

10878 Physik der Kondensierten Materie  
Summary

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# Introduction

## On this document

This document are my personal notes on the lecture 10878 Physik der Kondensierten Materie at Uni Basel in the Fall semester 2020. I type this document during the lectures (while the Professors speak) with minimal editing between lectures - there are A LOT of spelling mistakes, typos and errors of various severity.

## Information about this Course

all lectures online, exercises online Tue 8:15 to 10:00 in two groups

[nanophononics.physik.unibas.ch/pages/teaching.htm](https://nanophononics.physik.unibas.ch/pages/teaching.htm) and adam will hold the presentations

Exercises 30 points per sheet 60% of points required, hand in on thursday till 12am before lectures. Released on Friday. Links for exercise class will be sent by assistants.

Written exam on 20.01.2021 10:00 - 12:00, 5 pages double a4 handwritten notes and non-programmable calculator. Exam is based on exercise sheets.

Course mostly based on The Oxford Solid State Basics, Steven H. Simon. Some parts based on other books.

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# Chapter 1

## Heat in Solids

### 1.1 Introduction

What is condensed Matter physics and why is it important?

We deal in solids and liquids Condensed: interactions between atoms win over entropy. Key question is how to reproduce the properties of matter which is a many body problem. Currently the largest subfield of Physics.

Intro question: *Which keeps coffee warmest?*

- Steel Cup
- Ceramic Cup
- Glass Cup

The Steel cup loses, electrical conductors are also good thermal conductors. Glass actually has the lowest thermal conductivity, yet the ceramic cup still wins. Why?

It's not only the thermal conductivity that matters, it's also the heat capacity that matters, and in this regard ceramics are almost unbeatable.

### 1.2 Heat Capacity - Definitions

Heat Capacity is how much heat can you store in a material per change in temperature.

$$C = \frac{\partial U}{\partial T}.$$

Specific heat  $c_v$  is the heat capacity per unit volume.  $C_{N_a}$  is the heat capacity per mole, as it's otherwise per atom. We will only deal with the Heat Capacity at constant pressure and volume.

$$C_P \approx C_V = C.$$

We will now derive the Heat Capacity in a solid from the Heat Capacity in a Gas

$$\frac{C_V}{N} = \frac{3}{2} K_B \text{ for a mono-atomic gas.}$$

Multiplying by  $N_A$  yields:  $C_V = \frac{3}{2} R$ .

## 1.3 Dulong-Petit

Purely observational law for many solids:

$$C = 3K_B \text{ per atom}$$

$$C = 3R.$$

Works pretty well for many materials at room temperature (notable exception is Diamond).

Where did the law for Gas come from? Equipartition system, each degree of freedom gets equal amount of the internal energy  $\frac{1}{2}K_B T$ . In a Gas there are 3 degrees of Freedom (three elements of momentum).

For solids there are an additional 3 degrees of freedom from the potential energy (positional degrees of freedom)

3 degrees of freedom from kinetic energy + 3 degrees of freedom from potential energy = 6 degrees of freedom.

$$U = \frac{3+3}{2} K_B T.$$

$$C = \frac{\partial U}{\partial T} = 3K_B \text{ per atom.}$$

$$\implies C = 3R \text{ per mole.}$$

This 'law' fails for very low temperatures for all materials, and for some (notably diamond) it even fails completely.

## 1.4 Einstein Model

Assumes the particles are oscillating around their positional groundstate, and uses the expectation value of the potential energy instead of its' groundstate energy.

We assume the atoms are in a harmonic potential well and behave like harmonic oscillators with energy eigenstates  $E_n = \hbar\omega \left(n + \frac{1}{2}\right)$ . (Einstein did not include the factor  $+\frac{1}{2}$  as QM was not yet discovered).

We introduce the *Partition Function*  $Z$ , which describes the distribution among all possible microstates of a system. This function is extremely important in statistical mechanics and can be used to derive most macroscopic properties. It is given by:

$$Z = \sum_{n=0}^{\infty} \exp(-\beta E_n).$$

With  $\beta = \frac{1}{K_B T}$ .

We can use the partition function for example to calculate the probability to find a system in a certain microstate as  $P(E_n) = \frac{\exp(-\beta E_n)}{Z}$ . We can thus write the expectation value of the energy as

$$\langle E \rangle = \sum_{n=0}^{\infty} P(E_n) E_n = \frac{1}{Z} \sum_{n=0}^{\infty} E_n \exp(-\beta E_n).$$

Looking at this result we realize that

$$\begin{aligned} \frac{\partial Z}{\partial \beta} &= \sum_{n=0}^{\infty} -E_n \exp(-\beta E_n) \\ \implies \langle E \rangle &= -\frac{1}{Z} \frac{\partial Z}{\partial \beta}. \end{aligned}$$

This is the more general form of the energy expectation in terms of the partition function.

For the harmonic oscillator where  $E_n = \hbar\omega \left(n + \frac{1}{2}\right)$  we therefore get:

$$\begin{aligned} Z &= \sum_{n=0}^{\infty} \exp\left(-\beta \hbar\omega \left(n + \frac{1}{2}\right)\right) \\ &= \exp\left(-\frac{\beta \hbar\omega}{2}\right) * \sum_{n=0}^{\infty} \exp(-\beta \hbar\omega n) = \exp\left(-\frac{\beta \hbar\omega}{2}\right) * \sum_{n=0}^{\infty} \exp(-\beta \hbar\omega n) \end{aligned}$$

we recognize a geometric series and rewrite

$$Z = \exp\left(-\frac{\beta\hbar\omega}{2}\right) * \frac{1}{1 - \exp(-\beta\hbar\omega)} = \frac{1}{\exp\left(\frac{\beta\hbar\omega}{2}\right) + \exp\left(-\frac{\beta\hbar\omega}{2}\right)} = \frac{1}{2 \sinh\left(\frac{\beta\hbar\omega}{2}\right)}.$$

For the energy expectation value we thus get

$$\langle E \rangle = -\frac{1}{Z} \frac{\partial Z}{\partial \beta} = -\frac{\hbar\omega}{2} \coth\left(-\frac{\beta\hbar\omega}{2}\right).$$

Introducing the *Bose occupation factor*  $n_B(x) = \frac{1}{\exp(x)-1}$  we get

$$\langle E \rangle = \hbar\omega \left( n_B(\beta\hbar\omega) + \frac{1}{2} \right).$$

In this calculation we have only considered a one-dimensional harmonic oscillator, in reality we have to consider the three-dimensional case. In three dimensions the expectation value for the energy is three times higher

$$\langle E_{3d} \rangle = 3 \langle E_{1d} \rangle = 3\hbar\omega \left( n_B(\beta\hbar\omega) + \frac{1}{2} \right).$$

According to definition the Heat capacity is given by

$$c = \frac{\partial \langle E \rangle}{\partial T} = K_B (\beta\hbar\omega)^2 \frac{\exp(\beta\hbar\omega)}{[\exp(\beta\hbar\omega) - 1]^2}.$$

We will now investigate this new expression in the limit for very big and very small temperatures:

$$\lim_{T \rightarrow \infty} c = \lim_{\beta \rightarrow 0} c = 3K_B \text{ for big temperatures we recover the Dulong-Petit Law as expected}$$

$$\lim_{T \rightarrow 0} c = \lim_{\beta \rightarrow \infty} c = 0 \text{ for very small Temperatures the model predicts an exponential decay.}$$

This approximation models all materials at high and low temperatures, but still fails at very low temperatures, where it predicts an exponential decay, where in experiments a decay  $\propto T^3$  is measured.

## 1.5 Debye Model of Solids

Assumes that atoms at the bottom of their potential well oscillate not independently from each other, but are coupled in a collective motion. Similar to sound waves.

QM was not yet discovered, but the quantisation of light was known, Debye thus applied that same line of thinking to sound waves. It's important to note, that sound waves, contrary to light waves can also have a longitudinal part. We will treat waves in a finite solid with periodic boundary conditions. We define a periodic space on which there is a periodicity such that all waves are same at  $x = 0$  and  $x = L$ . Thus only wavelengths of the form

$$\lambda_m = \frac{2L}{m} \text{ are allowed.}$$

The Wavelengths of the 'allowed' waves are given by

$$K = \frac{2\pi m}{L}.$$

Whenever we want to treat the sum over 'all possible' values of  $k$  we can replace the sum over all  $k$  with an integral (provided that  $L$  is big)

$$\sum_k \rightarrow \frac{L}{2\pi} \int_{-\infty}^{\infty} dk.$$

In three dimensions this integral is over the three dimensional  $k$ -space

$$\sum_k \rightarrow \left( \frac{L}{2\pi} \right)^3 \int d^3k.$$

Debye assumed that the oscillation modes in a solid behave like sound waves, and therefor have a linear dispersion relation

$$\omega(k) = v|k|.$$

The only adaptation to Einstein we have to do, is to integrate over all possible  $k$ .

$$\begin{aligned} \langle E \rangle &= 3 \sum_k \hbar\omega(\vec{k}) \left( n_B(\beta\hbar\omega(\vec{k})) + \frac{1}{2} \right). \\ \implies \langle E \rangle &= 3 \frac{L^3}{(2\pi)^3} \int \hbar\omega(\vec{k}) \left( n_B(\beta\hbar\omega(\vec{k})) + \frac{1}{2} \right) d\vec{k}. \end{aligned}$$

By switching to spherical coordinates we get  $\int d\vec{k} \rightarrow 4\pi \int_0^\infty k^2 dk$

$$\implies \langle E \rangle = \frac{3(4\pi)L^3}{(2\pi)^3} \int_0^\infty k^2 \hbar\omega \left( n_B(\beta\hbar\omega) + \frac{1}{2} \right) dk.$$

