

Macroscopic Electrodynamics

Introduction to Dielectrics

Think about the interaction of the electromagnetic field with matter. The quanta of the EM field are photons, which can be emitted or absorbed ^{or scattered} by ~~the~~ charged particles such as electrons or protons (or atoms - made of charged particles). The passage of EM radiation through a medium like a solid is a complicated process ~~involving~~ involving the continual absorption, and re-radiation of photons. scattering,

Even when we think about this process classically, it remains quite complicated. The solid body contains charges and current loops which interact with the EM waves, absorbing, ~~re-emitting~~ scattering, or radiating them.

are the ~~permittivity~~ electric and magnetic permeabilities ϵ and μ , and it also involves a new set of fields, \vec{P} the polarization and \vec{M} the magnetization, which describe the

space-time evolution of the electrodynamic properties of matter. ^(course-ground) In a formal sense this effective theory is different from the kind of electrodynamics we have heretofore considered - we can call it "macroscopic electrodynamics." We treat all our fields (\vec{E} , \vec{B} , \vec{P} , \vec{M} , and the two useful derived fields \vec{D} , \vec{H}) as if they were garden variety fields, defined at a point, continuous, differentiable --- (occasionally you have to be careful) and they encode effects of physics at short distance scales as seen by long distance scales. All done by averaging over short distance physics.

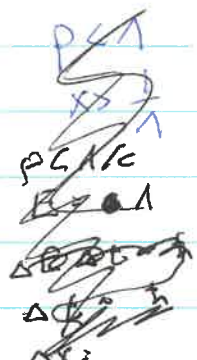
2nd ingredient - accuracy - How accurate is

EFT? depends on how many terms you keep, how accurately we do averaging. Example follows..

"integrating out"

And yet - over distances large compared to atomic sizes, this complicated process can be parameterized by a few numbers (possibly frequency dependent) - a dielectric constant $\epsilon(\omega)$, permeability μ , conductivity σ ... We can short-circuit the hard problem of scattering etc by ~~parameterizing~~ ^{computing} these numbers, and expressing the propagation in terms of them. Or - if the calculation is too hard - we perform experiments to measure these auxiliary quantities once, then use these numbers to make predictions for other ~~propagated~~ processes.

There is a modern language to describe this procedure. We say that we are constructing an "effective field theory" which is not a fundamental theory of Nature valid on all length scales, instead it has a built-in cutoff, ~~a constant~~ ^{in energy} scale Λ or ~~an~~ space cutoff $\sim \hbar/c\Lambda$, and it is not supposed to describe scales shorter than $1/\Lambda$. Often these effective theories ~~inherit~~ ^{properties} inherit parameters from a "more fundamental theory," but they also contain parameters which within the effective field theory are supposed to be fundamental, but which might represent ~~errors~~ ^{corrections} at the long distance / low momentum tail of the "real physics" (whatever it is.) In ~~the~~ classical electrodynamics, ~~and~~ μ ~~are~~ examples of these second class of parameters. Typically these theories break down as $\langle k \rangle \rightarrow \Lambda$ and ~~one's experiments probe distances close to the cutoff~~

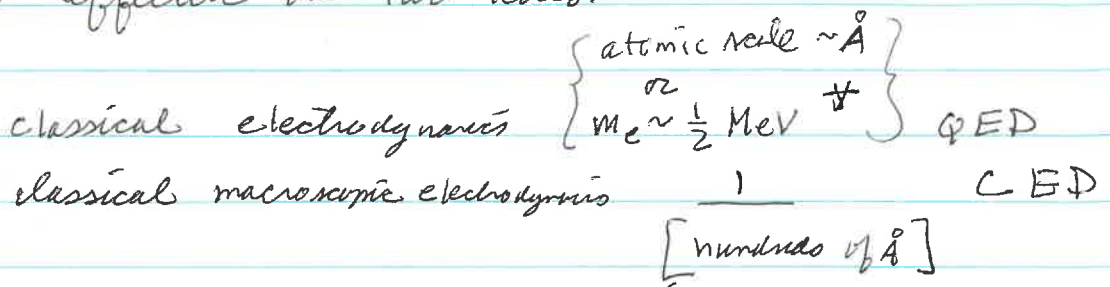


$E \ll \Lambda$
 $p \ll \frac{\Lambda}{c}$
 $\Delta x \Delta p \sim \hbar$
 $\Delta x \sim \frac{\hbar}{\Lambda}$

Examples of "effective field theories"

the effective theory	Λ	the more fundamental theory
classical general relativity	10^{19} GeV (Planck scale) $M_{Pl}^2 = \hbar c^3 / G_N$	strings quantum gravity
the standard model Gauge unification theories?	10^{16} GeV	"
the standard model	100 to 1000 GeV	?? SUSY GUT?
Weak interactions in isolation QED in isolation	} ~ 100 GeV or M_W	electroweak
pions and nucleons ⋮	~ 1 GeV ⋮	QCD ⋮
classical macroscopic electrodynamics	 [hundreds of Å]	QED

In fact, classical macroscopic electrodynamics is effective on two levels:



* plus a long story ~~and~~ incomplete story

$$\frac{GM^2}{R} = E = \frac{\hbar c}{\lambda}$$

$\lambda = R$

$$M^2 = \frac{\hbar c}{G}$$

$$(M_{Pl})^2 = \frac{\hbar c^3}{G}$$

From Microscopic to Macroscopic Electrodynamics

Following Jackson's notation, sec 6.6,

Maxwell's eqns are

$$\begin{aligned} \vec{\nabla} \cdot \vec{e} &= \eta / \epsilon_0 & \vec{\nabla} \cdot \vec{b} &= 0 \\ \vec{\nabla} \times \vec{e} + \frac{\partial \vec{b}}{\partial t} &= 0 & \vec{\nabla} \times \vec{b} - \frac{1}{c^2} \frac{\partial \vec{e}}{\partial t} &= \mu_0 \vec{j} \end{aligned}$$

 $b, e \equiv$ microscopic electric + magnetic fields $\eta, \vec{j} \equiv$ " " charge + current densityall fluctuating over distances 10^{-8} cmtimes $10^{-13} - 10^{-15}$ sec $10^{23 \pm}$ atoms/cm³

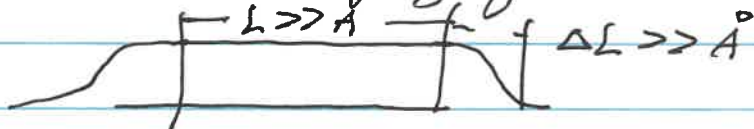
(Keeping track of all these DoF's is overkill.)

