

or

$$-\frac{\hbar^2}{2m} \frac{\partial^2 \phi}{\partial y^2} + \frac{1}{2m} \left(\hbar k_x - \frac{e}{c} H_y \right)^2 \phi = \left(\varepsilon - \frac{\hbar^2 k_z^2}{2m} \right) \phi$$

define y_0 such that $\hbar k_x \equiv \frac{eH}{c} y_0$

$$-\frac{\hbar^2}{2m} \frac{\partial^2 \phi}{\partial y^2} + \frac{1}{2m} \left(\frac{eH}{c} \right)^2 (y - y_0)^2 \phi = \left(\varepsilon - \frac{\hbar^2 k_z^2}{2m} \right) \phi$$

define cyclotron frequency $\omega_c \equiv \frac{eH}{mc}$

(a classical charged particle in uniform \vec{H} moves in a circular orbit with angular velocity ω_c)

Finally we get

$$\left[-\frac{\hbar^2}{2m} \frac{\partial^2}{\partial y^2} + \frac{1}{2} m \omega_c^2 (y - y_0)^2 \right] \phi(y) = \left(\epsilon - \frac{\hbar^2 k_z^2}{2m} \right) \phi(y)$$

This is just the Hamiltonian for a single harmonic oscillator of frequency ω_c that is centered at $y = y_0$.

We know the eigenvalues of energy of the harmonic oscillator are just

$$\hbar \omega_c (n + \frac{1}{2}) \quad n = 0, 1, 2, \dots$$

so we then have

$$\epsilon - \frac{\hbar^2 k_z^2}{2m} = \hbar \omega_c (n + \frac{1}{2})$$

or the energy eigenvalues of our particle are

$$\epsilon = \underbrace{\frac{\hbar^2 k_z^2}{2m}}_{T_z} + \underbrace{\hbar \omega_c (n + \frac{1}{2})}_{T_{xy}}$$

T_z kinetic energy of motion along \vec{z}
 T_{xy} kinetic energy of orbital motion in xy plane \perp to \vec{H}

The wavefunctions $\phi_n(y)$ are the usual harmonic oscillator wavefunctions (gaussian \times Hermite polynomial) only centered at y_0 .

We can therefore write our solution in terms of 3 quantum numbers, k_x, k_z, n

$$\left\{ \begin{array}{l} \psi_{k_x, k_y, n}(x, y, z) = e^{ik_x x} e^{ik_z z} \tilde{\Phi}_n(y - y_0) \\ E(k_x, k_y, n) = \frac{\hbar^2 k_y^2}{2m} + \hbar \omega_c (n + 1/2) \end{array} \right.$$

where $y_0 = \hbar k_x \frac{c}{eH} = \frac{\hbar k_x}{m \omega_c}$

Note E is independent of k_x so for fixed k_z and n there are many degenerate states corresponding to the different possible choices for k_x .

What are the possible values of k_x ?

If we take periodic boundary conditions along x ,

$$\psi(x + L_x, y, z) = \psi(x, y, z) \text{ then we must have}$$

$$e^{ik_x x} = e^{ik_x (x + L_x)} \Rightarrow k_x = \frac{2\pi}{L_x} \times (\text{integer})$$

But k_x also determines the value of y_0 about which the wavefunction is centered in the y -direction therefore we must have

$$0 \leq y_0 \leq L_y \Rightarrow 0 \leq \frac{\hbar k_x}{m \omega_c} \leq L_y$$

$$\Rightarrow k_{x \max} = L_y \frac{m \omega_c}{\hbar} = \frac{L_y m e H}{\hbar m c} = \frac{L_y e H}{\hbar c}$$