We now use the dispersion relation and insert  $\omega = vk$

$$\begin{aligned} \implies vdk &= d\omega \text{ and } k^2 = \frac{\omega^2}{v^2}. \\ \implies \langle E \rangle &= 3 \frac{4\pi L^3}{(2\pi)^3} \int_0^\infty \omega^2 \frac{1}{v^3} \hbar\omega \left( n_B(\beta\hbar\omega) + \frac{1}{2} \right) d\omega. \end{aligned}$$

If a solid has  $n$ -atoms per unit volume then a cube of side  $L$  has a total of  $N$  atoms.  $\rightarrow N = nL^3 \implies L^3 = \frac{N}{n}$ .

$$\implies \langle E \rangle = \int_0^\infty \left( \frac{12\pi\omega^2 N}{(2\pi)^3 n} \right) \frac{1}{v^3} \hbar\omega \left( n_B(\beta\hbar\omega) + \frac{1}{2} \right) d\omega.$$

If we define the density of States as

$$g(\omega) = N \left( \frac{12\pi\omega^2}{(2\pi)^3 nv^3} \right).$$

We can rewrite the energy expectation as

$$\langle E \rangle = \int_0^\infty g(\omega) \hbar\omega \left( n_B(\beta\hbar\omega) + \frac{1}{2} \right) d\omega.$$

The density of states  $g(\omega)$  is a factor weighing ow many states that there are at a given frequency. The Integral is thus just equal to the density of the states times the energy of the states at a given frequencyp.

If we further define a Debye frequency  $\omega_D = 6\pi^2 nv^3$  we can further simplify the density  $g(\omega) = N \frac{9\omega^2}{\omega_D^3}$ .

The final result for the energy expectation is thus:

$$\langle E \rangle_{tot} = \frac{9N\hbar}{\omega_D^3} \int_0^\infty \frac{\omega^3}{\exp(\beta\hbar\omega) - 1} d\omega.$$

Note: we have left out the factor of  $\frac{1}{2}$ , because we will differentiate later and it'd fall out anyway (also leaving it in creates unnecessary mathematical difficulties).

To solve the integral we make a change of variables

$$x = \beta\hbar\omega \rightarrow \frac{dx}{\beta\hbar} = d\omega.$$

The solution is still very tedious, the final result is given by

$$\langle E \rangle = 9N \frac{(K_B T)^4}{(\hbar\omega_D)^3} \left( \frac{\pi^4}{15} \right).$$

We therefore get

$$C = \frac{\partial \langle E \rangle}{\partial T} = \dots \propto T^3.$$



Debye thus correctly models the low temperature regime ( $T^3$  dependence) but fails for high temperatures, and does not converge to the Dulong-Petit.

The problem of the Debye model is that we were integrating over an infinite number of modes, but in reality there can never be an infinite amount of modes.

$$\text{No. of modes} = \int_0^\infty g(\omega) d\omega \rightarrow \infty.$$

Debye thus introduced a 'cutoff frequency' such that the total number of modes are equal to the total number of degrees of freedom

$$3N = \int_0^{\omega_{cutoff}} g(\omega) d\omega.$$

With this correction we get

$$\langle E \rangle = \int_0^{\omega_c} g(\omega) \hbar \omega n_B(\beta \hbar \omega) d\omega.$$

At high temperatures we thus get:

$$\begin{aligned} T \rightarrow \infty &\implies n_B(\beta \hbar \omega) \rightarrow K_B \frac{T}{\hbar \omega}. \\ \implies \langle E \rangle &= K_B T \int_0^{\omega_c} g(\omega) d\omega = 2K_B T N. \\ \implies C &= 3NK_B = 3R \text{ (Dulong-Petit!)}. \end{aligned}$$

It's a bit ad-hoc, but it works.

We actually find that the cutoff frequency is equal to the Debye frequency

$$\omega_{cutoff} = \omega_D.$$

The final result of Debye is thus:

$$\langle E \rangle_{tot} = \int_0^{\omega_c} g(\omega) \left( n_B(\beta \hbar \omega) + \frac{1}{2} \right) d\omega.$$

The Debye model is the first model that does not feature a fitting parameter. Unfortunately there are still some problems:

- $\omega_{cutoff}$  is an unexplained ad hoc correction
- The dispersion relation used  $\omega = vk$  does not hold at high temperature
- There are still some exotic materials that are not modelled correctly
- For metals the heat capacity  $c$  at low temperatures is different:  $c \propto \alpha T^3 + \gamma T$ .  
Only the  $T^3$  dependence is modeled correctly by the Debye model.

# Chapter 2

## Drude Theory

Developed three years after the discovery of electrons. To accurately describe metals we have to take the electron  $e^-$  into account.

Drude applies the kinetic theory of Boltzmann to electrons, a few assumptions are necessary:

- There are collisions among electrons with a scattering time  $\tau$ . The probability of scattering is given by  $\frac{\delta t}{\tau}$
- After scattering the momentum  $\vec{p}_{final} = 0$ . This obviously only holds on average across many collisions
- If there are electric and magnetic fields  $\vec{E}$   $\vec{B}$  the electrons will interact with them

While assumptions 1 and 3 are also valid in gasses, the third assumption obviously only holds for electrons.

### 2.1 Relaxation Time Approximation

We consider an electron traveling through a crystal with momentum  $\vec{p}$ . After some time a scattering event occurs, after this event it has a new momentum  $\vec{p}(t + \delta t)$ . We first consider all the electrons that collide in the time  $\delta t$ . The probability is given as  $\frac{\delta t}{\tau}$ , according to the second assumption, those electrons will end up with  $\vec{p} = 0$ .

Electrons that don't collide. The probability of no collision is given as  $1 - \frac{\delta t}{\tau}$ . When an electron does not collide, it will accelerate due to the force acting on it  $\vec{F} = m_e a = \frac{d\vec{p}}{dt}$ . The force can naturally vary with time  $\rightarrow \vec{F} + O(t)$ .

$$\implies \delta \vec{p} = \vec{F} \delta t + O(\delta t^2).$$

What we search is  $\vec{p}(t + \delta t)$ .

$$\vec{p}(t + \delta t) = \frac{\delta t}{\tau} \vec{p}_{final} + \left(1 - \frac{\delta t}{\tau}\right) \left[\vec{p}(t) + \vec{F}(t) \delta t + O(\delta t^2)\right].$$

The first term is 0 because of assumption 2

$$\begin{aligned} \implies \vec{p}(t + \delta t) &= \left(1 - \frac{\delta t}{\tau}\right) \left[\vec{p}(t) + \vec{F}(t) \delta t\right] = \vec{p}(t) + \vec{F}(t) \delta t - \frac{\delta t}{\tau} \vec{p}(t). \\ \implies \frac{d\vec{p}(t)}{dt} &= \vec{F} - \frac{\vec{p}(t)}{\tau}. \end{aligned}$$

This result is called the relaxation time equation of the Drude model. We know that

$$\vec{F} = -e \left( \vec{E} + \vec{v} \times \vec{B} \right).$$

The other term  $\vec{p}/\tau$  can be understood as a kind of drag on the electrons.

As an example we consider  $\vec{F} = 0$  and get

$$\frac{d\vec{p}(t)}{dt} = -\frac{\vec{p}(t)}{\tau} \implies \vec{p}(t) = \vec{p}_0 \exp\left(-\frac{t}{\tau}\right).$$

On average the momentum  $\vec{p}$  tends to zero (due to scattering) in absence of no external force  $\vec{F}$ .

## 2.2 $e^-$ in electric fields - Drude conductivity

We recall ohm's law  $\vec{j} = \sigma \vec{E}$ . We assume that we have  $\vec{E} \neq 0$  but  $\vec{B} = 0 \implies \vec{F} = -e\vec{E}$ . We thus get

$$\frac{d\vec{p}}{dt} = -e\vec{E} - \frac{\vec{p}}{\tau}.$$

We are interested in a steady state  $\frac{d\vec{p}}{dt} = 0$

$$\implies \vec{p} = -e\tau \vec{E} = m\vec{v}.$$

$$\implies \vec{v} = -\frac{e\tau}{m_e} \vec{E}.$$

We now use ohm's law with current density  $\vec{J} = -ne\vec{v}$ , with the electron density  $n$ .

$$\vec{J} = -ne\vec{v} = -ne \left( -\frac{e\tau}{m_e} \vec{E} \right) = \frac{ne^2\tau}{m_e} \vec{E} = \sigma \vec{E}.$$

$$\implies \sigma_{drude} = \frac{ne^2\tau}{m_e}.$$

The conclusion of this part, is that if you can measure  $\sigma_{drude}$  you can use it to measure the electron density in a material.

## 2.3 $e^-$ in $\vec{E}$ and $\vec{B}$ - Hall effect in the Drude model

$$\vec{F} = -e \left( \vec{E} + \vec{v} \times \vec{B} \right).$$

$$\implies \frac{d\vec{p}}{dt} = -e \left( \vec{E} + \vec{v} \times \vec{B} \right) - \frac{\vec{p}}{\tau}.$$

Again we are interested in the steady state.

$$-e\vec{E} - e\vec{v} \times \vec{B} - \frac{\vec{p}}{\tau} = 0.$$

$$\implies -e\vec{E} + \frac{\vec{J} \times \vec{B}}{n} + \frac{m_e}{ne\tau} \vec{J} = 0.$$

We assume that  $\vec{B} = -B_z \vec{e}_z$ . We then take a piece of metal with dimensions  $L_x$ ,  $L_y$ , and  $L_z$ , and run a current through the metal along the  $X$ -axis  $\implies \vec{J} = J_x \vec{e}_x$ , thus we also know that  $\vec{E} = E_x \vec{e}_x + E_y \vec{e}_y$  will only have components in the  $X/Y$ -Plane.

$$\text{X component: } eE_x = 0 + \frac{m_e}{ne\tau} J_x \implies E_x = \frac{m_e}{ne^2\tau} J_x$$

$$\text{y component: } eE_y = \frac{J_x B_z}{n} + 0 \implies E_y = \frac{B_z J_x}{ne}.$$

$$\text{z component: } 0 = 0 + 0.$$

We know that  $\vec{J} = \sigma \vec{E} = \frac{1}{\rho} \vec{E}$  but we now see that the resistance  $\rho$  depends on the direction (aka. is a matrix).

$$\rho_{xx} = \frac{E_x}{J_x} = \frac{ne^2\tau}{m_e}$$

$$\rho_{xy} = \frac{E_y}{J_x} = -\frac{B}{ne}$$

The really important coefficient is  $\rho_{xy}$  (the longitudinal resistivity  $\rho_{xx}$  is expected). We call  $\rho_{xy}$  the Hall resistivity, which means that there is a voltage along  $Y$  that we call Hall voltage.