Instead, do a spatial average over

 $L \sim$ few hundred \AA . (Want to deal with

light in range of kV, so no time avg)

$$\langle F(\vec{x}, t) \rangle = \int d^3x' f(x') F(x-x', t)$$

 $f(x)$ a smoothing function

$$\frac{\partial}{\partial t} \langle F(x, t) \rangle = \left\langle \frac{\partial F}{\partial t} \right\rangle$$

$$\frac{\partial}{\partial x_i} \langle F(x, t) \rangle = \int d^3x' f(x') \frac{\partial F(x-x', t)}{\partial x_i}$$

$$= \left\langle \frac{\partial F}{\partial x_i} \right\rangle$$

Now define macroscopic fields (by averaging with f)

$$\vec{E}(x,t) = \langle \vec{e}(x,t) \rangle$$

$$\vec{B}(x,t) = \langle \vec{b}(x,t) \rangle$$

The homogeneous eqns are simple

$$\vec{\nabla} \cdot \vec{B} = 0 \quad \vec{\nabla} \times \vec{E} + \frac{1}{c} \frac{\partial \vec{B}}{\partial t} = 0$$

inhomogeneous ones less so!

$$\epsilon_0 \vec{\nabla} \cdot \vec{E} = \langle \rho(x,t) \rangle$$

$$\frac{1}{\mu_0} \vec{\nabla} \times \vec{B} - \frac{\partial \vec{E}}{\partial t} = \langle \vec{j}(x,t) \rangle$$

Let's work on the two RHS's. We make a distinction between "bound charges" (which stay put, more or less in the material) and "free charges" which move around - and can be injected or removed from the material

$$\rho = \rho_{\text{free}} + \rho_{\text{bound}}$$

$$\rho_{\text{free}} = \sum q_i \delta(x-x_i)$$

$$\rho_{\text{bound}} = \sum_{n \text{ molecules}} \rho_n(x,t) = \sum_n \sum_j q_j \delta(x-x_j(n))$$

Let's average molecular charges
 $= \int d^3x' f(x') \rho_n(x-x',t)$

$$\langle \rho_n(x,t) \rangle = \int d^3x' f(x') \sum_j q_j \delta(x-x'-x_{jn}-x_n)$$



interatomic spring
 molecules - could be \AA or hundreds of \AA ...

$$= \sum_{j(n)} q_j f(x-x_n-x_{jn})$$

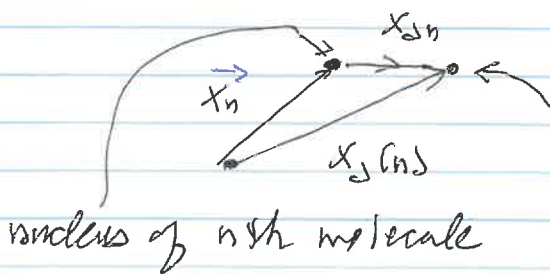
$$\eta = \eta_{\text{free}} + \eta_{\text{bound}}$$

$$\eta_{\text{free}} = \sum_i g_i \delta(x - x_i)$$

$$\rho_{\text{free}} = \langle \eta_{\text{free}} \rangle = \langle \sum_j g_j \delta(x - x_j) \rangle$$

$$\eta_{\text{bound}} = \sum_{n \text{ molecules}} \eta_n(x, t) = \sum_n \sum_{j \in n} g_j \delta(x - x_j(n))$$

long
call this g_j



offset of j th electron in n th molecule changes
 $\vec{x}_j(n) = \vec{x}_n + \vec{x}_{jn}$

For n th molecule (no sum on n yet)

$$\langle \eta_n(x, t) \rangle = \int d^3x' f(x') \eta_n(x - x', t)$$

$$= \int d^3x' \sum_{j \in n} g_j \delta^3(x - x' - x_{jn} - x_n) f(x')$$

$$= \sum_{j \in n} g_j f(\underbrace{x - x_n - x_{jn}}_{\substack{\text{nonzero over some} \\ \text{big range}}})$$

$$= \sum_{j \in n} g_j \left[f(x - x_n) - \vec{x}_{jn} \cdot \vec{\nabla}_x f(x - x_n) + \frac{1}{2} \sum_{\mu\nu} x_{jn}^\mu x_{jn}^\nu \frac{\partial^2 f}{\partial x_\mu \partial x_\nu} + \dots \right]$$

Now sum over n :

The first term contributes

$$\begin{aligned} \langle \eta_{\text{bound}} \rangle &= \left\langle \sum_n \underbrace{q_j^{(n)}}_{\text{total charge } q_n} f(x-x_n) \right\rangle \\ &= \left\langle \sum_n q_n \delta(x-x_n) \right\rangle \\ &= \rho_b \quad \text{total bound charge} \end{aligned}$$

The 2nd term

$$\begin{aligned} & - \sum_n \sum_{j \in n} q_j \vec{x}_{jn} \cdot \vec{\nabla}_x f \\ &= - \vec{\nabla}_x \cdot \sum_n \sum_{j \in n} \vec{x}_{jn} q_j \cdot f \quad \text{exchange } \sum \leftrightarrow \nabla \\ &= - \vec{\nabla}_x \cdot \left\langle \sum_n \vec{P}_n \right\rangle \quad \text{polarization} \end{aligned}$$

$$\langle \eta_b \rangle = - \vec{\nabla} \cdot \vec{P}(x)$$

New field

$$\begin{aligned} & \equiv \text{polarization } \vec{P} \equiv \text{average dipole moment} \\ & \vec{P}_n = \sum_{j \in n} \vec{x}_{jn} q_j \equiv \text{molecular dipole moment} \end{aligned}$$

