

14/05/19

4. de Haas - van Alphen effect.

→ allows us to measure Fermi surface which is crucial for various transport and electronic properties of the material.

→ In 1930 de Haas & van Alphen measured the magnetization of bismuth as a function of magnetic field at 4.2 K and found oscillatory behaviour.

Refs: Ashcroft and Mermin.

Block electrons in uniform magnetic field

semiclassical equations of motion (EOM) are given as

$$\dot{\mathbf{r}} = \mathbf{v}(\mathbf{k}) = \frac{1}{\hbar} \frac{\partial E(\mathbf{k})}{\partial \mathbf{k}}$$

$$\hbar \dot{\mathbf{k}} = -\frac{e}{c} \mathbf{v}(\mathbf{k}) \times \mathbf{H}$$

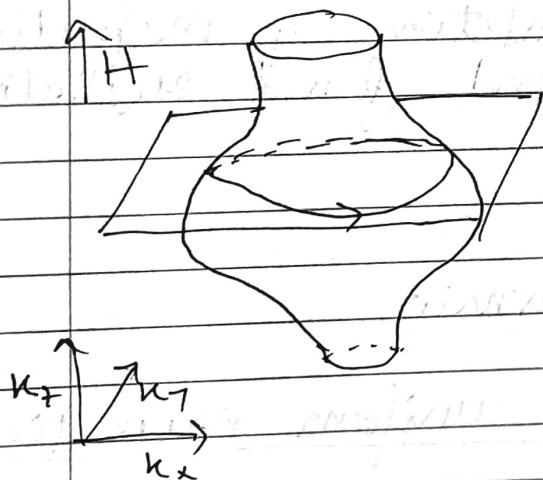
$$\Rightarrow (\dot{\mathbf{k}})_{\parallel} = 0 \Rightarrow (\mathbf{k})_{\parallel} = \text{const.}$$

component of $\mathbf{k}$ in the direction of the magnetic field.

and

$$\frac{d\epsilon(\mathbf{k})}{dt} = \frac{\partial \epsilon(\mathbf{k})}{\partial \mathbf{k}} \cdot \dot{\mathbf{k}} = \frac{-e}{c} \mathbf{v}(\mathbf{k}) \cdot [\mathbf{v}(\mathbf{k}) \times \mathbf{H}] = 0$$

This shows both $(\mathbf{k})$ and $\epsilon(\mathbf{k})$ are constants of motion.



Electrons will move along curves in the plane perpendicular to the magnetic field at constant energy.

→ The orbital period (T) of the electron can be calculated as

$$t_2 - t_1 = \int_{t_1}^{t_2} dt = \int_{k_1}^{k_2} \frac{dk}{|\dot{\mathbf{k}}|}$$

(plug in $E = 0$)

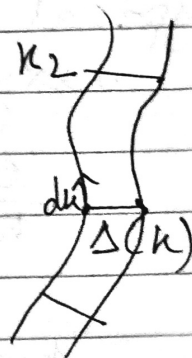
$$= \frac{\hbar c}{eH} \int_{k_1}^{k_2} \frac{dk}{\left| \left(\frac{\partial \epsilon}{\partial \mathbf{k}} \right)_\perp \right|}$$

↑
component $\perp$ to the magnetic field.

→ geometrical interpretation of $\left(\frac{\partial \epsilon}{\partial k}\right)_\perp$

$$\Delta \epsilon = \left(\frac{\partial \epsilon}{\partial k}\right)_\perp \Delta(k)$$

Since $\left(\frac{\partial \epsilon}{\partial k}\right)_\perp$ is $\perp$ to sec k_1



and $\perp$ to the orbit, it is $\parallel$ to $\Delta(k)$

So,

$$\Delta \epsilon = \left| \left(\frac{\partial \epsilon}{\partial k}\right)_\perp \right| \Delta(k)$$

$$t_2 - t_1 = \frac{\hbar^2 c}{eH} \frac{1}{\Delta \epsilon} \int_{k_1}^{k_2} \Delta(k) dk$$

in the limit $\Delta \epsilon \rightarrow 0$ $T = \frac{\hbar^2 c}{eH} \frac{\partial A_{1,2}}{\partial \epsilon}$

for a closed orbit we get

$$T(\epsilon, k_z) = \frac{\hbar^2 c}{eH} \frac{\partial}{\partial \epsilon} A(\epsilon, k_z)$$

Onsager's formula

→ typical quantum number in metals is very high

$$\nu = \frac{E_F}{\hbar \omega_c} = \frac{E_F}{\frac{e\hbar}{mc} \cdot H}, \quad \frac{e\hbar}{mc} \sim 10^{-8} \text{ cm/g}$$

so for typical metal with $E_F \sim 1 \text{ eV}$, even for 1 T or 10^4 G
 $\nu \sim 10^4$.

Therefore reasonable to use Bohr's correspondence principle

$$\Delta E_n = \frac{h}{T}, \quad T \text{ is the period of semi-classical motion.}$$

$$\rightarrow E_{\nu+1}(k_z) - E_{\nu}(k_z) = \frac{h}{T(E_{\nu}, k_z)}$$

$$\Rightarrow (E_{\nu+1} - E_{\nu}) \frac{\partial A(E_{\nu})}{\partial E} = \frac{2\pi e H}{\hbar c}$$

$$A(E_{\nu+1}) - A(E_{\nu}) = \boxed{\frac{2\pi e H}{\hbar c} = \Delta A}$$

using Bohr-Sommerfeld quantization condition

$$A(E_{\nu}, k_z) = (\nu + \gamma) \Delta A$$

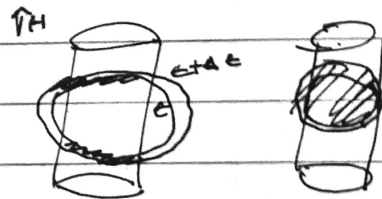
Origin of Oscillation

$$A = (\nu + \gamma) \frac{2\pi e H}{hc}$$

→ oscillatory structure in the electronic density of levels imposed by quantization condition.

→ For an orbit with extremal area A_e , a small variation of H will have large change in area.

$$\Delta\left(\frac{1}{H}\right) = \frac{2\pi e}{hc} \frac{1}{A_e(t_F)}$$



Conditions to observe dHvA

1. low T . $k_B T < \hbar \omega_c = \frac{\hbar e H}{m^* c}$
large magnetic field.

2. finite scattering time mixes different Lls, hence
 $\omega_c \tau > 1$.