The Hall effect thus is the effect, that when applying a magnetic field perpendicular to a current density we suddenly measure a voltage perpendicular to both. This effect is used for a variety of measurements.

We define the Hall coefficient:

$$R_H = \frac{\rho_{xy}}{|B|} = -\frac{1}{ne}.$$

How can the Hall effect can actually be used?

- Hall sensors to measure magnetic fields  $\vec{B}$ .

If  $n$ ,  $e$  and  $\vec{J}$  are known we can use the Hall effect to measure a magnetic field. Note: High Hall voltages are easier to measure, for such sensors we want to choose materials with a big Hall coefficient  $R_H = -\frac{1}{ne}$ , aka. with small  $n$  (semiconductors).

- If we know  $\vec{B}$  and  $\vec{J}$  we can measure the electron density.

## 2.4 Heat Capacity due to $e^-$

We study a cylinder that we are heating on one end. There will be a heat flow  $\vec{Q}$  and a heat flux, or heat current density,  $\vec{J}_Q = \frac{\text{energy}}{\text{time} \cdot \text{area}}$ .

$$\vec{J}_Q = -\kappa \nabla T.$$

Drude assumes that the heat in metals is only transported by the electrons, and he describes electrons with the kinetic theory of gasses.

$$\kappa = \frac{1}{3} n C \langle v \rangle \lambda = \frac{1}{2} n c \langle v \rangle^2 \tau.$$

With  $\lambda = \langle v \rangle \tau$  the distance an electron will travel before scattering. From the last Lecture we know that  $\frac{C}{N} = \frac{3}{2} K_B$ . We also know an expression for the velocity  $\langle v^2 \rangle = \frac{3 K_B T}{m_e}$

$$\Rightarrow \kappa = n \frac{3}{2} \frac{K_B^2 T}{m_e} \tau.$$

Note: we assumed  $\langle v^2 \rangle = \langle v \rangle^2$  which does not hold generally and will lead to problems later on. The question now is: how do we find  $\tau$ ?

From earlier we know  $\sigma_{Drude} = \frac{ne^2\tau}{m_e}$  we thus consider

$$\text{Lorenz number: } L_{Drude} = \frac{\kappa}{\sigma T} = \frac{3}{2} \left( \frac{K_B}{e} \right)^2 \frac{1}{T}.$$

We call the ratio between  $\kappa$  and  $\sigma$  the Wiedeman-Franz ratio

$$\frac{\kappa}{\sigma} = \frac{3}{2} \left( \frac{K_B}{e} \right)^2.$$

The Wiedeman-Franz law states that

$$LT = \frac{\kappa}{\sigma}.$$

Thus we know that in metals thermal conductivity  $\kappa$  and electrical conductivity  $\sigma$  are proportional to each other. The interesting thing is that the Lorenz number has another definition

$$L = \frac{\pi^2 K_B^2}{3e^2} \approx 2 * L_{Drude}.$$

Even with the factor of 2 the Drude model is still a success because previously it was unexplained why the Wiedeman-Franz law even holds (previously it was only known experimentally). The exact value of the Lorenz number given above comes from modern quantum mechanical calculations.

## 2.5 Thermoelectric Properties - how the Drude model fails

Conversion of Heat into electricity. Electrical currents always carry heat. But an electrical current is always heating the conductor via Joule heating  $I^2 R$ , for use in thermoelectric devices one wants to minimize  $R$ .

$$\Delta V = S \Delta T \text{ with } S \text{ the Seebeck coefficient.}$$

$$Q = \Pi I \text{ with } \Pi \text{ the Peltier coefficient.}$$

We get the relationship

$$\vec{J}_q = \Pi \vec{J}.$$

With  $\vec{J}_q$  the heat current density and  $\vec{J}$  the electrical current density. We already know:

$$\vec{J} = -en\vec{v}$$

$$\Rightarrow \Pi = \frac{J_q}{J} = -\frac{1}{3} \frac{C_v T}{e} = -\frac{C_v T}{3e} = -\frac{K_B T}{2e}.$$

There is also an expression for  $S$  the Seebeck coefficient

$$S = \frac{\Pi}{T} = -\frac{K_B}{2e} = -4.3 * 10^{-4} \frac{V}{K}.$$

But the real values of  $S$  are on the order of  $\propto 10^{-6} \frac{V}{K}$ . We now found a significant failure of the Drude system. But why did it work so well for the heat capacity but fails so badly for thermoelectrical effects?

Drude committed two errors, that cancelled each other out in the case of the heat capacity.

- He assumed  $\langle v \rangle^2 = 3 \frac{K_B T}{m_e} \gg$  actual  $\langle v \rangle^2$
- He assumed  $\frac{C_v}{N} = \frac{3}{2} K_B \gg$  the actual heat capacity

## 2.6 Success and Failures of the Drude model

### 2.6.1 Mean free path of $e^-$ - $\lambda$

$$\lambda = \langle v \rangle * \tau.$$

$$\sigma = \frac{ne^2 \tau}{m_e}, \quad \rho = \frac{1}{\sigma}.$$

$\rho$  can be measured in materials and can thus be used to measure  $\tau$ .

$$\Rightarrow \tau = \frac{m_e}{ne^2 \rho} \approx 1 - 10 \text{ fs}$$

$$\langle v \rangle \approx \sqrt{\frac{2K_B T}{m_e}}$$

$$\Rightarrow \lambda = \langle v \rangle \tau \approx 0.1 - 1 \text{ nm}.$$

Close to the lattice constants in materials  $\rightarrow$  success of Drude.

### 2.6.2 Measured Values

With the Hall effect we can measure the electron density

$$n = -\frac{1}{eR_H}.$$

We can now compare the values measured electron density to the theoretical values for the electron density. We call the density of atoms in a solid  $n_a$ . For monovalent atoms (with only one atom in the valence shell) we find

$$n_a = n.$$

We can infer a general quantity  $\alpha = \frac{n}{n_a}$ . For Valence = 1  $\Rightarrow \alpha = 1$ , for valence  $V \Rightarrow \alpha = V$ .

When we compare the known valence of various Metals to the theoretically measured  $\alpha$ 's we find a good match for Li or Ne (with a valence of  $V = 1$ ), but fails for elements with  $V > 1$ .

The Drude model works pretty well for monovalent atoms, but fails for higher valence atoms (sometimes even getting the sign of  $\alpha$  wrong).

Other successes:

- $\delta T$  respects the Wiedemann-Franz law (with only an error factor of 2)

Other failures:

- Seebeck coefficient is too big (by two orders of magnitude).

## Chapter 3

# Sommerfeld Theory for free electrons in metals

The mistake of Drude, was that he applied the kinetic theory of gasses (based on Bose-Einstein statistics) to electrons. He thus implicitly assumed electrons to be bosons, but they are fermions.

### 3.1 Fermi-Dirac Statistics

Historical context

- 1925: Pauli exclusion principle (two  $e^-$  cannot be in the exact same state)
- 1926: Schroedinger Equation and Fermi-Dirac statistics
- 1927: Sommerfeld applies FD statistics to metals

We have to define a Fermi occupation function  $n_F(\epsilon - \mu) = \frac{1}{\exp(\beta(\epsilon - \mu)) + 1}$ . It's the probability that the fermion occupies a state with energy  $\epsilon$ .  $\mu$  is the chemical potential, is defined by the Fermi occupation function. We use this Fermi occupation function  $n_F$  instead of the Bose-Einstein occupation function  $n_B$ .

$$n_B = \frac{g_i}{\exp\left(\frac{\epsilon_i - \mu}{k_B T}\right) - 1} \text{ with } g_i \text{ the states degeneracy.}$$

$$n_F = \frac{1}{\exp\left(\frac{\epsilon_i - \mu}{k_B T}\right) + 1}.$$

Fermi Statistics cannot allow for degeneracy, since fermions can *never* be in the exact same state. We have to differentiate Bosons (indistinguishable quantum particles) and Fermions (distinguishable quantum particles). Sommerfeld used FD and quantum mechanics to describe the electrons in a metal as a "Fermi-Gas". He does two upgrades to the Drude Model:

- calculates the energy quantum mechanically
- uses FD statistics

### 3.2 Quantum States of $e^-$ in solids

We consider a solid with regularly spaced atoms and look at the potential. The real potential is complicated so Sommerfeld made some approximations

- He neglected core electrons and considered only valence electrons.
- He replaced the true coulomb-potential with a constant, average, potential  $V_0$ .

- He ignored interactions between electrons.
- He assumes that the surface of the solid act like a barrier of height  $\infty$ .