The 3rd term involves the molecular quadrupole moment

$$(\varphi_n)_{\mu\nu} = \left\langle \sum_{j \in n} q_j x_{jn}^\mu x_{jn}^\nu \right\rangle$$

$$\text{and } \langle \eta_b \rangle = \dots + \frac{\partial^2}{\partial x_\mu \partial x_\nu} \varphi_{\mu\nu}(x, t)$$

and we re-arrange the $\nabla \cdot E$ equation to write with the charge on one side

$$\nabla \cdot [\epsilon_0 \vec{E} + \vec{P} - \sum_v \frac{\partial \phi_{jv}}{\partial x_j} + \dots] = \rho_{\text{macro}}$$

$$\rho_{\text{macro}} = \rho_f + \rho_{\text{bound}}$$

generally negligible for atomic systems - or larger -

which is why we usually don't see them.

so we have the "usual" equation
(note it's not fundamental!)

$$\nabla \cdot \vec{D} = \rho_{\text{macro}}$$

where

$$\vec{D} = \epsilon_0 \vec{E} + \vec{P}$$

ρ_f = "macroscopic" charge density

(the "free" charge density of undergrad E & M)

Now for magnetism ...

$$\vec{j}(\vec{x}, t) = \sum_j q_j \vec{v}_j \delta(\vec{x} - \vec{x}_j(t))$$

Molecular currents average into (for n th molecule)

$$\langle \vec{j}_n(\vec{x}, t) \rangle = \sum_{\Delta L_n} q_n (\vec{v}_{on} + \vec{v}_n) \delta(\vec{x} - \vec{x}_n - \vec{x}_{on})$$

A (my story, similar to what we just did

~~I don't have the strength to grind this out~~

For component l

$$\langle j_{le}(\vec{x}, t) \rangle = \langle j_{le}(\vec{x}, t) \rangle + \frac{\partial}{\partial t} [D_{le}^{-\epsilon_0} E_{le}] + (\vec{\nabla} \times \vec{M})_{le} + \dots$$

where

$$\vec{m}_n = \sum_{j \in n} \frac{q_j}{2} (\vec{x}_{jn} \times \vec{v}_{jn})$$

\equiv molecular magnetic moment in MKS

$$\text{CGS } \vec{m}_n = \sum_{j \in n} \frac{q_j}{2c} (\vec{x}_{jn} \times \vec{v}_{jn})$$

$$\vec{M}(\vec{x}, t) = \left\langle \sum_n \vec{m}_n \delta(\vec{x} - \vec{x}_n) \right\rangle \equiv \text{magnetization}$$

or plug into

$$\frac{1}{\mu_0} \nabla \times \vec{B} - \epsilon_0 \frac{\partial \vec{E}}{\partial t} = \vec{J} + \frac{\partial}{\partial t} (\vec{D} - \epsilon_0 \vec{E}) - \nabla \times \vec{M}$$

$$\nabla \times \vec{H} - \frac{\partial \vec{D}}{\partial t} = \vec{J}$$

$$\vec{H} = \frac{1}{\mu_0} \vec{B} - \vec{M} \quad \text{or} \quad \vec{B} = \mu_0 (\vec{H} + \vec{M})$$

$\vec{J} =$ macroscopic current density

- Derivation assumes medium isn't moving
- Says nothing about spin (must add by hand to \vec{M})

Recap: Macroscopic Electrodynamics

$$\nabla \cdot \mathbf{B} = 0 \quad \nabla \times \mathbf{E} + \frac{\partial \mathbf{B}}{\partial t} = 0$$

$$\nabla \cdot (\epsilon_0 \mathbf{E} + \mathbf{P}) = \rho_{\text{macro}} \text{ or } \rho_{\text{free}} = \nabla \cdot \mathbf{D}$$

one bond

\mathbf{P} = polarization = $\frac{\text{electric dipole moment}}{\text{unit volume}}$

$$\vec{J}(x,t) = \sum_n \sum_{\alpha n} q_{\alpha} (\vec{v}_n + \vec{v}_{\alpha n})$$

For component e

$$\langle J_e(x,t) \rangle = J_M(x,t)_e + \frac{\partial P_e(x,t)}{\partial t} + (\nabla \times \vec{M})_e$$

$\vec{M}(x,t)$ = Magnetization = $\frac{\text{magnetic dipole moment}}{\text{unit vol}}$

$$\vec{M} = \left\langle \sum_n \vec{m}_n \delta(\vec{x} - \vec{x}_n) \right\rangle$$

$$\vec{m}_n = \sum_{\alpha n} \frac{q_{\alpha}}{2} (\vec{x}_{\alpha n} \times \vec{v}_{\alpha n}) \text{ MKS}$$

$$= \sum_{\alpha n} \frac{q_{\alpha}}{2c} (\vec{x}_{\alpha n} \times \vec{v}_{\alpha n}) \text{ CGS}$$

c says nothing about spin - put in by hand!

$$\frac{1}{\mu_0} \vec{\nabla} \times \vec{B} - \epsilon_0 \frac{\partial \vec{E}}{\partial t} = \vec{J}_M + \frac{\partial}{\partial t} (\vec{D} - \epsilon_0 \vec{E})$$

$$+ \vec{\nabla} \times \vec{M}$$

or

$$\vec{\nabla} \times \vec{H} - \frac{\partial \vec{D}}{\partial t} = \vec{J}_M$$

or
board

$$\vec{H} = \frac{1}{\mu_0} \vec{B} - \vec{M} \quad \text{or} \quad \vec{B} = \mu_0 (\vec{H} + \vec{M})$$

so 2 new fields (\vec{P}, \vec{M})
 2 useful constructs (\vec{D}, \vec{H})
 e, \vec{J} macroscopic.

Back to electrostatics!

Dielectric Possibilities

Recall that $\nabla \times \vec{E} = 0, \nabla \cdot \vec{E} = \rho_{free} / \epsilon_0$

~~$\vec{D} = \epsilon_0 \vec{E} + \vec{P}$~~ ~~$\nabla \cdot \vec{D} = \rho_{free}$~~

~~$\vec{D} = \epsilon_0 \vec{E} + \vec{P}$~~ ~~$\nabla \cdot \vec{D} = \rho_{free}$~~ but $\vec{D} = \epsilon_0 \vec{E} + \vec{P}$

Need relationships between \vec{D} and \vec{E} to proceed...

Common ones are...

