

# Content of this lecture

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1. Electromagnetic wave
  - Maxwell's equations
  - Constitutive relations
  - Time- and spatial-harmonic field
  - Boundary conditions
  - Wave equation
2. Dispersion of materials
  - What is dispersion?
  - Phase and group velocities (in  $k$ - $\omega$  plot)
  - $k$ - $\omega$  dispersion relation
3. Microscopic and macroscopic theories of materials
  - Free and bound electrons
  - EM response of insulator/dielectric: **Lorentz model**
  - EM response of metal: **Drude model** (Lecture 5)
  - Band structures of materials
4. Example of engineering light-matter interaction with nanostructures – **form birefringence**

# 1. Electromagnetic wave

How to describe the wave property of light? – Maxwell's equations

$$\nabla \cdot \mathbf{D} = \rho_{\text{ext}}$$

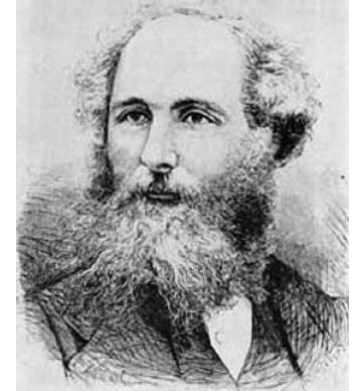
$$\nabla \cdot \mathbf{B} = 0$$

$$\nabla \times \mathbf{E} = -\partial \mathbf{B} / \partial t$$

$$\nabla \times \mathbf{H} = \partial \mathbf{D} / \partial t + \mathbf{J}_{\text{ext}}$$

Divergence Eqs.

Curl Eqs.



(1831 – 1879)

If no external charges and currents:

$$\nabla \cdot \mathbf{D} = 0$$

$$\nabla \cdot \mathbf{B} = 0$$

$$\nabla \times \mathbf{E} = -\partial \mathbf{B} / \partial t$$

$$\nabla \times \mathbf{H} = \partial \mathbf{D} / \partial t$$

linking four  
macroscopic fields  
 $\mathbf{E}$ ,  $\mathbf{H}$ ,  $\mathbf{D}$ ,  $\mathbf{B}$

$\mathbf{E}$  – electric field vector

$\mathbf{H}$  – magnetic field vector

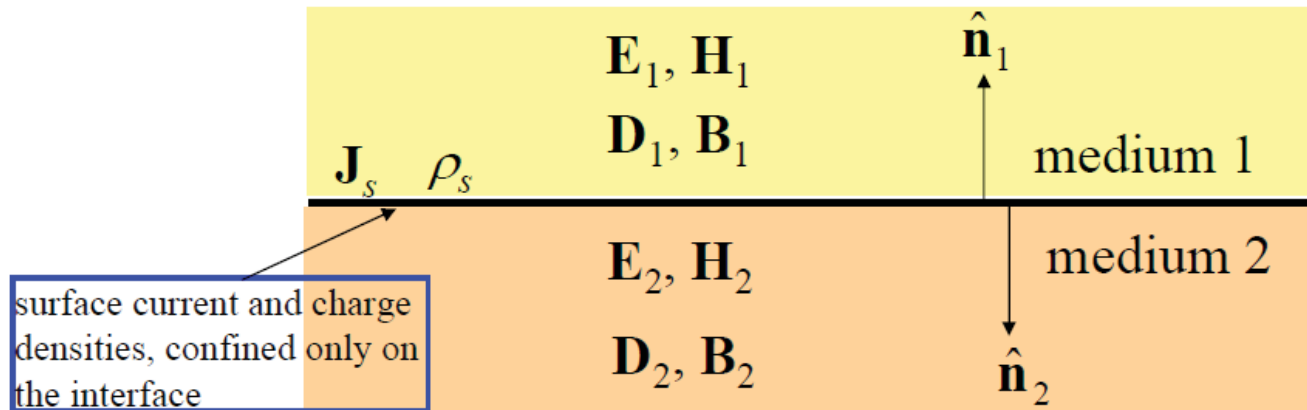
$\mathbf{D}$  – electric flux density

$\mathbf{B}$  – magnetic flux density

$\rho_{\text{ext}}$  – external charge density

$\mathbf{J}_{\text{ext}}$  – external current density

## Boundary conditions



For tangential components:

$$\begin{aligned}\hat{n}_1 \times (\mathbf{E}_1 - \mathbf{E}_2) &= 0 \\ \hat{n}_1 \times (\mathbf{H}_1 - \mathbf{H}_2) &= \mathbf{J}_s\end{aligned}$$

For normal components:

$$\begin{aligned}\hat{n}_1 \cdot (\mathbf{B}_1 - \mathbf{B}_2) &= 0 \\ \hat{n}_1 \cdot (\mathbf{D}_1 - \mathbf{D}_2) &= \rho_s\end{aligned}$$

If no external surface charges and currents:

$$E_{1t} = E_{2t}, H_{1t} = H_{2t}$$

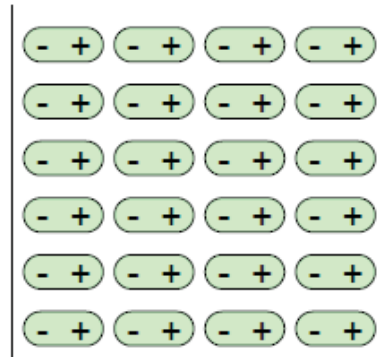
$$B_{1n} = B_{2n}, D_{1n} = D_{2n}$$

# Constitutive relations

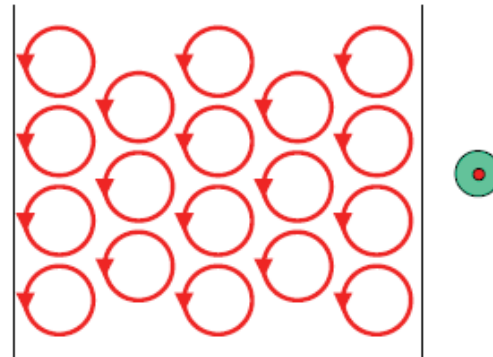
What are the relations between **E** and **D**, and **H** and **B**?