With these approximations we can write the Schroedinger Equation in 1d:

$$\frac{\hbar^2}{2m_e} \frac{\partial^2}{\partial x^2} \psi(x) = V\psi(x)$$

Because of the assumption that the surface of the solid has a potential of  $V_{boundary} = \infty$  we get the boundary conditions

$$\begin{aligned} \psi(0) &= \psi(L) = 0 \\ \implies \psi(x) &= A \sin(k_x x). \end{aligned}$$

We get standing waves with  $k_x = \frac{2\pi}{L} n_x$  with  $n_x \in \mathbb{Z}$ . We get the Energy of a state as

$$E_n = \frac{\hbar^2 k_{x,n}^2}{2m_e} = \frac{\hbar^2 4\pi^2 n_x^2}{2m_e L^2}.$$

We have a discrete, quadratic potential. The energy entirely determined by the quantum number  $n_x$ . According to the Pauli exclusion principle each electron has to have unique quantum numbers. There are multiple quantum numbers, they are

- $n_x, n_y$ , and  $n_z$
- spin number  $n_s = \pm \frac{1}{2}$

We know that if two electrons have the same energystate (given by  $\vec{n}$ ) must have opposite spin. Every value of the three energy state numbers ( $\vec{n}$ ) can hold two electrons (one with spin 'up' and one with spin 'down').

### 3.3 Filling the $k$ -space with electrons

We now consider the  $K$ -space, in which only discrete values are allowed. We know that the spacing between the allowed values in the  $i$ -axis is given by  $\frac{2\pi}{L_i}$ , with  $L_i$  the length of the solid in the  $i$ -axis. The area of a discrete value in a two dimensional  $K$ -space is given by

$$A = \frac{2\pi}{L_x} \frac{2\pi}{L_y} = \frac{(2\pi)^2}{L_x L_y}$$

In 3d we get a volume

$$V = \frac{(2\pi)^3}{L_x L_y L_z}.$$

We consider a two dimensional system and lable the  $K$ -states

$$\begin{aligned} K_x &= \left(\frac{2\pi}{L_x}\right) n_x \quad K_y = \left(\frac{2\pi}{L_y}\right) n_y \\ E &= \frac{\hbar^2 \vec{K}^2}{2m_e} = \frac{\hbar^2}{2m_e} (K_x^2 + K_y^2) = \frac{\hbar^2 (2\pi)^2}{2m_e} \left( \frac{n_x^2}{L_x^2} + \frac{n_y^2}{L_y^2} \right). \end{aligned}$$

We thus have to fill the  $K$ -space from the center outwards, up to a certain maximum  $\vec{K}_F$ , that we call Fermi  $K$ -Vector. We have to choose  $\vec{K}_F$  such that we can fill the sphere in the  $K$ -space with radius  $|\vec{K}_F|$  with all of our electrons.

$$E = \frac{\hbar^2 K^2}{2m_e} \implies E_F = \frac{\hbar^2 \vec{K}_F^2}{2m_e} \text{ Fermi energy.}$$

We thus get a sphere containing allowed energy states, we call this sphere *Fermi Sphere*, the maximum allowed energy in this Fermi Sphere is the Fermi Energy  $E_F = \frac{\hbar^2 K_F^2}{2m_e}$ . No matter what dimensionality our problem has, we can define the Fermi Sphere and get the occupation probability

$$n_F = \frac{1}{\exp\left(\frac{E-E_F}{K_B T}\right) + 1}.$$

At  $T = 0$  this probability is a step-function with  $n_F = 1 \forall E < E_F$  and  $n_F = 0 \forall E > E_F$ . For any other temperatures the step function is 'smoothed' out and we get  $n_F = \frac{1}{2}$  at  $E = E_F$ .

### 3.3.1 Fermi-Energy, -Wavevector and -surface

We assume that there are  $N$  electrons in our system.

$$N = 2 * \sum_{\vec{k}} n_{FD}(E) = 2 \frac{V}{(2\pi)^3} \int n_{FD} d\vec{k}.$$

We know that at  $T = 0$  K the Fermi-Dirac function is a step function

$$n_{FD}(E) = 1 \forall E \leq E_F, 0 \text{ otherwise.}$$

$$\begin{aligned} \Rightarrow N &= 2 \frac{V}{(2\pi)^3} \int_0^{K \leq K_F} dK \\ &= \frac{2V}{(2\pi)^3} * \frac{4}{3} \pi K_F^3. \end{aligned}$$

We would like to know the electron density  $n = \frac{N}{V}$

$$\Rightarrow n = \frac{N}{V} = \frac{K_F^3}{3\pi^2}.$$

We can use this fact to easily calculate  $K_F$  and thus the Fermi-Energy from the electron density.

$$K_F = (3\pi^2 n)^{\frac{1}{3}}.$$

### 3.3.2 Fermi-Momentum and Velocity

We define the Fermi-velocity via:

$$\begin{aligned} \vec{P} &= \hbar \vec{K}_f = m_e \vec{v}_F \\ v_f &= \frac{\hbar}{m_e} (3\pi^2 n)^{\frac{1}{3}}. \end{aligned}$$

For typical electron densities we find  $v_F \approx 0.5$  to  $1\% C$  with  $C$  the speed of light. We know

$$E_F = \frac{\hbar^2 K_F^2}{2m_e} = \frac{\hbar^2}{2m_e} (3\pi^2 n)^{\frac{2}{3}}.$$

from this we find the Fermi-Temperature

$$T_F = \frac{E_F}{K_B}.$$

For typical metals the Fermi-Temperatures we find is somewhere between  $6 * 10^4$  K and  $10^5$  K.



### 3.3.3 Average Energy of an electron in a Fermi-Gas

To get the average energy we first calculate the total energy of our system:

$$\begin{aligned} E_{tot} &= 2 \int_0^{K \leq K_F} \frac{V}{(2\pi)^3} \frac{\hbar^2 k^2}{2m_e} d^3k \\ &= \frac{\hbar^2 V}{(2\pi)^3 m_e} \int_{06} K_f k^2 d\vec{K}. \end{aligned}$$

Substituting  $d\vec{K} = 4\pi k^2 dk$  (our  $\vec{K}$  are packed in a sphere, over which we integrate)

$$\begin{aligned} &= \frac{\hbar^2 V}{(2\pi)^3 m_e} 4\pi \int k^4 dk = \frac{\hbar^2 V}{2\pi^2 m_e} \frac{1}{5} K_F^5 \\ E_{tot} &= \frac{\hbar^2 V}{10\pi^2 m_e} K_F^5. \end{aligned}$$

Using

$$N = \frac{2V}{(2\pi)^3} \frac{4}{3} K_F^3$$

the average energy  $E_{avg}$  is given by

$$\frac{E_{tot}}{N} = \frac{3}{5} \frac{\hbar^2}{m_e} K_F^2 = \frac{3}{5} E_F.$$

## 3.4 Density of States

$g$  is the number of (quantum) states for electrons per unit volume and per unit energy. We can define the density of electrons at a porticular energy and temperature as

$$n(E, T) = \frac{g(E)}{V} f(E_F, T).$$

With  $f(E, T)$  the Fermi-Dicrac distribution.

As defined above  $g(k)$  is the number of  $k$ -states per total volume. We can now write

$$g(E)dE = g(k)d^3k = \left(\frac{L}{2\pi}\right)^3 d^3k$$

We also know already

$$\begin{aligned} E = \frac{\hbar^2 k^2}{2m_e} &\implies k = \sqrt{\frac{2m_e}{\hbar^2} E} \implies dk = \sqrt{\frac{2m_e}{\hbar^2} E} \frac{1}{2} dE. \\ &\implies g(E)dE = \left(\frac{L}{2\pi}\right)^3 4\pi k^2 dk \\ &= \left(\frac{L}{2\pi}\right)^3 4\pi \frac{2m_e E}{\hbar^2} \sqrt{\frac{2m_e}{\hbar^2} E} \frac{1}{2} dE \\ &\implies g(E) = \frac{V}{(2\pi)^2} \left(\frac{2m_e}{\hbar^2}\right)^{\frac{3}{2}} \sqrt{E}. \end{aligned}$$

$g(E)$  per volume is thus given by

$$\frac{g(E)}{V} = \frac{1}{(2\pi)^2} \left(\frac{2m_e}{\hbar^2}\right)^{\frac{3}{2}} \sqrt{E} = \frac{3}{2} \frac{n}{E_F} \sqrt{\frac{E}{E_F}}.$$

The key takeaways here are that  $g(E) \propto \sqrt{E}$  and that

$$g(E_F) = \frac{3}{2} \frac{n}{E_F}.$$

### 3.5 Electronic Heat Capacity

We now want to calculate the contribution of the electrons to the Heat Capacity. In a gas of electrons we would expect  $U_{el} = 3 * \frac{1}{2} K_B T \Rightarrow c_{el} = \frac{3}{2} K_B$  per electron. But in a fermi-gas only the electrons close to the Fermi-surface can actually "gain" internal energy and contribute to the heat capacity.