— a) Ferroelectricity: $\vec{P} \neq 0$ when $\vec{E} = 0$

Typically \vec{E} specified external to problem
or analogy of "hard ferromagnet"

— b) linear isotropic material $\vec{P} \propto \vec{E}$

$$\vec{P} = \epsilon_0 \chi_e \vec{E}$$

$\chi_e \equiv$ electric susceptibility

$E =$ total field (external + induced)

i.e. \vec{P} vs \vec{E} is linear } not true in general
 $\vec{P} \parallel \vec{E}$

$$\text{then } \vec{D} = \epsilon_0 (1 + \chi_e) \vec{E} \equiv \epsilon \vec{E}$$

$\frac{\epsilon}{\epsilon_0} =$ "dielectric constant"

"relative electric permittivity"

Note if entire ~~medium~~ medium filled with material of dielectric constant ϵ , independent of position.

$$\nabla \cdot \vec{D} = \rho_{free} \Rightarrow \nabla \cdot \vec{E} = \rho_{free} / \epsilon$$

Same as free space, but $\epsilon_{free} \Rightarrow \epsilon_{free} / \epsilon$ is renormalized by polarization of medium

$\epsilon_{free} \rightarrow$  $\Rightarrow \epsilon' = \epsilon_{free} / \epsilon < \epsilon$

— c) Anisotropic media (crystals)

$$D_i = \epsilon_{ij} E_j \quad \epsilon_{ij} = \text{dielectric tensor, } \epsilon_{ij} = \epsilon_{ji}$$

— d) "other"

~~Josephson 5.3.1 → the solution -~~

Always

$$\nabla \cdot \vec{E} = \frac{\rho_{\text{tot}}}{\epsilon_0}$$

$$\nabla \times \vec{E} = 0 \quad \leftarrow \quad \vec{E} = -\nabla \phi$$

$$\nabla \cdot \vec{D} = \rho_{\text{macro}}$$

$$\vec{D} = \epsilon_r \vec{E} + \vec{P}$$

Linear isotropic material

$$\vec{P} = \epsilon_r \chi_e \vec{E}$$

$$\vec{D} = \epsilon_r (1 + \chi_e) \vec{E} = \epsilon \vec{E}$$

$$\therefore \vec{P} = (\epsilon - \epsilon_0) \vec{E}$$

Static

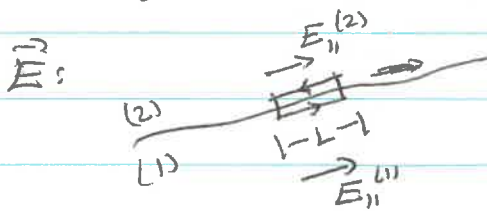
Boundary Conditions and Boundary Value Problems, with Dielectrics

It's easiest to ~~begin~~ begin with

$$\vec{\nabla} \cdot \vec{D} = \rho_{ext} \quad (1)$$

$$\vec{\nabla} \times \vec{E} = 0 \quad (2)$$

Now we ~~begin~~ consider a boundary between two dielectric material, which possibly contains a free surface charge density σ

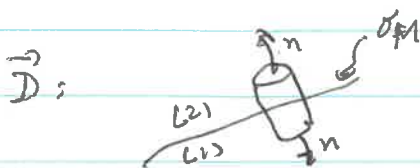


use $\vec{\nabla} \times \vec{E} = 0$

consider a ~~loop~~ line \int as shown:

$$\int \vec{E} \cdot d\vec{l} = 0 = L \cdot (E_{||}^1 - E_{||}^2)$$

$\therefore E_{||}^1 = E_{||}^2$: tangential component of \vec{E} is continuous



consider a ~~pillbox~~ Gaussian pillbox, use Gauss' law for \vec{D}

$$(\vec{D}_2 - \vec{D}_1) \cdot \vec{n} = \sigma_{ext}$$

(discontinuity in normal component of \vec{D} given by ~~free~~ ^{macro} surface charge density.)

$$\vec{D} = \epsilon \vec{E}$$

Also: note from (2)

$$\vec{E} = -\vec{\nabla} \Phi \quad \text{so} \quad \vec{\nabla} \cdot [\epsilon \vec{\nabla} \Phi] = -\rho_{ext}$$

and if ϵ is piecewise constant, $\epsilon \nabla^2 \Phi = -\rho_{ext}$

~~These 2 b.c.~~ These 2 b.c. ~~are~~ + ~~the~~ Poisson's equation get us through most boundary value problems.

$$\nabla \cdot (\epsilon_0 \vec{E} + \vec{P} + \dots) = \rho_{\text{macro}}$$

$$\epsilon_0 \nabla \cdot \vec{E} = \rho_{\text{macro}} - \nabla \cdot \vec{P}$$

suppose $\rho_{\text{micro}} = 0$, $\vec{E} = -\nabla \Phi$

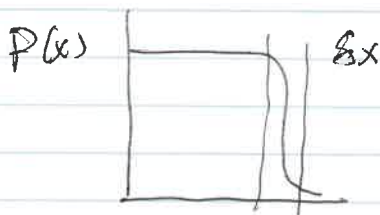
$$\nabla^2 \Phi = \frac{\nabla \cdot \vec{P}}{\epsilon_0}$$

then (formally)

$$\Phi(x) = - \int_{\text{vol of dielectric}} d^3x' \frac{1}{4\pi\epsilon_0} \frac{\nabla' \cdot \vec{P}(x')}{|x-x'|} \quad (*)$$

$\Rightarrow -\nabla \cdot \vec{P} =$ volume charge density in analogy with ρ

Also, this assumes $\nabla \cdot \vec{P}$ is smooth. It's sometimes convenient to treat \vec{P} as if it were a constant but this is continuous - almost



$$\nabla \cdot \vec{P} \sim \frac{1}{\delta} \vec{P}(x) \cdot \hat{n}$$

$$\int d^3x \nabla \cdot \vec{P} = \int \delta d^3x \frac{1}{\delta} \vec{P} \cdot \hat{n}$$

i.e. $\sigma_{\text{pol}} = \vec{P} \cdot \hat{n}$

$$\Phi = \frac{1}{4\pi\epsilon_0} \int d^3x' \frac{\sigma_{\text{pol}}(x')}{|x-x'|}$$