– determined by the EM response of materials

Polarization **P**



Magnetization **M**



relative permittivity

$$\mathbf{D} = \underline{\epsilon_0} \mathbf{E} + \mathbf{P} = \epsilon_0 \underline{\epsilon} \mathbf{E}$$

permittivity in  
vacuum

polarization

relative permeability

$$\mathbf{B} = \underline{\mu_0} \mathbf{H} + \mu_0 \mathbf{M} = \mu_0 \underline{\mu} \mathbf{H}$$

permeability in  
vacuum

magnetization

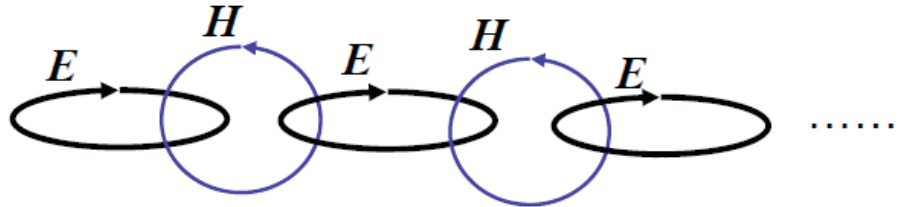
Meaning: total electric/magnetic flux density = flux from external field + flux due to material polarization/magnetization

# Wave equation

According to the curl equations:

$$\begin{aligned}\nabla \times \mathbf{E} &= -\partial \mathbf{B} / \partial t \\ \nabla \times \mathbf{H} &= \partial \mathbf{D} / \partial t\end{aligned}$$

changing electric field results in changing magnetic field and vice versa → **electromagnetic wave**



In homogeneous ( $\epsilon$  and  $\mu$  are spatially independent) media, we have:

$$\nabla \times \nabla \times \mathbf{E} = -\mu_0 \mu \frac{\partial}{\partial t} \nabla \times \mathbf{H} = -\epsilon_0 \mu_0 \epsilon \mu \frac{\partial^2 \mathbf{E}}{\partial t^2}$$



$$\nabla(\nabla \cdot \mathbf{E}) - \nabla^2 \mathbf{E} = -\frac{n^2}{c^2} \frac{\partial^2 \mathbf{E}}{\partial t^2}$$

**Wave Equation**

Light speed in vacuum:

$$c = \frac{1}{\sqrt{\epsilon_0 \mu_0}}$$

Refractive index:

$$n = \sqrt{\epsilon \mu}$$

## Harmonic field

Solution to wave equation:

$$\nabla(\nabla \cdot \mathbf{E}) - \nabla^2 \mathbf{E} = -\frac{\varepsilon\mu}{c^2} \frac{\partial^2 \mathbf{E}}{\partial t^2}$$

Henceforth, we consider only nonmagnetic media ( $\mathbf{M}=0, \mu=1$ )

→ time- and spatial-harmonic field:

$$\mathbf{E}(\mathbf{r}, t) = \mathbf{E}_0 \exp(i\mathbf{k} \cdot \mathbf{r} - i\omega t)$$

$\mathbf{k}$  – wave vector

(check this is a solution)

In this case,  $\nabla \square \square \mathbf{k}$ ,  $\partial / \partial t \square -\square\square$  (derive by yourself)

Therefore, the wave equation turns to

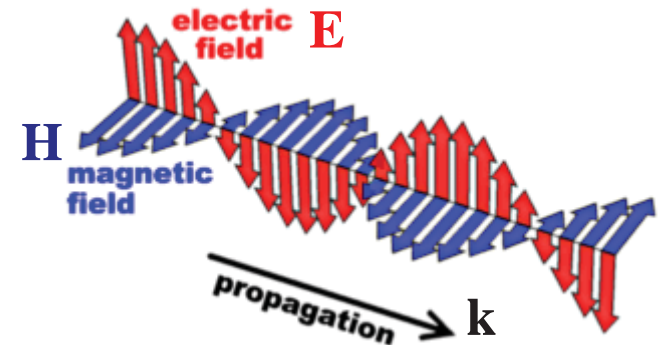
$$\mathbf{k}(\mathbf{k} \cdot \mathbf{E}) - \square^2 \mathbf{E} = -\varepsilon \frac{\square^2}{c^2} \mathbf{E}$$

- If transverse wave →  $\mathbf{k} \cdot \mathbf{E} = 0 \rightarrow$

$$k = \sqrt{\varepsilon} \frac{\omega}{c} \equiv nk_0$$

- If longitudinal wave →  $\mathbf{k}(\mathbf{k} \cdot \mathbf{E}) = k^2 \mathbf{E}$

→  $\varepsilon = 0$  ← more discussion in Lecture 5



## 2. Dispersion of materials

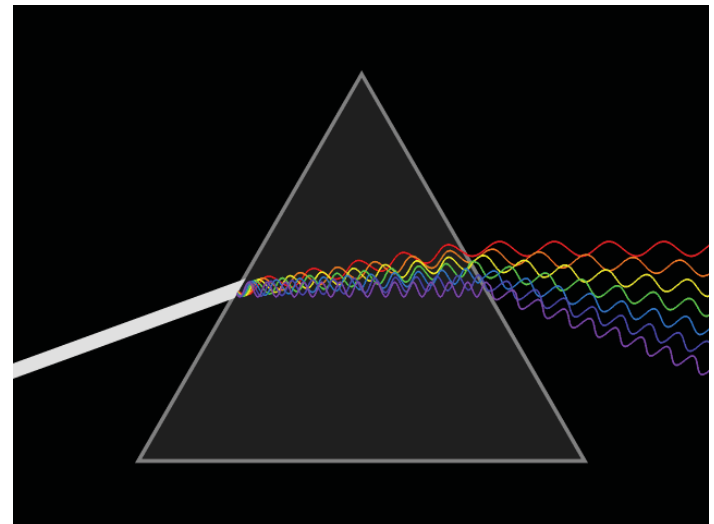
- In most media, the permittivity  $\epsilon$  is not a constant, which varies with respect to light frequency  $\omega$ , i.e.,  $\epsilon = \epsilon(\omega)$ .
- Therefore,  $\mathbf{D} = \epsilon(\omega)\mathbf{E}$  does not hold a linear relation; this frequency-dependent property is called **dispersion**.
- Any real material has more or less dispersion.

Example:

Observation of dispersion in prism

$$n(\omega) = \sqrt{\epsilon(\omega)}$$

refractive index is  $\omega$ -dependent



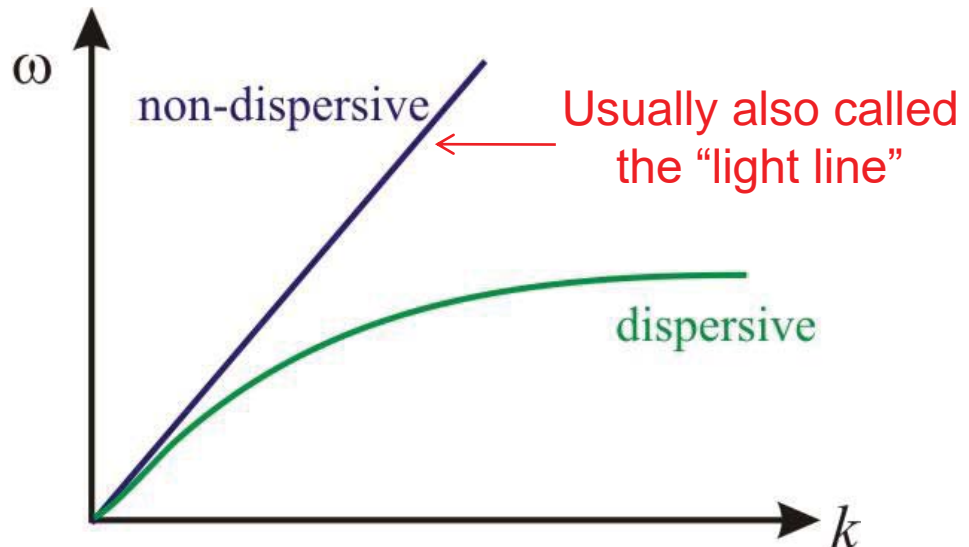
## $k$ - $\omega$ dispersion relation

For transverse EM wave:  $k = \sqrt{\epsilon} \frac{\omega}{c} \Rightarrow \omega = \frac{kc}{\sqrt{\epsilon}}$

- For **non-dispersive** media (for example, in vacuum),  $\epsilon$  is constant,  $k$ - $\omega$  relation is **linear**;
- Otherwise, for **dispersive** media,  $\epsilon$  is  $\omega$ -dependent so that the  $k$ - $\omega$  relation is **nonlinear**.