A fermigas will thus "accept" heat in a very different way. We consider the number of electrons

$$N = V \int_0^\infty g(E) n_F(\beta(E - \mu)) dE$$

$$E(T) = V \int_0^\infty g(E) E n_F(\beta(E - \mu)) dE$$

$$C(T) = \frac{\partial E(T)}{\partial T}.$$

This integral for  $E(T)$  is very hard and usually solved numerically. To solve it analytically we make a few assumptions:

- $\mu$  is independent from  $T$ . This only holds as long as we consider small Temperature changes
- $E(T) = E(T=0) +$  number of electrons that can be excited times amount of energy absorbed

These assumptions yield

$$E(T) = E(T=0) + \frac{\gamma}{2} V g(E_F) K_B T * K_B T$$

$$\Rightarrow c = \frac{\partial E}{\partial T} = \frac{\gamma}{2} V g(E_F) K_B^2 2T$$

Using  $g(E_F) = \frac{3}{2} \frac{n}{E_F}$  and  $V = \frac{N}{n}$

$$c = \gamma N \frac{3}{2} K_B \left( \frac{K_B T}{E_F} \right).$$

An accurate calculation yields  $\gamma = \frac{\pi}{3}$ . The final result of the heat capacity of the electrons is thus

$$c = \frac{\pi}{2} N K_B \left( \frac{K_B T}{E_F} \right).$$

The factor  $\frac{K_B T}{E_F}$  is usually extremely small, as the Fermi-Energy is usually high. For the total internal energy of the Fermi-Gas we get

$$U_{el} = \int_0^\infty E n_F(E, T) g(E) dE.$$

# Chapter 4

## The form of solids

### 4.1 LCAO - Linear Combination of Atomic Orbitals

Is a quantum mechanical theory, also referred to as molecular orbital theory.

We consider two atoms, atom  $A$  and atom  $B$ . We look that there's a electron orbiting the two atoms, in a molecule it will certainly interact with both atoms. For now we only consider a single electron.

We define the distance between the atoms  $\vec{R}$  and the distance between the atoms and the electrons  $\vec{R}_A$  and  $\vec{R}_B$  respectively.

We assume that there is only one type of orbital. We will now try to write the hamiltonian of this system and deduce the Schroedinger Equation.

$$\hat{H} = \hat{H}_{core} + \hat{H}_{valence}.$$

Where  $\hat{H}_{core}$  only describes the nuclei and the core electrons and  $\hat{H}_{valence}$  only describes the valence electrons.

We introduce a common assumption in Condensed Matter Physics called the Born-Oppenheimer approximation. It assumes:

- nuclei in fixed positions
- kinetic energy of the nuclei  $E_{kin,nuc} = 0$

These assumptions are valid, as the nuclei are much much heavier and move much slower than the electrons.

For the valence hamiltonian we get:

$$H_{valence} = -\frac{\hbar^2}{2m_e} \left( \frac{\partial^2}{\partial x^2} + \frac{\partial^2}{\partial y^2} + \frac{\partial^2}{\partial z^2} \right) - \frac{e^2}{4\pi\epsilon_0} \left( \frac{1}{R_A} + \frac{1}{R_B} \right).$$

With the first term the kinetic energy of the electron and the second term it's coulomb energy. Since we assume that the nuclear kinetic energy is zero our hamiltonian only consists of this valence hamiltonian.

In LCAO we only search for solutions of the form

$$|\psi\rangle = c_A |\phi_A\rangle + c_B |\phi_B\rangle.$$

With  $|\phi_A, B\rangle$  the single atom orbitals.

To find the coefficients  $c_A$  and  $c_B$  we use a variational approach.

$$\langle\psi| H_{val} |\psi\rangle = E \langle\psi|\psi\rangle.$$

We want to minimize the Energy  $E$ .

$$\begin{aligned} E &= \frac{\langle\psi| H_{val} |\psi\rangle}{\langle\psi|\psi\rangle} = \frac{\langle c_A\phi_A + c_B\phi_B | H_{val} | c_A\phi_A + c_B\phi_B \rangle}{\langle c_A\phi_A + c_B\phi_B | c_A\phi_A + c_B\phi_B \rangle} \\ \Rightarrow E &= \frac{(c_A^2 \langle\phi_A| H_v |\phi_A\rangle + c_B^2 \langle\phi_B| H_v |\phi_B\rangle + 2c_Ac_B \langle\phi_B| H_v |\phi_A\rangle)}{c_A^2 \langle\phi_A|\phi_B\rangle + 2c_Ac_B \langle\phi_B|\phi_A\rangle + c_B^2 \langle\phi_B|\phi_B\rangle}. \end{aligned}$$

We introduce the binding integrals  $H_{ij}$  as

$$H_{ij} = \langle\phi_i| H_v |\phi_j\rangle = \int \phi_i^*(\vec{r}) \hat{H}_v \phi_j(\vec{r}) d\vec{r}.$$

For example  $H_{AA}$  describes the energy of the energy of an electron around atom  $A$  that still feels the presence of atom  $B$ .

We further introduce the overlap integrals

$$S_{ij} = \langle \phi_i | \phi_j \rangle = \int \phi_i^* (\vec{r}) \phi_j (\vec{r}) d\vec{r}.$$

For all of these  $0 \leq S_{ij} \leq 1$ . With  $S_{ij} = 0$  meaning that the two orbitals do not overlap at all and  $S_{ij} = 1$  meaning that the orbitals overlap perfectly.

These definitions help with our goal to minimize  $E$ .

$$E = \frac{\langle \psi | H_v | \psi \rangle}{\langle \psi | \psi \rangle}.$$

To find the minimum we set the derivative to 0

$$\begin{aligned} \frac{\partial E}{\partial c_A} &= \frac{\partial E}{\partial c_B} = 0 \\ E &= \frac{(c_A^2 \langle \phi_A | H_v | \phi_A \rangle + c_B^2 \langle \phi_B | H_v | \phi_B \rangle + 2c_A c_B \langle \phi_B | H_v | \phi_A \rangle)}{c_A^2 \langle \phi_A | \phi_B \rangle + 2c_A c_B \langle \phi_B | \phi_A \rangle + c_B^2 \langle \phi_B | \phi_B \rangle} \\ &= \frac{c_A^2 H_{AA} + 2c_A c_B + c_B^2 H_{BB}}{c_A^2 + 2c_A c_B + c_B^2} \\ &\implies \frac{\partial E}{\partial c_A} = 0 = \dots \\ &\implies E = \frac{c_A H_{AA} + c_B H_{AB}}{c_A + c_B S_{AB}} \end{aligned}$$

If we also consider the derivative with respect to  $c_B$  we get:

$$c_A (H_{AA} - E) + c_B (H_{AB} - S_{AB} E) = c_A (H_{AB} - E S_{AB}) + c_B (H_{BB} - E) = 0.$$

If we assume  $H_{ii} = E_0$  (which holds as long as atom  $A$  and  $B$  are of the same type) we get two equations

$$\begin{aligned} c_A (H_{AA} - E) + c_B (H_{AB} - S_{AB} E) &= 0 \\ c_A (H_{AB} - S_{AB} E) + c_B (H_{BB} - E) &= 0 \\ \implies (E_0 - E)^2 - (H_{AB} - S_{AB} E)^2 &= 0. \\ \implies E_{\pm} &= \frac{E_0 \pm H_{AB}}{1 \pm S_{AB}}. \end{aligned}$$

Bonding thus introduces new energy eigenstates  $E_+$  and  $E_-$ , one above and one below the original level  $E_0$ . We call the lower level  $E_+$  the bonding energy and the higher level  $E_-$  the antibonding energy. (since  $H_{AB} \ll 0$  we get  $E_+$  as the lower level). We define  $V_1 = E_0 - E_+$  and  $V_2 = E_0 - E_-$ . In general we find

$$|V_1| < |V_2|.$$

The antibonding level is thus always enough to 'counteract' the bonding level.

We can thus define the binding energy  $E_B^{molecule}$ . If only the bonding level is occupied we get

$$E_B^{molecule} = 2|V_1| = 2(E_0 - E_+).$$

With the factor of 2 coming from two electrons occupying the level.

$$\begin{aligned} S_{AB} \ll 1 &\implies E_+ = \frac{E_0 + H_{AB}}{1 + S_{AB}} \approx E_0 + H_{AB} \\ \implies E_B &\approx 2(E_0 - (E_0 + H_{AB})) = -2H_{AB}. \end{aligned}$$

We consider the splitting energy  $E_- - E_+ = -2H_{AB}$ .

Until now we have not included the coulomb repulsion of the nuclei, for a correct approach we would have to include this.

We now want to determine  $c_A$  and  $c_B$ . For this we now assume  $S_{AB} \approx 0$ .

$$c_A (E_0 - E) + c_B H_{AB} = 0.$$

We find:

- For  $E = E_+ : c_A = c_B = \frac{1}{\sqrt{2}}$
- For  $E = E_- : c_A = -c_B = \frac{1}{\sqrt{2}}$

We therefore write

$$\text{Bonding Wavefunction: } |\psi_+\rangle = \frac{1}{\sqrt{2}} (|\phi_A\rangle + |\phi_B\rangle)$$

$$\text{Antibonding Wavefunction: } |\psi_-\rangle = \frac{1}{\sqrt{2}} (|\phi_A\rangle - |\phi_B\rangle).$$

We call this kind of bonding  $\sigma$ -bonding. A  $\sigma$ -bond is defined as any bond where only one orbital-lobe from each atom contributes (the types of lobes interacting does not matter, it could be S type orbital-lobes or P type orbital-lobes). If two lobes are contributing to the bond, we call it a  $\pi$ -bonding.

## 4.2 Crystal Structures and Symmetries

In Crystals we get Periodic Potentials

$$V(\vec{r}) = V(\vec{r} + \vec{R}).$$

## Chapter 5

# Vibrations of a 1D-Atomic Chain

If you assume stationary atoms, you cannot explain certain physical properties:

- Thermal Expansion
- thermal conductivity and specific heat (can be modelled but not correctly)

We call a quantum of the atomic vibration a *Phonon*. In classical mechanics this would be called a normal mode. We have seen, that the interatomic potential has a asymmetric shape, with the attraction being less steep than the repulsion. As an approximation we assume that the potential is well approximated by a parabola, at least around the equilibrium position.

$$V(x) = V(x_{eq}) + \frac{c}{2}(x - x_{eq})^2 + \frac{c}{3!}O(x^3).$$

In the harmonic approximation we now only consider the first two terms:

$$V_{Harm} = V(x_{Eq}) + \frac{2}{2}(x - x_{Eq})^2.$$

At high temperatures the harmonic approximation fails.