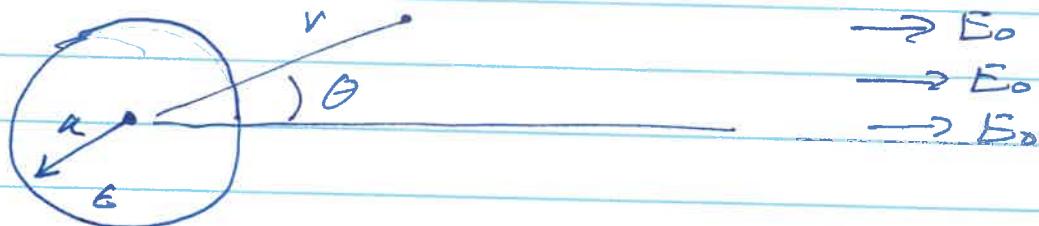
Be careful not to over count with (*)

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

$$\vec{E} = -\nabla \Phi$$

BCD-2

Electrostatics with linear dielectrics is only a small variation on what we've done before. Just solve for Φ within & outside the dielectric, then match b.c.s, continuity in normal D , tangential E . A classical problem: Dielectric sphere in external E -field E_0



$$\Phi_{\text{out}}(r, \theta) = \sum_{l=0}^{\infty} \left(B r^l + \frac{C_l}{r^{l+1}} \right) P_l(\cos \theta)$$

$$\longrightarrow \left(-E_0 r + \frac{C_1}{r^2} \right) \cos \theta$$

(what happened to the other l 's?) and

$$\Phi_{\text{in}} = A r \cos \theta \quad (\vec{E} = -\nabla \Phi)$$

continuity in Φ : $-E_0 a + \frac{C_1}{a^2} = A a$ (1)

$D \cdot n$ continuous $-\epsilon \frac{\partial \Phi_{\text{in}}}{\partial r} \Big|_{r=a} = -\epsilon_0 \frac{\partial \Phi_{\text{out}}}{\partial r} \Big|_{r=a}$

$$-\epsilon A = \epsilon_0 \left[E_0 + \frac{2C_1}{a^3} \right] \quad (2)$$

or $\left[A = -E_0 + \frac{C_1}{a^3} \right] \cdot 2\epsilon_0$ from (1)

$$(2\epsilon_0 + \epsilon) A = -3\epsilon_0 E_0$$

$$2\epsilon_0 A = -2\epsilon_0 E_0 + \frac{2C_1}{a^3}$$

$$A = \frac{-3\epsilon_0}{2\epsilon_0 + \epsilon} E_0 \quad C_1 = a^3 E_0 \cdot \left(\frac{\epsilon - \epsilon_0}{\epsilon + 2\epsilon_0} \right)$$

$$\Phi_{in} = \frac{-3\epsilon_0}{\epsilon + 2\epsilon_0} E_0 r \cos\theta$$

$$\Phi_{out} = -E_0 \left[r - \left(\frac{\epsilon - \epsilon_0}{\epsilon + 2\epsilon_0} \right) \frac{a^3}{r^2} \right] \cos\theta$$

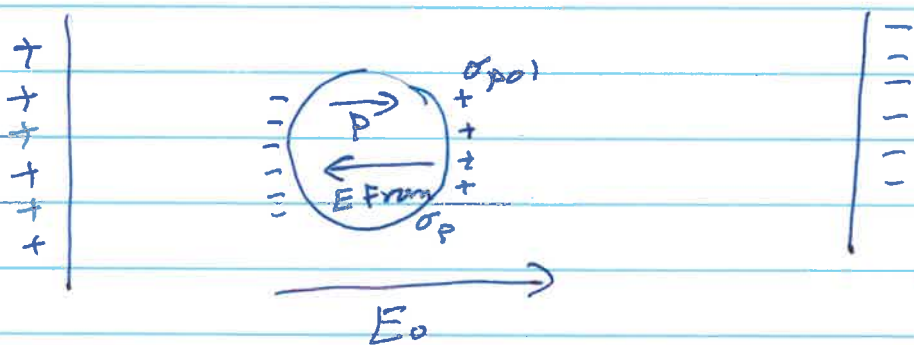
Notice inside $\vec{E}_{in} = \hat{z} \left(\frac{3}{2 + \frac{\epsilon}{\epsilon_0}} \right) E_0$

constant and smaller than E_0

$$\text{Polarization: } \vec{P} = (\epsilon - \epsilon_0) \vec{E}_{in} = 3\epsilon_0 \left[\frac{\frac{\epsilon}{\epsilon_0} - 1}{\frac{\epsilon}{\epsilon_0} + 2} \right] \vec{E}_0$$

$$\sigma_{pol} = \vec{P} \cdot \hat{n} = \vec{P} \cdot \hat{r} = 3\epsilon_0 \left(\frac{\frac{\epsilon}{\epsilon_0} - 1}{\frac{\epsilon}{\epsilon_0} + 2} \right) E_0 \cos\theta$$

σ_{pol} generates an E-field which is oppositely oriented to E_0 , reducing the internal E



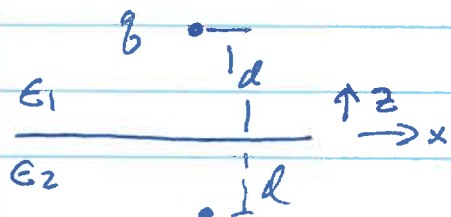
$$\vec{E}_{in} = \vec{E}_{out} \left[\frac{3}{2 + \epsilon/\epsilon_0} \right] = -\nabla\Phi$$

$$\frac{3\epsilon_0 \left(1 + \frac{\epsilon}{\epsilon_0} - 1 \right)}{\frac{\epsilon}{\epsilon_0} + 2}$$

$$\vec{D}_{in} = \epsilon_0 \vec{E}_{out} \left[\frac{3\epsilon/\epsilon_0}{2 + \epsilon/\epsilon_0} \right] = \epsilon \vec{E}_{in}$$

$$\vec{P}_{in} = \vec{D}_{in} - \epsilon_0 \vec{E}_{in} = \epsilon_0 \vec{E}_{out} \times \frac{3 \left(\frac{\epsilon}{\epsilon_0} - 1 \right)}{\left(\frac{\epsilon}{\epsilon_0} + 2 \right)}$$