We can plot the  $k$ - $\omega$  dispersion relation:

**Important!!!**





## Phase and group velocities

Practical light cannot be ideally monochromatic (single  $\omega$ ).  
Multi-frequency components mixture → **wave package**



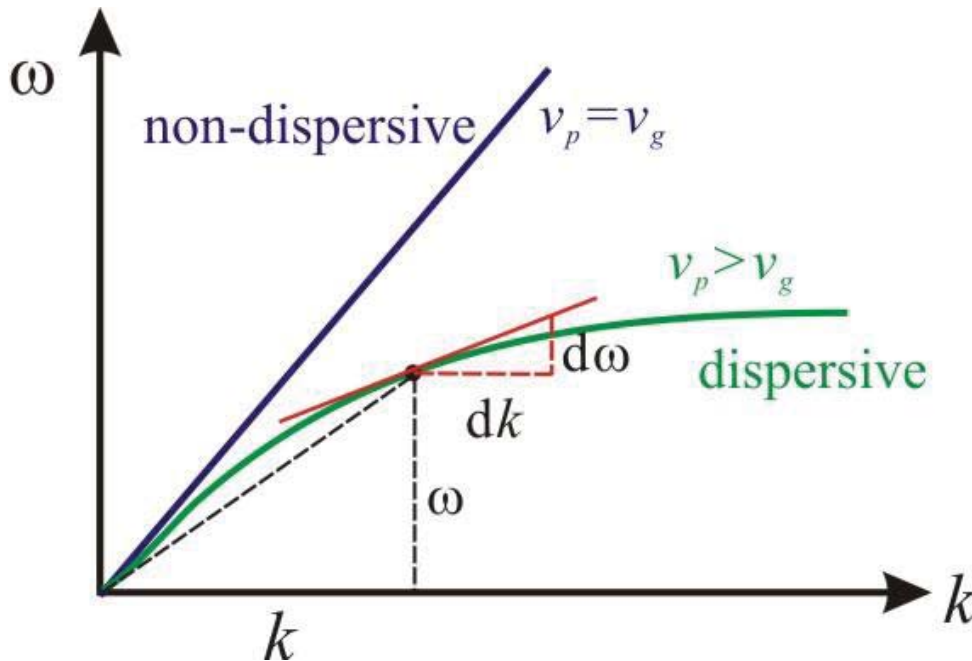
- **Phase velocity:** velocity of the oscillation or velocity of the equi-phase plane (red point)

$$v_p = \frac{1}{\sqrt{\epsilon_0 \mu_0 \epsilon \mu}} = \frac{c}{n} = \frac{\omega}{k} \quad \because \mathbf{E}(\mathbf{r}, t) = \mathbf{E}_0 \exp(\underbrace{i\mathbf{k} \cdot \mathbf{r} - i\omega t}_{\text{Let it be constant}})$$

- **Group velocity:** velocity of the waveform envelope or velocity of the equi-amplitude plane (green points)

$$v_g = \frac{d\omega}{dk} \quad (\text{derive by yourself. Hint: you may consider the simplest case of two-wave mixture.})$$

## Phase and group velocities in $k$ - $\omega$ plot



$$v_p = \frac{\omega}{k}, \quad v_g = \frac{d\omega}{dk}$$

– In non-dispersive media:

$$v_p = v_g$$

– In dispersive media:

$$v_p \neq v_g$$

- $v_g$  is the velocity of the transported signal or carried energy, therefore it must be  $v_g < c$  according to **Relativity**!
- There is no such restriction on  $v_p$ , therefore it is possible that  $v_p > c$

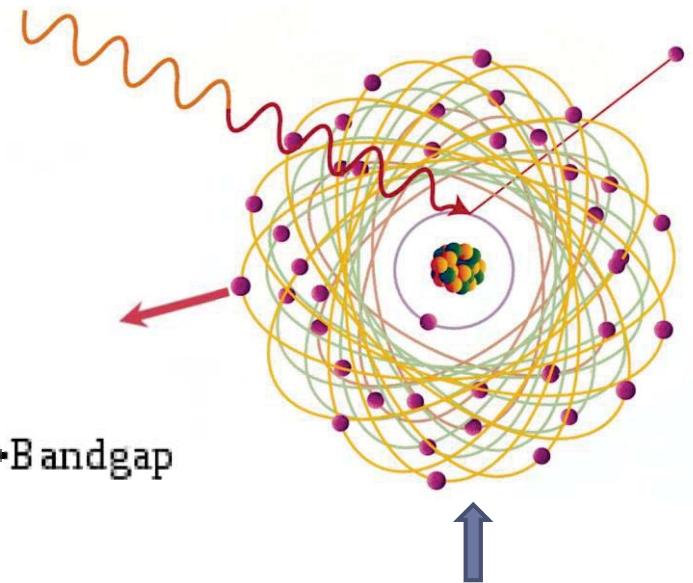
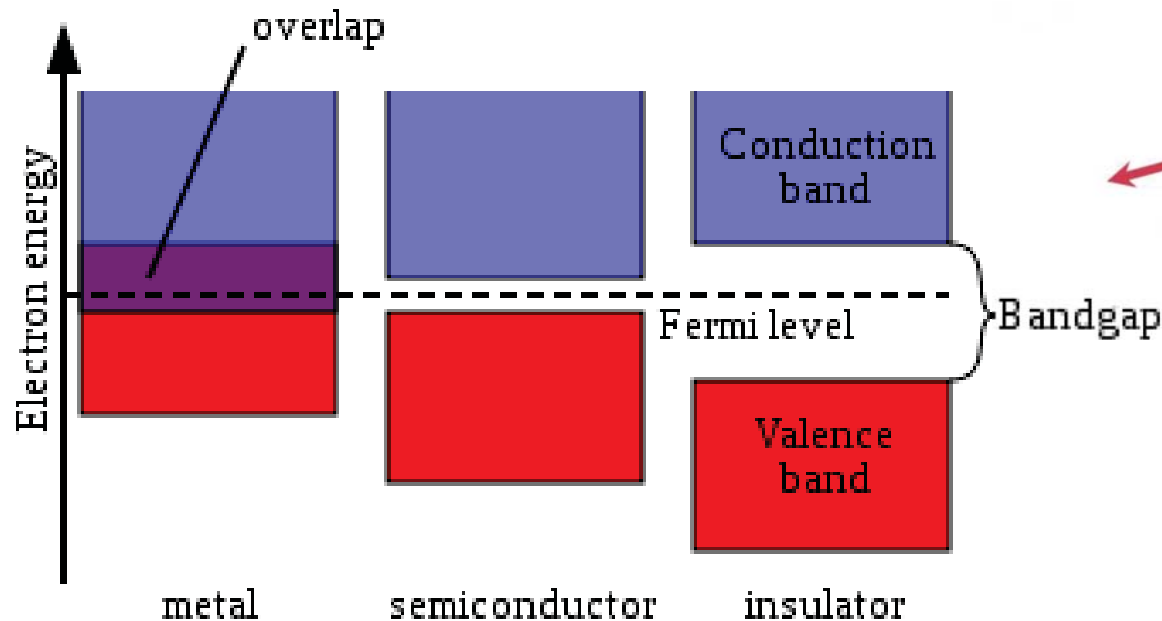
### 3. Micro- and macroscopic theories of materials