In the asymmetrical potential we can already see an intuitive explanation for the thermal expansion:

$$\overline{x_{Eq}}(T) = \frac{x_{min} + x_{max}}{2} > x_{Eq}.$$

This effect however is not modelled in the harmonic approximation, which is symmetrical.

### 5.1 Monoatomic Chain (Harmonic)

We model a chain of identical Atoms (same mass, charge etc.) in a harmonic potential. This situation is analogous to a chain of mass points connected by springs. As long as we are in the harmonic approximation, we can imagine the atoms connected by springs.

We call the atom in 'the center' of the chain  $n$ , its' left neighbour  $n - 1$  and its' right neighbour  $n + 1$ . We call the equilibrium distance between two of the atoms  $x_{Eq} = a$  the lattice constant. We call the position of atom  $i$   $x_i$ . We thus know the equilibrium position of atom  $i$ ; it is given by

$$x_i^0 = i * a.$$

We now consider vibrations (for example due to temperatures)

$$\delta x_i = x_i - x_i^0.$$

We can now apply Hooke's law:  $\vec{F} = -c\Delta x$

$$\implies \vec{F} = m\vec{a} = -c\Delta x$$

$$\implies m\delta\ddot{x}_n = -c\delta x_n = -c([\delta x_n - \delta x_{n+1}] + [\delta x_n - \delta x_{n-1}]) = c(\delta x_{n+1} + \delta x_{n-1} - 2\delta x_n).$$

We only considered atoms moving along the chain, we only consider longitudinal waves for now.  
As an ansatz for a solution we take:

$$\delta x_i = A \exp\{i\omega t - ikx_n\}.$$

With the frequency  $\omega \geq 0$ ; note there's only one  $\omega$ , as in a normal mode all atoms vibrate with the same frequency.  
With the wavevector  $k$  and  $x_n = n * a$  as before. To find  $\omega$  we write

$$\begin{aligned} m\delta x_n'' &= Am (i\omega)^2 \exp\{i\omega t - ikna\} = CA \exp\{i\omega t\} (\exp\{-ika(n+1)\} + \exp\{-ika(n-1)\} - 2\exp\{-ikna\}) \\ &= C \exp\{-ikna\} (2 \cos ka - 2) \\ \implies -m\omega^2 \exp\{-ikna\} &= C \exp\{-ikna\} (2 \cos ka - 2) \\ \implies \omega^2 &= \frac{4C}{m} \sin^2 \frac{ka}{2}. \end{aligned}$$

We thus find the frequency  $\omega$  as:

$$\omega(k) = 2\sqrt{\frac{C}{m}} * \left| \sin \frac{ka}{2} \right|.$$

We call the relation between  $\omega$  and the wavevector  $k$  a dispersion relation. We have this found above the dispersion relation of a 1-D Monoatomic Chain.

We see that this function is periodic with  $k \rightarrow k \pm \frac{2\pi}{a}$ .

In general we know that: If a system is periodic in real space  $(x, y, z)$  then it is also periodic in the reciprocal space  $(k_x, k_y, k_z)$ .

This is a useful fact that will come in handy many times.

We call one of the periodic zones in the reciprocal space a Brillouin zone (BZ). The first Brillouin zone reaches from  $-\frac{\pi}{a}$  to  $\frac{\pi}{a}$ . It is usually sufficient to plot any function within the first BZ.

We thus find that  $\delta x_n(k) = \delta x_n(k + \frac{2\pi}{a})$ .

$$\begin{aligned} \delta x_n(k) &= A * \exp\{i\omega t - ikna\} \\ \delta x_n(k \pm \frac{2\pi}{a}) &= A \exp\left\{i\omega t - i\left(k \pm \frac{2\pi}{a}\right)na\right\} = A \exp\{i\omega t - ikna\} * \exp\{-i2\pi n\}. \\ \implies n &\in \mathbb{Z}. \end{aligned}$$

We say that  $\vec{k}$  is defined up to any lattice vector. In our case the lattice vector is  $\frac{2\pi}{a}$ . We thus define the reciprocal lattice vector  $G_m$

$$G_m = \frac{2\pi}{a} * m, \forall m \in \mathbb{Z}.$$

We get

$$\exp\{iG_m \vec{x}_m\} = 1.$$

We can thus turn the definition around and say that this exponential gives the definition of the reciprocal lattice.

### 5.1.1

We found the dispersion relation

$$\omega = 2\sqrt{\frac{C}{m}} * \sin \frac{ka}{2}.$$

We now think about this relation

- $k \rightarrow 0$  we find:  $\implies \omega = 2\sqrt{\frac{C}{m}} \frac{ka}{2} = \sqrt{\frac{C}{m}} * ka$   
We thus find a linear dispersion relation  $\omega \propto k$  with a constant called the speed of sound:  $\omega = k * v_{\text{sound}}$ .  
Any time we find a linear dispersion we call it a sound like dispersion.

We define certain velocities:

- Sound Velocity  $V_s = a\sqrt{\frac{C}{m}}$  is the velocity of the wave at  $k = 0$ .

- Phase Velocity  $v_{phase} = \frac{\omega}{k}$  speed at which the maxima and minima travel
- Group Velocity  $v_{group} = \frac{d\omega}{dk}$  speed of the wave packet

For materials with linear dispersion we find  $v_p = \frac{\alpha k}{k} = \alpha = \frac{d(\alpha k)}{k} = v_g$ . Phase and group velocities are equal only for linear dispersion.

For example we find in the case of the monoatomic chain, that the group velocity  $v_g = 0$  for  $k = \frac{\pi}{a}$ . How do we count the number of normal modes?

$$\# \text{modes} = \frac{\# \text{all possible } k}{\text{spacing between neighbouring } k} = \frac{\frac{2\pi}{a}}{?}$$

To find the spacing between the possible  $k$ , we use periodic boundary condition:  $x_i = x_{i+N}$ . Meaning that we assume the monoatomic chain to be a closed loop. We get that:

$$\begin{aligned} A \exp\{i\omega t - iKna\} &= A \exp\{i\omega t - iK(n+N)a\} \\ \implies \exp\{-iKNa\} &= 1 \\ \implies K &= P \frac{2\pi}{Na}, \forall P \in \mathbb{Z}. \end{aligned}$$

We thus know the number of modes as:

$$\# \text{modes} = \frac{\frac{2\pi}{a}}{\frac{2\pi}{Na}} = N.$$

We find that the total number of modes is equal to the number of atoms, since we are considering only longitudinal waves in our 1D system this is also the number of degrees of freedom. While Debye had to introduce a cutoff frequency to get this result, we now found it generally.

### 5.1.2 Types of Vibrations

Up to now we only considered longitudinal waves in our 1D system. In addition to the  $\delta x$  we already considered there are also  $\delta y$  and  $\delta z$  to consider. Since  $\delta y, \delta z \perp \vec{K}$  these waves are Transverse modes. In general we find

$$\omega(\text{Transverse mode}) < \omega(\text{longitudinal mode}).$$

## 5.2 Diatomic linear chain

We still assume a harmonic potential, but now we assume that we have two different types of atoms, repeating in a ABABABA manner. We thus get different spring constants  $C_1$  and  $C_2$  and different masses  $m_1$  and  $m_2$ . The first step to solve such a problem, is to select a periodic unit cell. We thus select a unit cell of length  $a$  that includes one atom pair AB.

The next step is to identify a reference point, since the unit cell is no longer symmetrical the center of the unit cell is no longer the center of mass. Instead of describing the position of an atom like before, we now call the coordinate of that reference point  $\delta x_n$ . We thus get the position of the unit cell as  $r_n = n * a$ .

The third step is to describe the position of the atoms in relation to the reference point. We call the positions of the Atoms

$$\begin{aligned} x_n^{eq} &= an - \alpha a \\ y_n^{eq} &= an + \beta a. \end{aligned}$$

For simplicity we assume  $m_1 = m_2$  and  $c_1 \neq c_2$ , without loss of generality we assume  $c_2 > c_1$ . We now solve the problem similar as before. We consider the force on one of the atoms

$$\begin{aligned} F_1 &= m\delta\ddot{x}_n = c_2(\delta y_n - \delta x_n) + c_1(\delta y_{n-1} - \delta x_n) \\ F_2 &= m\delta\ddot{x}_n = c_2(\delta x_n - \delta y_n) + c_1(\delta x_{n+1} - \delta y_n). \end{aligned}$$

We choose the ansatz

$$\begin{aligned} \delta x_n &= A_x \exp\{i\omega t - iKna\} \\ \delta y_n &= A_y \exp\{i\omega t - iKna\}. \end{aligned}$$



We get two different amplitudes, but the same frequency. As the definition of a normal mode is per its' frequency. As before we get  $\omega > 0$ ,  $K$  positive or negative and the number of  $K$  values =  $\frac{\text{range of } K}{\text{spacing of } k} = \frac{\frac{2\pi}{a}}{\frac{2\pi}{Na}} = N$ . We have to remember that  $N$  here is the number of unit cells, not the number of atoms. By inserting the ansatz into the equation of motion we get:

$$-\omega^2 m A_x \exp\{i\omega t - iKna\} = c_2 A_y \exp\{i\omega t - iKna\} + c_1 A_y \exp\{i\omega t - iKna\} - (c_1 + c_2) A_x \exp\{i\omega t - iKa\}$$

$$\Rightarrow \omega^2 m \begin{bmatrix} A_x \\ A_y \end{bmatrix} = \begin{bmatrix} c_1 + c_2 & -c_2 - c_1 \exp\{ika\} \\ -c_1 \exp\{-ika\} - c_2 & c_1 + c_2 \end{bmatrix} \begin{bmatrix} A_x \\ A_y \end{bmatrix}.$$

We thus get an Eigenvalue problem the solution yields:

$$m\omega^2 = c_1 + c_2 \mp \sqrt{c_1^2 + c_2^2 + 2c_1 c_2 \cos(ka)}$$

$$\omega_{\pm} = \sqrt{\frac{c_1 + c_2}{m} \mp \frac{1}{m} \sqrt{c_1^2 + c_2^2 + 2c_1 c_2 \cos(ka)}}$$

For each  $K$  we thus get two possible  $\omega$ . We therefore have two dispersion relations  $\omega_+(K)$  and  $\omega_-(K)$ . We call these two dispersion relations *phonon branches*.