Images and dielectrics: point charge in medium 1, above medium 2 - both linear dielectrics



$$\epsilon_1 \nabla \cdot \mathbf{E} = \rho \quad z > 0$$

$$\epsilon_2 \nabla \cdot \mathbf{E} = 0 \quad z < 0$$

$$\nabla \times \mathbf{E} = 0$$

and D.n continuous $\epsilon_1 E_z^{(1)} = \epsilon_2 E_z^{(2)}$ at $z=0$
 $E_{||}^{(1)} = E_{||}^{(2)}$ at $z=0$

We guess that to solve for the potential in (1), an image at $z = -d$ will do the trick, but maybe the charge is different

$$\Phi^{(1)} = \frac{1}{4\pi\epsilon_1} \left\{ \frac{q}{R_1} + \frac{q'}{R_2} \right\}$$

$$R_1^2 = \underbrace{x^2 + y^2}_{\rho^2} + (d-z)^2 \quad R_2^2 = \rho^2 + (z+d)^2$$

There are no charges in region (2), but there will be a surface polarization charge on the interface. Let's guess that Φ in region (2) can be given by a point charge q'' at the location of the real charge

$$\Phi^{(2)} = \frac{1}{4\pi\epsilon_2} \frac{q''}{R_1}$$

Need 2 identities

$$\text{Derivatives: } \frac{\partial}{\partial z} \frac{1}{R_1} \Big|_{z=0} = -\frac{\partial}{\partial z} \frac{1}{R_2} \Big|_{z=0} = \frac{d}{[\rho^2 + d^2]^{\frac{3}{2}}}$$

(b) $\frac{\partial}{\partial e} \frac{1}{R_1} \Big|_{z=0} = \frac{\partial}{\partial e} \frac{1}{R_2} \Big|_{z=0} = \frac{-e}{[e^2 + d^2]^{3/2}}$

so D.n: $\epsilon_1 E_2^{(1)} = \epsilon_2 E_2^{(2)}$

$$\frac{\epsilon_1}{4\pi\epsilon_1} \left\{ \frac{\partial}{\partial z} \left[\frac{q}{R_1} + \frac{q'}{R_2} \right] \right\} = \frac{\epsilon_2}{4\pi\epsilon_2} \frac{\partial}{\partial z} \frac{q''}{R_1}$$

$\boxed{q = q' = q''}$ from (a)

$$E_x^{(1)} = E_x^{(2)} \quad \frac{1}{4\pi\epsilon_1} \frac{\partial}{\partial e} \left[\frac{q}{R_1} + \frac{q'}{R_2} \right] = \frac{1}{4\pi\epsilon_2} \frac{\partial}{\partial e} \frac{q''}{R_1}$$

$\boxed{\frac{1}{\epsilon_1} [q + q'] = \frac{1}{\epsilon_2} q''}$ from b

$$\frac{q - q'}{\epsilon_1} = \frac{q''}{\epsilon_1}$$

$$\frac{q}{\epsilon_1} + \frac{q'}{\epsilon_1} = \frac{q''}{\epsilon_2}$$

$$q'' = \frac{2\epsilon_2}{\epsilon_1 + \epsilon_2} q$$

o $q'' =$ "below" image

$$q' = - \left[\frac{\epsilon_2 - \epsilon_1}{\epsilon_2 + \epsilon_1} \right] q$$

note if $\epsilon_1 = \epsilon_2$, $q' = 0$, $q'' = q$

o $q' =$ "above" image

The ~~volume~~ volume polarization charge density is

$\rho_{pol} = -\vec{\nabla} \cdot \vec{P}$ - it is zero (except at the point charge q) - but, at the interface, there's a surface polarization charge $\sigma_p = (\vec{P}_1 - \vec{P}_2) \cdot \hat{n}$...

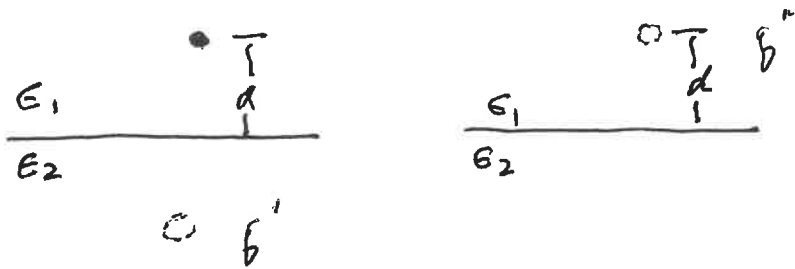
$$D = \epsilon E = \epsilon_0 E + P \quad \text{so} \quad \vec{P} = (\epsilon - \epsilon_0) \vec{E} = \left(\frac{\epsilon - \epsilon_0}{\epsilon} \right) \vec{D}$$

~~$\sigma_p = \left(\frac{\epsilon - \epsilon_0}{\epsilon} \right) D$~~

Recap:
$$\Phi^{(1)} = \frac{1}{4\pi\epsilon_1} \left[\frac{q}{R_1} + \frac{q'}{R_2} \right]$$

$$\Phi^{(2)} = \frac{1}{4\pi\epsilon_2} \left[\frac{q''}{R_1} \right]$$

$$R_1^2 = c^2 + (z-d)^2 \quad R_2^2 = c^2 + (z+d)^2$$



$\vec{D} \cdot \hat{n}$ continuous: $\epsilon_1 E_z^{(1)} = \epsilon_2 E_z^{(2)}$

$$-\epsilon_1 \left. \frac{\partial \Phi^{(1)}}{\partial z} \right|_{z=0} = -\epsilon_2 \left. \frac{\partial \Phi^{(2)}}{\partial z} \right|_{z=0}$$

\vec{E}_{tan} continuous

$$\left. \frac{\partial \Phi^{(1)}}{\partial c} \right|_{z=0} = \left. \frac{\partial \Phi^{(2)}}{\partial c} \right|_{z=0}$$

2 eqns, 2 unknowns q' & q''

(algebra ...)

$$q'' = \frac{2\epsilon_2}{\epsilon_1 + \epsilon_2} q \quad q' = - \left(\frac{\epsilon_2 - \epsilon_1}{\epsilon_1 + \epsilon_2} \right) q$$

check: if $\epsilon_1 = \epsilon_2$ $q' = 0$, $q'' = q$ (no interface)