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- So far, we have known that the EM properties of materials are characterized by several **macroscopic** parameters:
  - permittivity  $\epsilon$
  - permeability  $\mu$
  - conductivity  $\sigma$  ( $\mathbf{J} = \sigma \mathbf{E}$ , characterizes how easy charges can move;  $\sigma = 0 \rightarrow$  insulator,  $\sigma = \infty \rightarrow$  perfect conductor)
- In nanophotonics, we aim to **engineer** these **macroscopic** parameters with artificial nanostructure.
- For this, we should know the **microscopic** origin of these macroscopic parameters for different materials:
  - Insulator (dielectric)
  - metal

# Free and bound electrons in materials

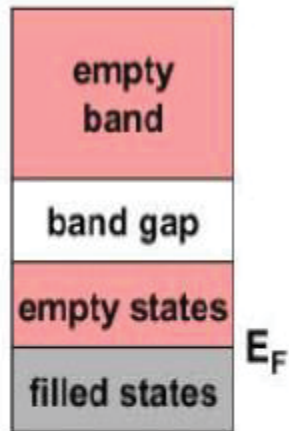
- In conduction band – **free electrons**
- In valence band – **bound electrons**



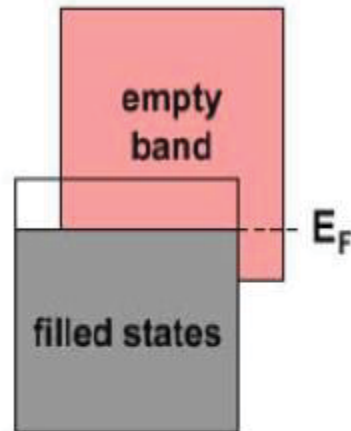
Electron cloud  
surrounding the  
atom cores

Energy needed for an electron to leap to higher  
energy level or the conduction band

# Typical band structures of materials



**metals:**  
available and  
filled states in the  
same band  
(Cu, Au, Ag)



**metals:**  
overlap between  
filled valence  
band and empty  
conduction band  
(Al, Mg)



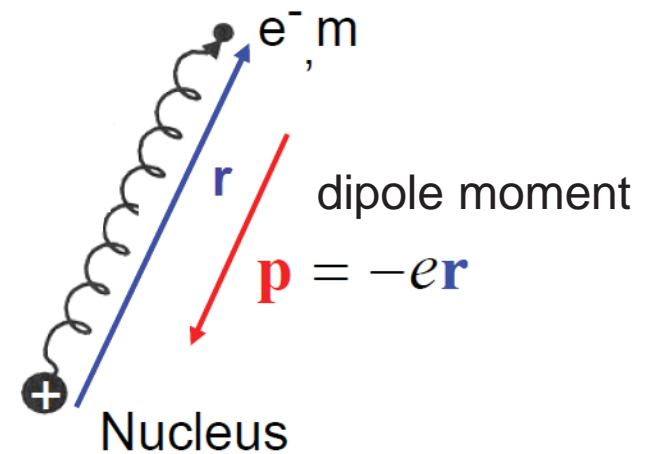
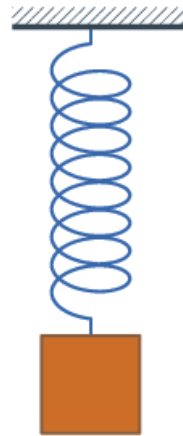
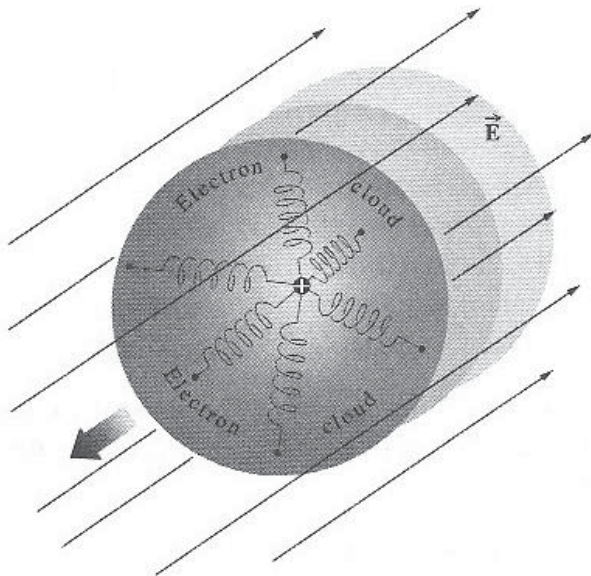
**semiconductors:**  
filled valence  
band separated  
from empty  
conduction band  
by a narrow band  
gap (< 2 eV)



**insulators:**  
filled valence  
band separated  
from empty  
conduction band  
by a large band  
gap (> 2 eV)

## Insulator – Lorentz model

- Insulator response is determined by the behavior of **bound electrons**.
- Under an external driving field  $\mathbf{E}$ , bound electrons can be treated as **harmonic oscillators** → **Lorentz model**



**Lorentz model**

How many forces on the bound electron?

- electric field force  $-e\mathbf{E}$
- damping force  $m\gamma\mathbf{v}$  ( $\gamma$ : damping frequency)
- restoring force  $K\mathbf{r}$  ( $K$ : restring-force constant)

Equation of motion:

$$m\ddot{\mathbf{r}} + m\gamma\dot{\mathbf{r}} + K\mathbf{r} = -e\mathbf{E}$$

For a time-harmonic stimulus  $\mathbf{E}(t) = \mathbf{E}_0 \exp(-i\omega t)$ , there is a time-harmonic solution  $\mathbf{r}(t) = \mathbf{r}_0 \exp(-i\omega t)$ , which is solved as:

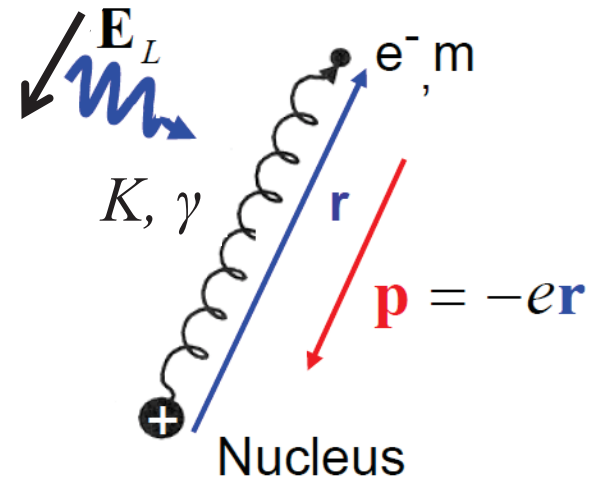
$$\mathbf{r} = \frac{e / m}{\omega^2 + i\omega\gamma - \omega_0^2} \mathbf{E}$$

$$\omega_0 = \sqrt{K / m}$$

(natural frequency of bound electron)