In this case the two phonon branches have famous names:

- $\omega_-$  is called the *acoustic branch*

$$\lim_{K \rightarrow 0} \omega_-(K) = \sqrt{\frac{c_1 c_2 a^2}{2(c_1 + c_2)}} |K|$$

We see that  $\omega_-$  has a linear dispersion (acoustic dispersion) at small  $K$ .

- $\omega_+$  is called the *optical branch*

$$\lim_{K \rightarrow 0} \omega K = \sqrt{\frac{2(c_1 + c_2)}{m}}$$

The optical branch has a group velocity of  $v_g = \frac{d\omega_+}{dK} = 0$ .

The reason that this branch is called the optical branch is, that we can measure it by using light as a perturbation.

We thus see that for  $K \rightarrow 0$  we get two different values. We get

$$\omega_-(K \rightarrow 0)$$

$$\Rightarrow \begin{bmatrix} 0 \\ 0 \end{bmatrix} = \frac{c_1 + c_2}{m} \begin{bmatrix} A_x - A_y \\ -A_x + A_y \end{bmatrix}$$

$$\Rightarrow A_x = A_y.$$

In acoustic branches the atoms vibrate in phase.

For the optical branch we get:

$$\dots \Rightarrow A_x = -A_y.$$

In optical branches the atoms vibrate out of phase.

Because the energy requirement of vibrating out of phase is much higher we find:

$$\omega(\text{optical branches}) > \omega(\text{acoustical branches}).$$

(We have demonstrated this only at  $K = 0$  but it holds in general).

We will now also demonstrate it at the BZ boundaries  $K = \pm \frac{\pi}{a}$ :

$$\omega_+ \left( k = \frac{\pi}{a} \right) = \sqrt{\frac{2c_2}{m}}$$

$$\omega_- \left( k = \frac{\pi}{a} \right) = \sqrt{\frac{2c_1}{m}}.$$

Since we know  $c_2 > c_1 \Rightarrow \omega_- < \omega_+$ .

In summary

- For the monoatomic Chain we got:

–

- For the diatomic chain we got ( $2N$  atoms):

–  $2N$  Branches

- in general:

If there are  $M$  atoms in a unit cell we will get  $M$  modes at each  $K$ .

We always get 1 acoustic mode and  $M - 1$  optical modes.

In  $d$  dimensions we get  $d * M$  modes at each  $\vec{K}$ . We find  $d$  acoustic modes and  $d(M - 1)$  optical modes.

## 5.3 Quantum Correspondence

Discrete quantum vibrations of phonons with energy  $\hbar\omega$ . We utilize the Bose occupation factor again, as phonons are bosons

$$n_B(\hbar\omega\beta) = \frac{1}{\exp(\hbar\omega\beta) - 1}.$$

We thus find the energy expectation of a single phonon as

$$\langle E_n \rangle = \hbar\omega_k \left( n_B + \frac{1}{2} \right).$$

For the total energy we have to sum over the first BZ

$$U_{tot} = \sum_{k=-\frac{\pi}{a}}^{\frac{\pi}{a}} \hbar\omega_k \left( n_B + \frac{1}{2} \right).$$

For example in the case of a mono-atomic chain we found:

$$\omega_k = 2\sqrt{\frac{c}{m}} \left| \sin\left(\frac{ka}{2}\right) \right|.$$

For many atoms  $N \rightarrow \infty$  we get  $\sum_k \rightarrow \frac{L}{2\pi} \int_{-\frac{\pi}{a}}^{\frac{\pi}{a}} dK$ .

This is a more general expression for the total energy, as for example we found in the Einstein or Debye models:

- Einstein:  $\omega_k = \omega_{einstein}$
- Debye:  $\omega_k = v_s * k$

But now we can correctly describe  $\omega_k$  without using these approximations.

As before we can introduce the density of states  $g(\omega)$ .

$$\frac{L}{2\pi} \int dK = \int g(\omega) d\omega$$

$$g(\omega) = 2 \frac{Na}{2\pi} \left| \frac{dK}{d\omega} \right| \propto \frac{1}{\left| \frac{d\omega}{dK} \right|}$$

For mono atomic chains we thus find

$$g(\omega) = \frac{1}{\cos\left(\frac{ka}{2}\right)}.$$

## 5.4 Crystal Momentum

$K$  is fully equivalent to  $K \pm G_m$ , as defined before.

We define crystal momentum as

$$\hbar \left( K \bmod \frac{2\pi}{a} \right).$$

We thus get that the conservation of momentum is still valid, but is not absolute, instead it occurs modulo  $\frac{2\pi}{a}$ . This basically means, that the momentum is always defined by the equivalent  $K$  in the first BZ.

## Chapter 6

# Tight binding chain

We consider electrons in periodic potentials. We assume equally spaced atoms, each with only one available orbital. We define  $|n\rangle$  as the orbital of the  $e^-$  on the atom  $n$ . We also assume  $\langle n|m\rangle = \delta_{mn}$ . As before we have LCAO with  $S_{AB} = \langle \phi_A | \phi_B \rangle = \delta_{mn}$ , this is only an approximation, but holds in this toy model. We write the (Born-Oppenheimer) hamiltonian as

$$\hat{H} = \frac{\vec{p}^2}{2m} + \sum_j V(\vec{r} - \vec{R}_j) = K + \sum_j V_j.$$

We make an Ansatz for a trial wavefunction as

$$|\psi\rangle = \sum_n \phi_n |n\rangle.$$

To find the factors  $\phi_n$  we solve the Schroedinger Equations:

$$\sum_m H_{nm} \phi_m = E \phi_n.$$

Proof:

$$E = \frac{\langle \psi | H | \psi \rangle}{\langle \psi | \psi \rangle}$$

Missing

We can define the energy of one atom as

$$E_m |m\rangle = H |m\rangle = (K + V_m) |m\rangle.$$

In the case where we have many nuclei we get

$$H = K + V_m + \sum_{j \neq m} V_j.$$

$$\implies E |m\rangle = E_m |m\rangle + \sum_{j \neq m} V_j |m\rangle.$$

$$\implies H_{nm} = \langle n | H | m \rangle = E_{atomic} \langle n | m \rangle + \langle n | \sum_{j \neq m} V_j | m \rangle.$$

- $n = m \implies \langle n | \sum_j V_j | m \rangle = \langle m | \sum_j V_j | n \rangle = V_0$
- $n \neq m \implies \langle n | \sum_{j \neq n} V_j | m \rangle = -t$  "Hopping energy"  
if  $|n - m| > 1 \implies$  "Hopping"  $\approx 0$ , we assume that only direct neighbours interact with each other.
- $|n - m| > 1 \implies \sum_{j \neq n} \langle n | V_j | m \rangle = 0$

$$\Rightarrow H_{nm} = E_{atomic}\delta_{nm} + \langle n | \sum_j V_j | m \rangle = (E_a + V_0)\delta_{nm} + (-t) * (\delta_{n,m+1} + \delta_{n,m-1})$$

$$\Rightarrow H_{nm} = E_0\delta_{nm} - t(\delta_{n,m+1} + \delta_{n,m-1})$$

We make the ansatz  $\phi_n = \frac{\exp(-ikna)}{\sqrt{N}}$ .

$$\sum_m H_{nm}\phi_m = E\phi_n$$

$$\frac{E_0 \exp(-ikna)}{\sqrt{N}} = \frac{t}{\sqrt{N}} (\exp(-ik(n-1)a) + \exp(-ik(n+1)a)) = E \frac{1}{\sqrt{N}} \exp(-ikna)$$

$$\Rightarrow E = E_0 - 2t \cos(ka).$$

We call this relation between  $E$  and  $K$  the electric dispersion. This gives us an energy band, aka. the possible energies for the electrons.

We have found  $\phi_n = \frac{\exp(-ikna)}{\sqrt{N}}$ . The wavefunction of an electron is therefore delocalized over the whole crystal.

If we expand  $E$  around  $K \approx 0$  we get:

$$E \approx E_0 - 2t \left( 1 - \frac{(ka)^2}{2} \right) = \text{const.} + tk^2 a^2.$$

Quadratic dependence!

For free electrons we know  $E = C + \hbar \frac{K^2}{2m}$ , at  $K \approx 0$  we therefore get a similar behaviour as for free electrons. We find that

$$\frac{\hbar^2}{2m^*} = a^2 t$$

$$E = \text{const.} + \frac{\hbar^2 K^2}{2m^*}.$$

With the 'effective mass'  $m^*$  which depends on  $t$ . For now this effective mass is only a mathematical constant with a dimension of a mass, later we will see that it also has a physical meaning. The effective mass has no relation with the real mass of the electron.

Above we spoke of an energy band, but is this justified, the  $K$  vectors are discrete, shouldn't we get discrete energy levels from this?