There could be a volume polarization charge density:

$$\rho_{pol} = -\vec{\nabla} \cdot \vec{P} \quad \text{but} \quad \vec{P} = (\epsilon - \epsilon_0) \vec{D} \quad \text{and} \quad \vec{\nabla} \cdot \vec{D} = 0$$

except at $z=0$. There is a surface charge density

$$\sigma_p = (\epsilon_1 - \epsilon_2) \cdot n \quad \text{due to discontinuity in } \epsilon.$$

Electrostatic Energy

If a point charge q_i is brought from infinity to a point x_i , against a potential Φ (which vanishes at infinity) the work done on it is

$$W_i = q_i \Phi(x_i)$$

Suppose Φ comes from $(n-1)$ point charges:

$$\Phi(x) = \frac{1}{4\pi\epsilon_0} \sum_{j=1}^{n-1} \frac{q_j}{|x-x_j|}$$

The total work done to assemble all the charges is

$$W = \frac{1}{4\pi\epsilon_0} \sum_{i=1}^n \sum_{j=1}^{i-1} \frac{q_i q_j}{|x_i - x_j|}$$

(they all fight each other.) This is a sum over pairs: sum over everybody, with a $1/2$:

$$W = \frac{1}{2} \sum_i \sum_{\substack{j \\ \{i \neq j\}}} \frac{q_i q_j}{|x_i - x_j|} \frac{1}{4\pi\epsilon_0}$$

pass to continuum be careful about $i \neq j$ (neglect self energy)

$$W = \frac{1}{4\pi\epsilon_0} \frac{1}{2} \int \frac{\rho(x) \rho(x')}{|x-x'|} d^3x d^3x'$$

$$\text{or } \frac{1}{2} \int \rho(x) \Phi(x) d^3x \quad \text{---} \rightarrow \rho = -\epsilon_0 \nabla^2 \Phi$$

$$\text{or } \frac{-\epsilon_0}{2} \int \Phi(x) \nabla^2 \Phi(x) d^3x$$

$$\text{or } \frac{1}{2} \frac{\epsilon_0}{2} \int |\nabla \Phi|^2 d^3x = \frac{\epsilon_0}{2} \int E^2 d^3x$$

energy density in electrostatic field $w = \frac{\epsilon_0}{2} E^2$

all very standard...

Energy and Dielectrics

Energy used to assemble charges all brought in from infinity is

$$W = \frac{1}{2} \int \rho(x) \Phi(x) d^3x$$

counts all the charge ~~at~~ (microscopic, macroscopic) and all the fields.

Typically we want to bring together pre-charges and pre-assembled dielectrics, and get a bill for them, not for the cost of the dielectric.

Start over

Make a change $\delta\rho$ in macroscopic charge density

$$\delta W = \int \delta\rho(x) \Phi(x) d^3x \quad \text{***} \quad (\text{keep this})$$

Φ is potential due to all charge already present,

$$\vec{E} = -\vec{\nabla}\Phi$$

but $\vec{\nabla} \cdot \vec{D} = \rho$, $\delta\rho = \vec{\nabla} \cdot \delta\vec{D}$

$$\delta W = \int [\vec{\nabla} \cdot (\delta\vec{D})] \Phi \xrightarrow{\text{parts}} \int \delta\vec{D} \cdot \vec{E} d^3x$$

2 - signs ($\vec{E} = -\vec{\nabla}\Phi$)

$$W = \int d^3x \int_0^D \delta\vec{D} \cdot \vec{E}$$

Formal total electrostatic energy, gotten by bringing D from 0 to its final value.

linear medium only $\vec{E} \cdot \delta\vec{D} = \frac{1}{2} \delta(\vec{E} \cdot \vec{D})$

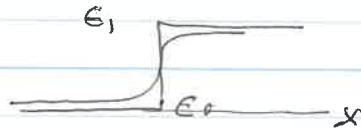
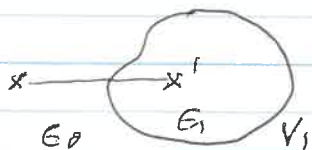
$$W = \frac{1}{2} \int \vec{E} \cdot \vec{D} d^3x$$

A different problem: What is the change in energy when a linear dielectric is placed in an E field, where sources (charges) are fixed?

Initially $\epsilon_0(x), E_0(x), \vec{E}_0(x), \vec{D}_0(x) = \epsilon_0 \vec{E}_0(x)$

$$W_0 = \frac{1}{2} \int \vec{E}_0 \cdot \vec{D}_0 d^3x$$

Now drop in a dielectric with volume V_1



$$\vec{E}_0 \rightarrow \vec{E}, \vec{D} = \epsilon \vec{E} \quad W_1 = \frac{1}{2} \int \vec{E} \cdot \vec{D} d^3x$$

$$W_1 - W_0 = \Delta W = \frac{1}{2} \int d^3x (\vec{E} \cdot \vec{D} - \vec{E}_0 \cdot \vec{D}_0)$$

$$= \frac{1}{2} \int \underbrace{(\vec{E} \cdot \vec{D}_0 - \vec{D} \cdot \vec{E}_0)}_{\text{zero outside } V_1} d^3x + \frac{1}{2} \int \underbrace{(\vec{E} + \vec{E}_0) \cdot (\vec{D} - \vec{D}_0)}_{\text{will be zero}} d^3x$$

zero outside V_1

$\vec{D} = \epsilon_0 \vec{E}$ outside, $\vec{D} = \epsilon_1 \vec{E}$ inside

$\vec{D}_0 = \epsilon_0 \vec{E}_0$ everywhere

so outside $\vec{E} \cdot \epsilon_0 \vec{E}_0 - \epsilon_0 \vec{E} \cdot \vec{E}$

will be zero

drop for now

$\equiv I_2$

$$\Delta W = \frac{1}{2} \int_{V_1} [\epsilon_0 - \epsilon_1] \vec{E} \cdot \vec{E}_0 d^3x \quad (\text{circled})$$

in free space

$$\vec{D} = \epsilon_0 \vec{E} + \vec{P} = \epsilon_1 \vec{E}$$

$$(\epsilon_1 - \epsilon_0) \vec{E} = \vec{P}$$

$$W = - \frac{1}{2} \int_{V_1} \vec{P} \cdot \vec{E}_0 d^3x$$

energy density of ^{linear} dielectric placed in field E_0 whose sources are fixed

$$w = -\frac{1}{2} \vec{P} \cdot \vec{E}_0$$

like $W = \dots - \vec{p} \cdot \vec{E}_0$ as for dipole \vec{p} with $\frac{1}{2}$ because medium is polarizable ($\frac{1}{2} \vec{D} \cdot \vec{E}$)

o.o of $\epsilon_1 > \epsilon_0$, dielectric sucked into region of increasing field E_0