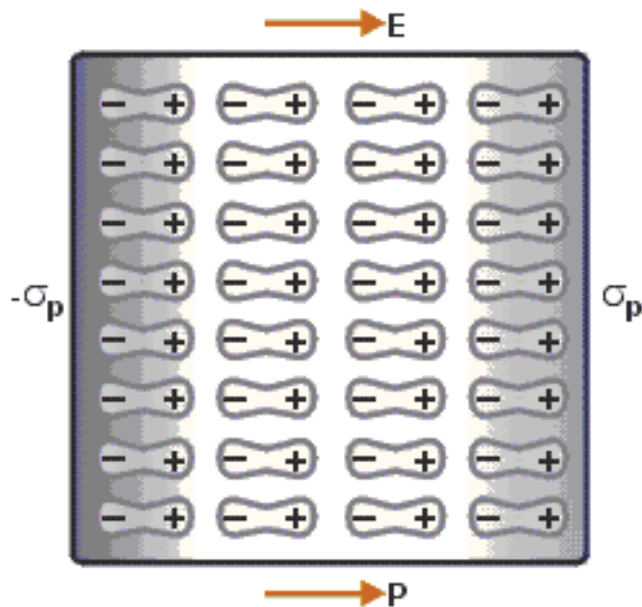
Then the **electric dipole moment** is:

$$\mathbf{p} = -e\mathbf{r} = \frac{e^2 / m}{\omega_0^2 - \omega^2 - i\omega\gamma} \mathbf{E}$$





Now we can connect the **microscopic electric dipole moment  $\mathbf{p}$**  to the **macroscopic polarization vector  $\mathbf{P}$** :



$N$  – density of electrons

$$\mathbf{P} = \sum \mathbf{p} = \frac{\sum e^2 / m}{\omega_0^2 - \omega^2 - i\omega\gamma} \mathbf{E}$$

$$\mathbf{D} = \epsilon_0 \mathbf{E} + \mathbf{P} = \epsilon_0 \epsilon \mathbf{E}$$

$\chi$ , electric susceptibility

$$\epsilon(\omega) = 1 + \frac{\omega_p^2}{\omega_0^2 - \omega^2 - i\omega\gamma}$$

$$\omega_p = \sqrt{\frac{Ne^2}{\epsilon_0 m}} \quad (\text{plasma frequency})$$

What does it mean?

More discussion in Lecture 5



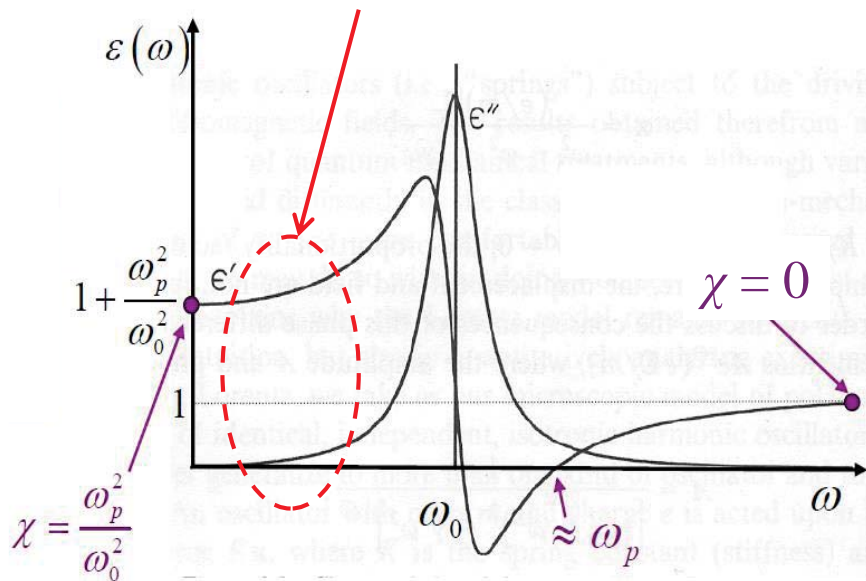
Now let's have a closer look at  $\varepsilon(\omega)$ :

If  $\gamma \neq 0$ , then  $\varepsilon$  is complex:  $\varepsilon = \varepsilon' + i\varepsilon'' = 1 + \frac{\omega_p^2}{\omega_0^2 - \omega^2 - i\omega\gamma}$

$$\varepsilon' = 1 + \chi' = 1 + \frac{\omega_p^2(\omega_0^2 - \omega^2)}{(\omega_0^2 - \omega^2)^2 + \gamma^2\omega^2}$$

$$\varepsilon'' = \chi'' = \frac{\omega_p^2\gamma\omega}{(\omega_0^2 - \omega^2)^2 + \gamma^2\omega^2}$$

Optical frequency  $\omega_0 \gg \omega \gg \gamma$



Several typical points:

- When  $\omega \rightarrow 0$ :  $\varepsilon' \rightarrow 1 + \frac{\omega_p^2}{\omega_0^2}$ ,  $\varepsilon'' \rightarrow 0$
- When  $\omega = \omega_0$ :  $\varepsilon' = 1$ ,  $\varepsilon'' \rightarrow \text{maximum}$
- When  $\omega \approx \omega_p$ :  $\varepsilon' = 0$
- When  $\omega \rightarrow \infty$ :  $\varepsilon' \rightarrow 1$ ,  $\varepsilon'' \rightarrow 0$

Note: usually  $\omega_p \gg \omega_0 \gg \gamma$ ,  $\gamma \sim 100\text{THz}$

# Refractive index $\hat{\epsilon}(\omega)$ – characterizing EM wave propagation

For nonmagnetic media:

$$n = n' + in'' = \sqrt{\epsilon}$$

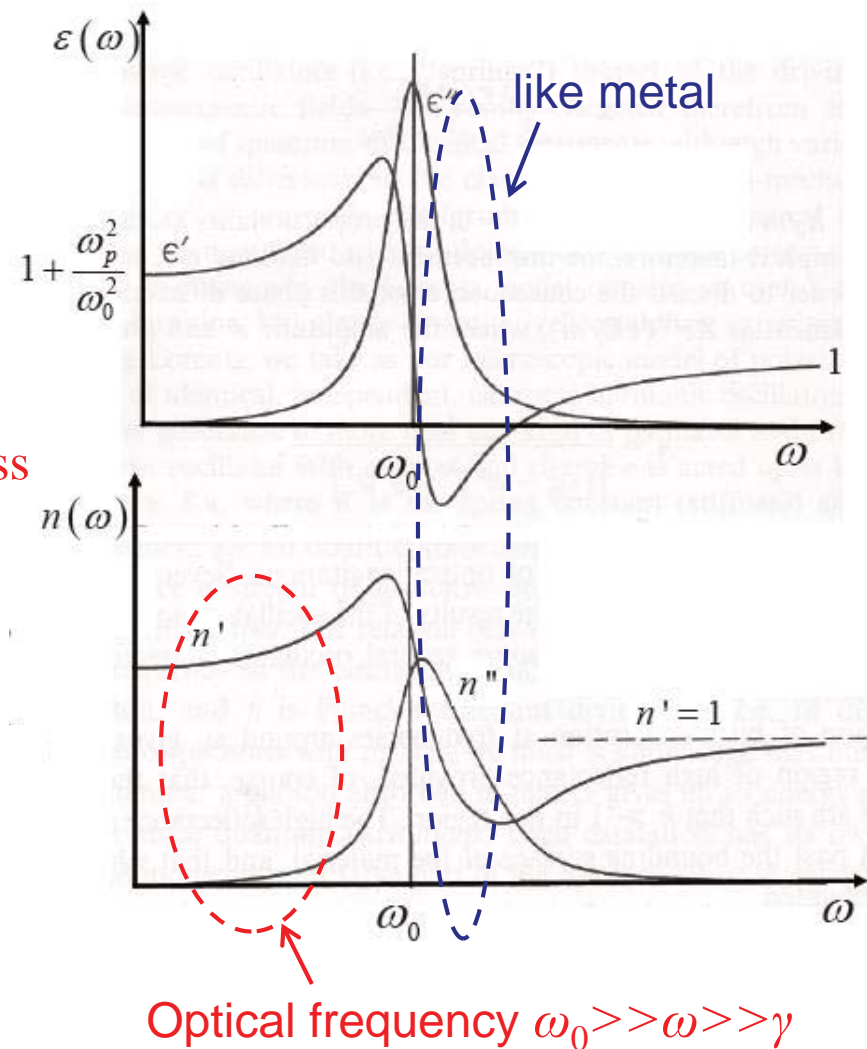
$$n' = \sqrt{\frac{\sqrt{\epsilon'^2 + \epsilon''^2} + \epsilon'}{2}}$$

$$n'' = \sqrt{\frac{\sqrt{\epsilon'^2 + \epsilon''^2} - \epsilon'}{2}}$$

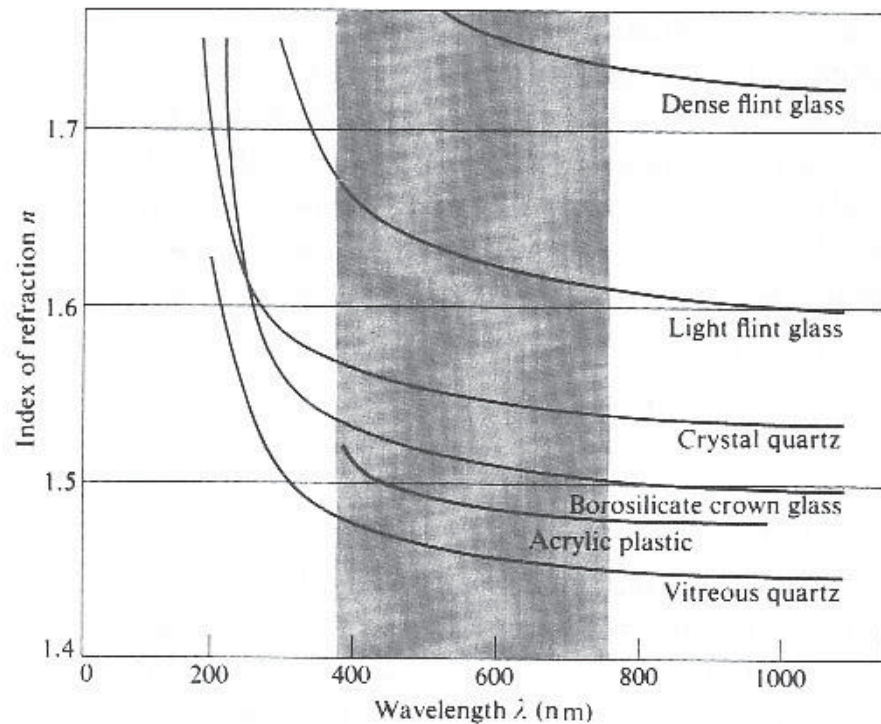
- When  $\omega \ll \omega_0$ :  
high  $n' \rightarrow$  low  $v_p = c/n'$ ,  $n'' \approx 0 \rightarrow$  lossless  
**↑ This is the so-called insulator!**

- When  $\omega \sim \omega_0$ :  
rapidly varying  $n' \rightarrow$  strong dispersion  
 $n''$  high  $\rightarrow$  large absorption  
**↑ like metal**

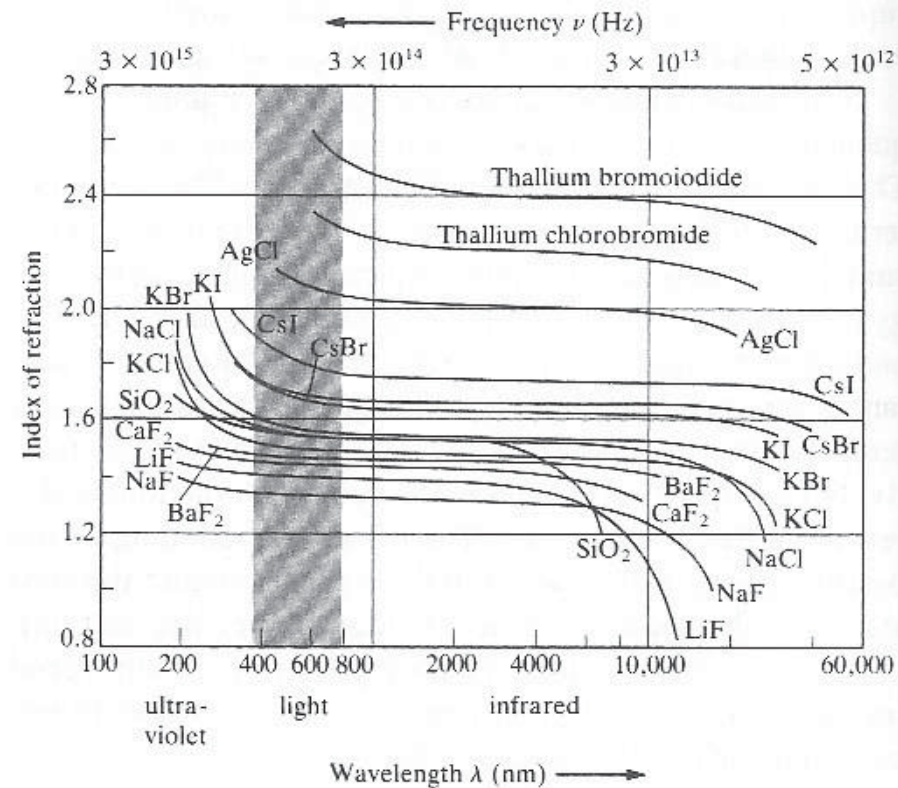
- When  $\omega \gg \omega_0$ :  
 $n' \approx 1 \rightarrow v_p \approx c$ ,  $n'' = 0 \rightarrow$  lossless  
**↑ like vacuum or air**



## Refractive indices of some typical insulators (at optical frequency)



**Figure 3.40** The wavelength dependence of the index of refraction for various materials.



**Figure 3.42** Index of refraction versus wavelength and frequency for several important optical crystals. (Adapted from data published by The Harshaw Chemical Co.)

