

Example

1D chain of ions connected by springs



nearest neighbor interaction only
 $\frac{1}{2}K(u_i - u_{i+1})^2$

u_i = displacement of ion i

$$M\ddot{u}_i = -K(u_i - u_{i+1}) - K(u_i - u_{i-1}) \quad \text{integer } n$$

Assume $u_n(t) = u_0 e^{i(kR_n - \omega t)}$ $R_n = na$
Substitute in and cancel common factors of $e^{i(kR_n - \omega t)}$

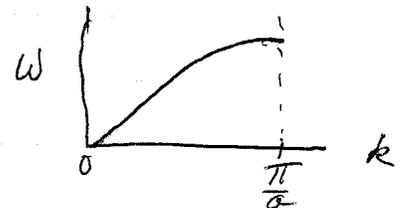
$$-\omega^2 M u_0 = -K(u_0 - u_0 e^{ika}) - K(u_0 - u_0 e^{-ika})$$

$$\begin{aligned} \Rightarrow -\omega^2 M &= -K(1 - e^{ika} + 1 - e^{-ika}) \\ &= -2K(1 - \cos ka) \end{aligned}$$

$$\omega = \sqrt{\frac{2K}{M} (1 - \cos ka)}$$

use $\frac{1 - \cos ka}{2} = \sin^2\left(\frac{ka}{2}\right)$

$$\omega = \sqrt{\frac{K}{M}} 2 \left| \sin\left(\frac{ka}{2}\right) \right|$$



at small $ka \ll 1$, $\sin ka \approx ka$

$$\omega \approx \sqrt{\frac{K}{M}} ka \Rightarrow \text{speed of sound}$$

$$c = \sqrt{\frac{K}{M}} a$$

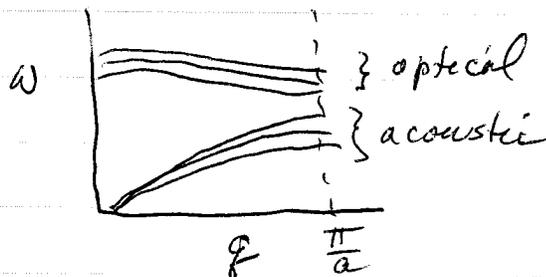
Previous discussion assumed monatomic B.L.
 When have BL with basis, the dynamic matrix must acquire an additional index that labels the n atoms in the basis at any BL site ~~lattice~~ R_i

\Rightarrow 3 modes for each atom in primitive cell of BL

$\rightarrow 3n$ elastic modes

of these, 3 are acoustic modes as before - one longitudinal, two transverse - with $\omega_s \approx c_s q$ as $q \rightarrow 0$.

The $3(n-1)$ remaining modes are "optical" modes where $\omega_s(q) \rightarrow \text{const}$ as $q \rightarrow 0$.



see A+M chpt 22
 and problem #1
 on Problem Set 6

optical modes correspond to ^{"internal"} vibrations of the atoms within a primitive cell of the BL with respect to each other.
 Acoustic modes correspond to motions of the primitive cell as a whole.

When we treat the elastic vibrations of the solid quantum mechanically, these "normal modes" of elastic vibration, i.e. the independent modes of harmonic oscillation, get quantized just like harmonic oscillators. Each degree of excitation of a given mode of oscillation is called a "phonon".

For example, if the vibration at wave vector \vec{q} , polarization s has energy

$$\hbar \omega_s(\vec{q}) (n + 1/2)$$

We say there are n phonons of wave vector \vec{q} polarization s .

As with excitations of any harmonic oscillator phonons behave as bosons, and ~~are not~~ their number is not conserved (chemical potential $\mu_{\text{phonon}} = 0$).

Electrons can scatter by absorbing or emitting phonons, while conserving energy and crystal momentum, i.e. for absorption

$$E(\vec{k}_f) = E(\vec{k}_i) + \hbar \omega_s(\vec{q})$$

$\vec{k}_f = \vec{k}_i + \vec{q} + \vec{K}$

\vec{k}_f final electron crystal momentum
 \vec{k}_i initial electron crystal momentum
 \vec{q} phonon crystal momentum
 \vec{K} Brillouin vector

But: Suppose we consider the conduction electrons as frozen, and uniform, and the ion-ion interaction therefore is Coulomb.

In our discussion of plasma oscillations we saw that the only longitudinal mode of oscillation of a Coulomb interacting set of charges is, as $q \rightarrow 0$, at the plasma frequency. For ions of mass M and density n_{ion} , this would be

$$\Omega_p = \sqrt{\frac{4\pi n_{ion} Q_{ion}^2}{M}} \quad Q_{ion} = \text{charge of ion}$$

This does not agree with the expectation above that the frequency of oscillation for a longitudinally polarized elastic vibration should be, $\omega_s = c_s q$, vanishing as $q \rightarrow 0$!

Why? Because if interaction between ions is pure Coulomb, then the sum $\sum_{R_i} (\hat{q} \cdot \vec{R}_i)^2 \frac{1}{R_i^3}$ does not converge, as we had assumed in the previous discussion!