Technically we do, but there are many states in a relatively small interval of energy, we can therefore assume a continuous energy band. We now want to count the states

$$\# \text{ of states} = 2 * 2 * \frac{dN}{dE}$$

$K$  can be  $\pm$  and there are two electrons per level  $\rightarrow 2 * 2$ . Rewriting

$$\frac{dN}{dE} = \frac{dN}{dK} \frac{dK}{dE}.$$

We know that  $\frac{dK}{dN} = \frac{2\pi}{L}$  and we also know the dispersion relation between  $K$  and  $E$ .

## 6.1 Monovalent Atoms

Monovalent Atoms have only 1 electron in the outer most shell.  $N \text{ atoms} \rightarrow N e^-$ .

But the energy Band can host  $2N$  electrons, it will thus only be half filled. The band will be filled from the bottom up, since there are free electron bands, there can be an excitation this results in the crystal being able to conduct electricity and heat really well.

## 6.2 Divalent Atoms

$N \text{ atoms} \rightarrow 2N$  electrons. All bands are filled, diatomic crystals are insulators and their electrons don't contribute to the heat capacity.

### 6.3 $N$ orbitals in the unit Cell

There are either more than one atoms per unit cells or each atom has more than one orbital. This is the most common situation.

This situation is analougous to the phonos in a diatomic chain. We also get two dispersion branches.

Example: 3 electrons in each unit cell.  $N$  atoms  $\rightarrow 2N$  possible energies in the first energy band but  $3N$  electrons.

We completely fill the lower band and then fill the remaining electrons into the higher band.

The filled lower band does, as above, not contribute to thermal or electrical conduction. However the higher band is not completely filled and thus excitations are possible, this higher band contributes to the thermal and electrical conductivity. The smallest distance between the two bands is called a 'band gap'.

# Chapter 7

## Crystalline Structures

A crystal is an arrangement of material, which is periodic in space. This can mean a periodic arrangement of atoms or whole molecules.

### 7.1 Important definitions

- A crystal is a PERIODIC arrangements of atoms
- A lattice is a set of points defined as integer sums of *primitive lattice vectors*

A lattice has three main characteristics:

1. A lattice is a set of points defined as integer sums of *primitive lattice vectors*
2. A lattice is a set of vectors such that addition of two valid lattice vectors gives a third valid vector
3. A lattice looks the same from every lattice point.

Any lattice describes a crystal, but not every crystal can be described by a lattice.

Any periodic structure can be transformed in a lattice by carefully choosing a unit cell. If we have a periodic structure we can choose a unit cell as a *basis* such that these cells form a lattice.

We can thus describe each crystal by choosing a suitable basis and it's corresponding lattice. The choice of the unit cell is not unique. We can therefore choose unitcells such that it's mathematically convinient.

A famous example of a unit cell are the so called Wigner-Seitz unit cell. It is defined as the region of space around a given lattice point, that is closer to this lattice point than to any other.

To fully describe a crystal we have to specify the basis and the positions of the atoms within these unit cells.

### 7.2 Symmetry Operations

Symmetry operations can be used to classify lattices.

- Translational Symmetry:  
There always exists a vector  $\vec{R}$  such that a translation by  $\vec{R}$  leaves the system unchanged. ALL CRYSTALS HAVE TRANSLATIONAL SYMMETRY
- Rotational Symmetry:  
We can rotate the system by an angle  $\sigma$  such that the system remains unchanged. Not all crystals have rotational symmetry.
- Reflection  
There can be one or many mirror planes such that a reflection across these planes leaves the system unchanged. Not all crystals have a Symmetry in reflexion.
- Inversion  
 $(x, y, z) \rightarrow (-x, -y, -z)$ . Not all crystals have a Inversion symmetry.

Some parts missing here

### 7.2.1 Summary of the properties of the reciprocal lattice

- For every direct lattice  $\exists$  a reciprocal lattice
- The reciprocal of a reciprocal lattice is the original direct lattice  
if we have a set of reciprocal lattices  $\{\vec{G}\}$  and it's reciprocal is  $\{\vec{M}\}$   
 $\implies e^{i\vec{G}\cdot\vec{M}} = 1$  i.e.  $\{\vec{M}\} = \{\vec{R}\}$
- $\vec{R} \cdot \vec{G} = 2\pi m$  with  $m \in \mathbb{Z}$
- $\vec{G}(h, k, l) = h\vec{b}_1 + k\vec{b}_2 + l\vec{b}_3$ , where  $h, k, l$  are the miller indecies.  $\vec{G} \perp (h \ k \ l)$  lattice plane.
- The separation between lattice planes of the direct lattice  $(h \ k \ l)$   
 $d(h, k, l) = \frac{2\pi}{|\vec{G}(h, k, l)|} = \frac{a}{\sqrt{h^2 + k^2 + l^2}}$

## 7.3 Brillouin Zone

We want to study waves in crystals (eg. electrons or phonons).

We define the Brillouin Zone (BZ) as the unit cell of the reciprocal lattice.

We call the  $N^{\text{th}}$  BZ: points in  $\vec{k}$  space where  $\vec{k} = 0$  is the  $N^{\text{th}}$  closest reciprocal lattice point are the  $N^{\text{th}}$  BZ. This corresponds to the Wigner Seitz unit cell of the reciprocal lattice. The number of  $\vec{k}$  states in a BZ is equal to the number of unitacells in the system.

We consider some examples in 3d:

- Lattice | Rep. Lattice | First BZ
- Simple cubic | Simple cubic | Cube
- BCC | FCC | WS cell for FCC
- FCC | BCC | WS cell for BCC

## Chapter 8

# Reciprocal Space



# Chapter 9

## Scattering Experiments

By sending a known wave with  $\vec{k}$  into a sample we usually get a scattered part  $\vec{k}'$  and an unscattered part  $\vec{k}$ . By studying the scattered wave  $\vec{k}'$  we can infer the structure.

The wave should be comparable in size to the structure you want to investigate. In CoMa we want to investigate crystal structures of solids, we thus need waves with a wavelength on the order of a few Angstroms. In particular we consider X-Rays and Neutrons. (In theory electrons are also feasible, but physically much more complex - in this lecture we only consider X-Rays and Neutrons).

The atoms in a crystal are producing a periodic potential. Our wave is interacting with this potential  $V(\vec{R})$  that is scattering the wave.

### 9.1 Fermi's Golden Rule

We look at the Scattering Rate  $\Gamma$ :

$$\Gamma(k, k') = \frac{2\pi}{\hbar} \left| \langle \vec{k}' | V | \vec{k} \rangle \right|^2 \delta(E_{k'} - E_k).$$

We assume ELASTIC Scattering; this means we assume  $E = E'$  and  $|\vec{k}| = |\vec{k}'|$

This is a reasonable assumption as the majority of a scattered signal results from elastic scattering and only a negligible part of the signal results from inelastic scattering.

We now investigate the matrix element

$$\begin{aligned} \langle k' | V | k \rangle &= \int d\vec{r} e^{-i\vec{k}' \cdot \vec{r}} V(\vec{r}) e^{i\vec{k} \cdot \vec{r}} = \int d\vec{r} e^{-i(\vec{k}' - \vec{k}) \cdot \vec{r}} V(\vec{r}) \\ &= FT[V(\vec{r})]. \end{aligned}$$

If we consider a periodic scattering potential we find

$$FT[V(\vec{r})] = \frac{(2\pi)^3}{V_{unitcell}} \sum_{\vec{G}} \delta^3(\vec{k} - \vec{k}' - \vec{G}) S(\vec{G}).$$

With the structure factor  $S(\vec{G})$  given as

$$S(\vec{G}) = \int_{unitcell} d^3\vec{r} e^{i\vec{G} \cdot \vec{r}} V(\vec{r}).$$

Any periodic potential has a FT with non-zero values only at the positions of the RECIPROCAL LATTICE VECTORS and its values are weighted by the structure factor. We thus find:

1. Scattering is non-zero only if  $\vec{k} - \vec{k}' = \vec{G}$   
this relation which expresses the conservation of crystal momentum is called the Laue condition.
2. Intensity of Scattering is proportional to  $|S(\vec{G})|^2$

## 9.2 Equivalence of Laue and Bragg Conditions

The Laue condition is equivalent to imposing constructive interference between waves scattered on different lattice planes.

We assume a wave incident on a lattice plane with an angle of  $\theta$ . Note: the wave is thus deflected by  $2\theta$ .

A wave reflected on the second lattice plane has to travel an additional distance equal to  $2d \sin(\theta)$ . To get constructive interference we need

$$2d \sin(\theta) = n\lambda.$$

We know that our  $\vec{G}$  is perpendicular on the lattice plane. Since we only consider elastic scattering we have  $|k| = |k'|$ . We define  $\hat{k}$ ,  $\hat{k}'$ ,  $\hat{G}$  as the unit vectors in the directions of the corresponding vectors. We find

$$\hat{k}' \cdot \hat{G} = -\sin(\theta)$$

$$\hat{k} \cdot \hat{G} = \sin(\theta).$$

We rewrite the Laue condition

$$\frac{2\pi}{\lambda} (\hat{k} - \hat{k}') = \vec{G}$$

$$\frac{2\pi}{\lambda} (\hat{G} \cdot \hat{k} - \hat{G} \cdot \hat{k}') = \hat{G} \cdot \vec{G}$$

$$\frac{2\pi}{\lambda} (\sin(\theta) - (-\sin(\theta))) = |\vec{G}|.$$

$$\frac{2\pi}{|\vec{G}|} 2 \sin(\theta) = \lambda$$

$$2d \sin(\theta) = n \cdot d \text{ with } n \in \mathbb{Z}.$$

We have now recovered the Bragg condition from the Laue condition. We have thus found that the conservation of momentum corresponds to constructive interference.

## 9.3 Intensity of Scattering

$$I \propto \left| S(\vec{G} = \vec{k} - \vec{k}') \right|^2$$

With  $S = FT[V(\vec{r})]$ .