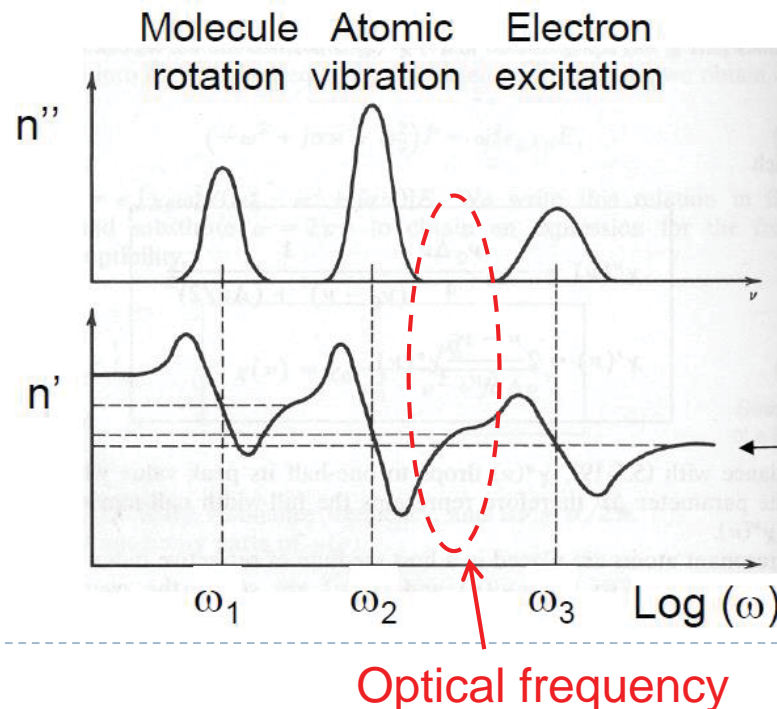
But, realistic media usually have multiple resonances:

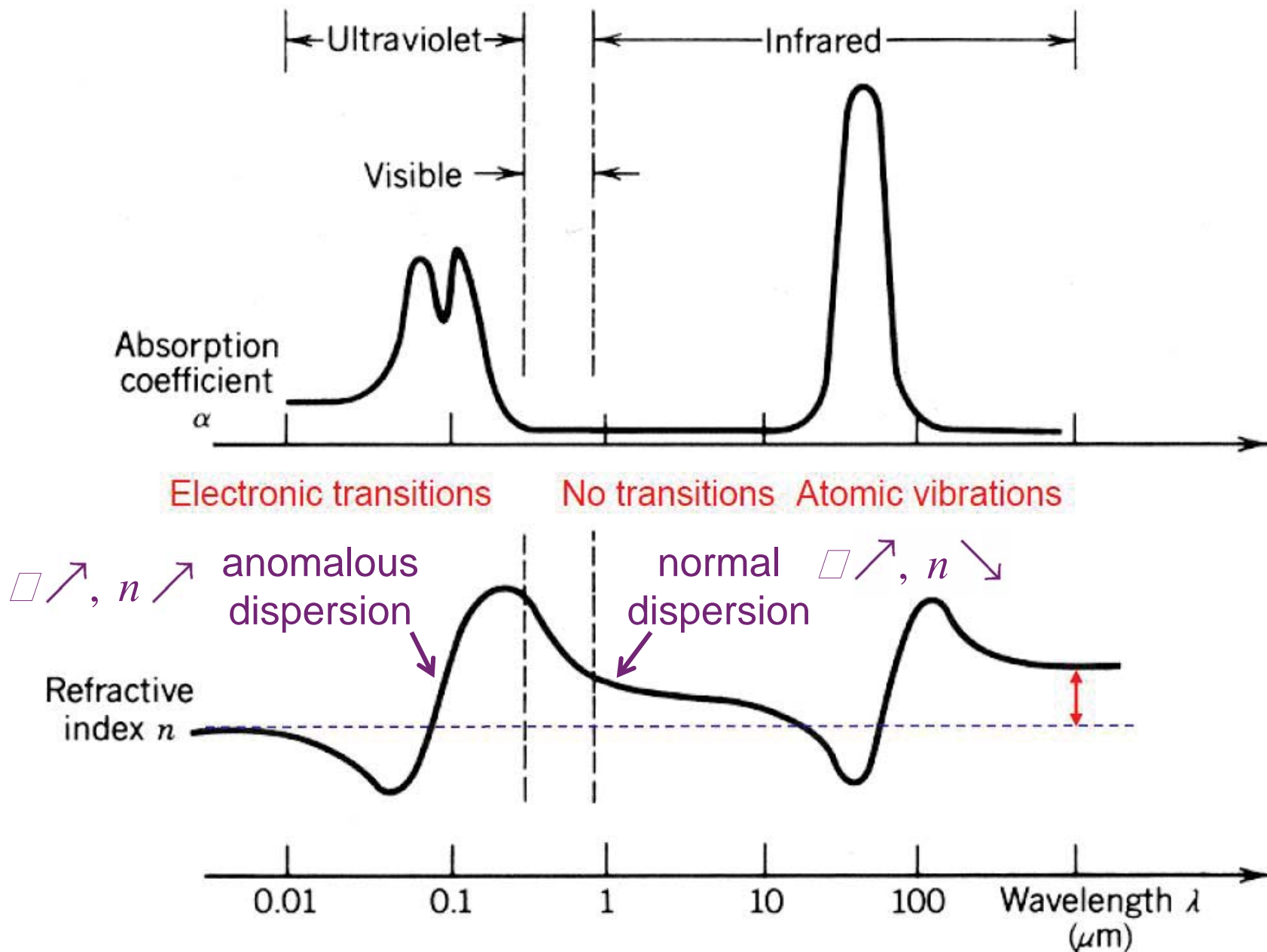
- Resonances occur due to motion of molecules (**very low**  $\omega_0$ ), atoms (**low**  $\omega_0$ ), or electron transitions (**high**  $\omega_0$ )
- Therefore, the Lorentz model should be modified by taking into account **multiple resonance terms**:

$$\varepsilon = 1 + \sum_j \frac{\omega_{jp}^2}{\omega_{j0}^2 - \omega^2 - i\omega\gamma_j}$$

$$\left( \omega_{jp}^2 = \frac{N_j e^2}{\varepsilon_0 m}, \quad \omega_{j0}^2 = \frac{K_j}{m} \right)$$