But we know from experiment and experience that longitudinal (acoustic) sound modes do exist with $\omega_s = c_s q$ linear dispersion relation! What is the resolution of this paradox?

also called
Born-Oppenheimer
approx

The answer is screening! We make the adiabatic approximation and assume that conduction electrons move so much faster than ions that they always relax to their minimum energy configuration corresponding to the instantaneous positions of the ions, as the ions move. The electrons will then screen the Coulombic ion-ion interaction and make it short ranged. The sum $\sum (\hat{g} \cdot \vec{R}_i)^2 \overleftrightarrow{D}(\vec{R}_i)$ now converges and we get the longitudinal elastic modes with $\omega_L = c_L q$. Moreover we can use this argument to estimate the speed of sound c_L .

The phonon freq for polarization s , wavevector \vec{q} was determined by

$$\omega^2 M \vec{E}_s = \overleftrightarrow{D}(\vec{q}) \cdot \vec{E}_s$$

If we let $\overleftrightarrow{D}^0(\vec{q})$ be the dynamical matrix due to bare Coulombic ion-ion interactions, then we expect for the longitudinal mode that $\omega_L = \Omega_L q$, i.e.

$$\Omega_L^2 M \vec{E}_L = \overleftrightarrow{D}^0(\vec{q}) \cdot \vec{E}_L$$

Now a longitudinal ionic vibration of wave vector \vec{q} sets up a charge density of wave vector \vec{q} , which sets up an electric field of wave vector \vec{q} . The electrons screen this field by a factor $\frac{1}{\epsilon(\vec{q})}$ where $\epsilon(\vec{q})$ is the electron dielectric function.

Since $\vec{D}(\vec{q})$, the dynamical matrix, is \propto to the ion-ion forces (effective ion-ion spring constant in the harmonic approx) we expect that these forces will get screened by the electrons and so the screened dynamical matrix \vec{D} is related to the bare \vec{D}^0 by

$$\vec{D}(\vec{q}) = \frac{\vec{D}^0(\vec{q})}{\epsilon(\vec{q})}$$

Hence we expect that

$$\begin{aligned} \Omega_{\vec{q}}^2 M \vec{\epsilon}_e &= \vec{D}^0(\vec{q}) \cdot \vec{\epsilon}_e \Rightarrow \frac{\Omega_{\vec{q}}^2}{\epsilon(\vec{q})} M \vec{\epsilon}_e = \frac{\vec{D}^0(\vec{q})}{\epsilon(\vec{q})} \cdot \vec{\epsilon}_e \\ \Rightarrow \frac{\Omega_{\vec{q}}^2}{\epsilon(\vec{q})} M \vec{\epsilon}_e &= \vec{D}(\vec{q}) \cdot \vec{\epsilon}_e \end{aligned}$$

so the freq of oscillation is now

$$\omega_e^2(\vec{q}) = \frac{\Omega_{\vec{q}}^2}{\epsilon(\vec{q})}$$

For small \vec{q} we can use the Thomas-Fermi approx

$$\epsilon(q) \approx 1 + k_0^2/q^2 \quad \text{where } k_0^2 = 4\pi e^2 g(E_F)$$

So

$$\omega_e^2(q) = \frac{\Omega_p^2}{1 + k_0^2/q^2} = \frac{\Omega_p^2 q^2}{k_0^2 + q^2} \approx \frac{\Omega_p^2}{k_0^2} q^2$$

for small $q \ll k_0$

$$\omega_e(q) = \left(\frac{\Omega_p}{k_0}\right) q \Rightarrow \text{speed of sound is}$$

$$c_e = \frac{\Omega_p}{k_0}$$

$$\Rightarrow c_e^2 = \frac{4\pi n_{\text{ion}} Q_{\text{ion}}^2}{M} \frac{1}{4\pi e^2 g(E_F)}$$

if n is conduction electron density and Z the valence number of conduction electrons, then

$$n_{\text{ion}} = \frac{n}{Z}, \quad Q_{\text{ion}} = Ze$$

$$c_e^2 = \frac{n Z}{M g(E_F)}$$

In the free electron approx, $g(E_F) = \frac{3}{2} \frac{M}{E_F}$

$$\text{So } c_e^2 = \frac{n Z}{M \left(\frac{3}{2} \frac{M}{E_F}\right)} = \frac{2 Z E_F}{3 M} = \frac{2 Z}{3 M} \frac{1}{2} m v_F^2$$

$$c_e^2 = \frac{Z m}{3 M} v_F^2$$

$$c_e = \sqrt{\frac{Z m}{3 M}} v_F$$

For ions ($\frac{m_{elec}}{m_{proton}} \sim \frac{1}{2000}$) we expect

$$\frac{c_e}{v_F} = \sqrt{\frac{Z}{3} \frac{m}{M}} \sim 0.01$$

Our result that $c_e \approx 0.01 v_F$ is consistent with the adiabatic approx that electrons move with speeds (v_F) much greater than the ions (c_e)

The above result is known as the Bohm-Staver relation

It gives results in correct order of magnitude agreement with experiment. For typical metals

$$v_F \sim 10^8 \text{ cm/sec}$$

$$c_e \sim 10^6 \text{ cm/sec}$$