Schematic of realistic  $n'$   
and  $n''$  with multiple  
resonances





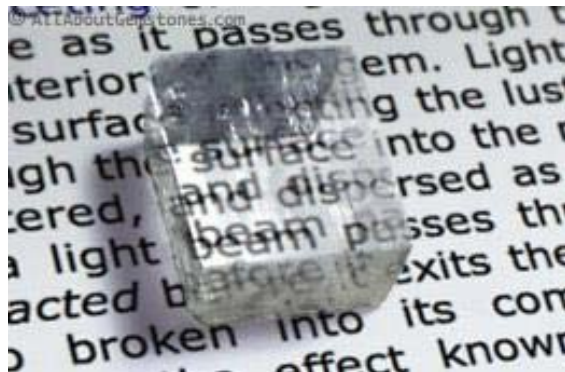
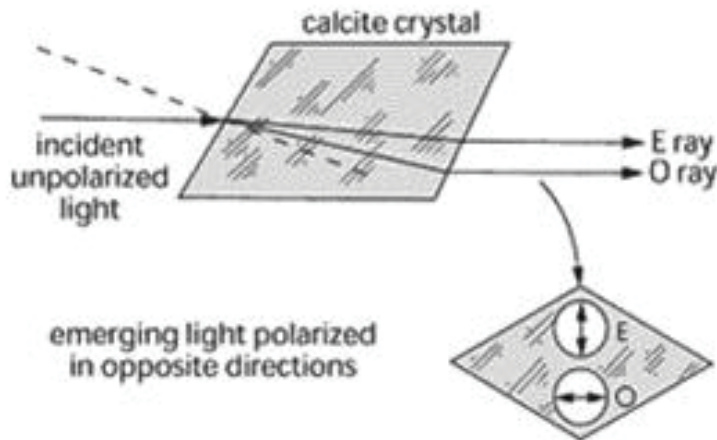
## 4. Example of engineering light-matter interaction with nanostructures – **form birefringence**

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- Having understood the EM response of materials from both microscopic and macroscopic aspects, we can try to **engineer (alter/modify/tailor)** the light-matter interaction with artificial nanostructures.
- In this course, we are mostly talking about this kind of things ...
- Here, let's see a typical example – **form birefringence**
- More will be discussed in the succedent lectures

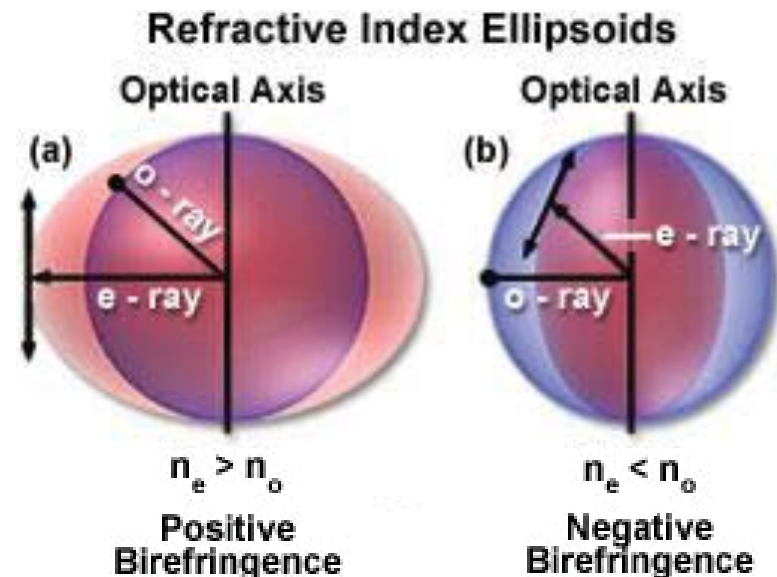


**Birefringence** (also called **double refraction**): the splitting of a light ray into two rays (an **ordinary** one and an **extraordinary** one) in an optically anisotropic medium such as calcite.



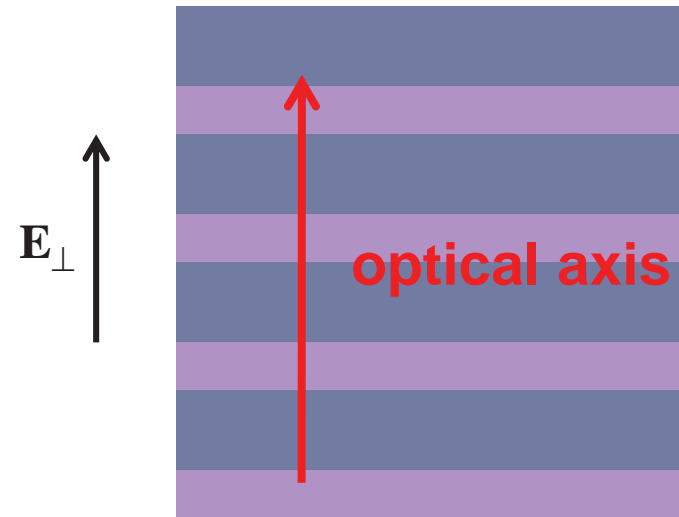
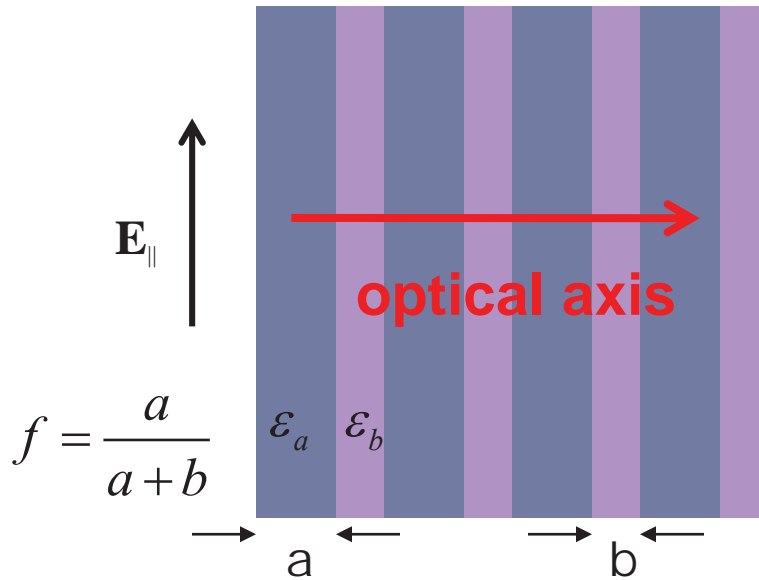
**Reason:** refractive index is directionally dependent

$$\Delta n = n_e - n_o$$



# What happens for stacked isotropic dielectric nano-plates?

molecule size  $\ll a, b \ll \lambda$  (can be treated as **effective media**)



The effective permittivity can be derived as:

**o-ray**  $\rightarrow \epsilon_{\parallel} = f\epsilon_a + (1-f)\epsilon_b$

**e-ray**  $\rightarrow \epsilon_{\perp} = \frac{\epsilon_a \epsilon_b}{f\epsilon_b + (1-f)\epsilon_a}$

**Uniaxial  
crystal!**

(Homework: derive these expressions and prove that it is a negative uniaxial crystal)

**Form birefringence** – birefringence/anisotropy induced by the structural arrangement rather than the composing materials themselves.



# Summary

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- ▶ Focus of this lecture: **understanding light-matter interaction**
- ▶ Electromagnetic wave:  
Review of Maxwell's equations, boundary conditions, constitutive relations, wave equation, time- and spatial-harmonic field
- ▶ Dispersion of materials:  
dispersion (frequency-dependent effect), understand  $k$ - $\omega$  plot, physical meaning of phase and group velocities
- ▶ Microscopic and macroscopic theories of materials:  
EM response of **insulator: Lorentz model** (bound electrons)
- ▶ Example of engineering light-matter interaction with artificial nanostructures: **form birefringence**