Again, this is for fixed charge, not fixed voltage (no batteries)

The term which is zero!

$$\nabla \times (\vec{E} + \vec{E}_0) = -\nabla \tilde{\Phi}$$

$$I_2 = -\frac{1}{2} \int \vec{\nabla} \tilde{\Phi} \cdot (\vec{D} - \vec{D}_0) d^3x$$

$$= \frac{1}{2} \int \tilde{\Phi} \vec{\nabla} \cdot (\vec{D} - \vec{D}_0) d^3x$$

but \vec{D} is unaffected by the dielectric;

$$\nabla \cdot \vec{D} = \nabla \cdot \vec{D}_0$$

$$I_2 = 0$$

Fixed charge: $W = \text{energy}$. If you shift a coordinate $\vec{x} \rightarrow \vec{x} + \delta\vec{x}$, the "principle of virtual work" says

$$\delta W = - \vec{F} \cdot \hat{n} \delta x$$

$\therefore \vec{F} = \text{force acting on dielectric} = - \nabla W$ fixed

Case of fixed Voltage is different. As ~~Φ is varied~~, as dielectric moves, wires carry charge in & out of battery to maintain potential. Energy is supplied by source.

Start with

$$W = \frac{1}{2} \int \epsilon \Phi^2 d^3x \quad \text{count all charges, as assembled.}$$

Change $\epsilon \rightarrow \epsilon + \delta\epsilon$, $\Phi \rightarrow \Phi + \delta\Phi$, imagine independent.

$$\delta W = \frac{1}{2} \int d^3x [\epsilon \delta\Phi + \Phi \delta\epsilon] \quad (*)$$

point to ~~***~~ on ED-1 If dielectric properties were changed this would be $\delta W = \int \delta\epsilon \cdot \Phi^2 d^3x$, so the 2 terms in (*) would be equal. But if dielectric properties change, $\epsilon(x) \rightarrow \epsilon(x) + \delta\epsilon(x)$, polarization charge changes, $\epsilon = \epsilon_{free} + \epsilon_{pol}$, we only want to deal with ϵ_{free} .

How to proceed:

1) Disconnect the battery, move the dielectric. Free Charges are fixed. We did this already

$$\delta W_1 = \frac{1}{2} \int \epsilon \delta\Phi_1 d^3x = - \frac{1}{2} \int [\epsilon_1 - \epsilon_0] \vec{E} \cdot \vec{E}_0 d^3x$$

in this process $\delta\Phi_1 \neq 0$ - potentials changed

2) Reconnect the battery. $\delta\Phi =$
 $\delta\Phi_2 = -\delta\Phi_1$ to restore
 all the potentials - but there is also a $\delta\epsilon$

$$\delta W_2 = \frac{1}{2} \int [\epsilon \delta\Phi_2 + \Phi \delta\epsilon] d^3x$$

but - dielectric properties are not changed - this is

$$\delta W = \int \delta\epsilon \cdot \Phi$$

$$\Rightarrow \epsilon \delta\Phi = \Phi \delta\epsilon$$

And $\delta\Phi = \delta\Phi_2 = -\delta\Phi_1$

$$\delta W_2 = -\frac{1}{2} \delta W_1$$

$$\Rightarrow \delta W = (1-2) \frac{1}{2} \int \epsilon \delta\Phi_1 d^3x' = -\delta W_1$$

the sign flips!

$$F = + \nabla W \Big|_{\text{fixed } V}$$

Horrible business - but "physical" description

is in Griffiths - see p-188

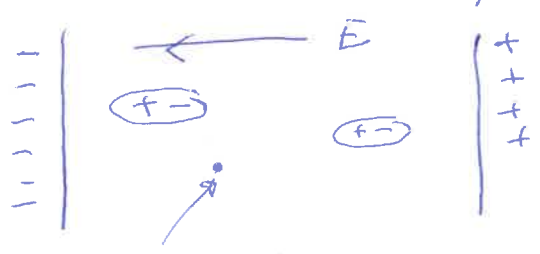
More dielectric w/ $\epsilon > 1$ into region of greater \vec{E}
 at fixed voltage, energy increases.

Macroscopic and microscopic quantities

$$\vec{P} = \epsilon_0 \chi_e \vec{E}$$

\vec{P} = polarization
 \vec{p} = dipole moment

χ_e is a macroscopic quantity. How is it related to microscopic properties of the material? Interestingly, somewhat complicated story of how might - see Zangwill, Ch 6. What follows is a story based on (semi) classical modelling of material as little dipoles



\vec{E} Field at a point = $\vec{E}_{tot} = \text{macro field} + \text{local variation}$

$$\vec{E}_{tot} = \vec{E} + \vec{E}_i$$

$$\vec{E}_i = \vec{E}_{near} (\text{from a real calculation}) - \vec{E}_p$$

(E_p = field from smooth background of polarization - this is a macroscopic quantity, already included in \vec{E})

$$- \vec{E}_{tot} = \vec{E} + \vec{E}_{near} - \vec{E}_p$$

Now recall the story about the dipole moment (S-fn in dipole field)

$$\int_{r < R} \vec{E}(x) d^3x = -\frac{\vec{P}}{3\epsilon_0}$$

$$\vec{P} = \text{dipole moment / unit vol} \Rightarrow \vec{P} = \frac{4\pi R^3}{3} \vec{P}$$

$$\vec{E}_p = \text{average field} = \frac{1}{\left[\frac{4\pi R^3}{3} \right]} \int_{r < R} E d^3x = -\frac{\vec{P}}{3\epsilon_0}$$

$$\Rightarrow \vec{E}_i = \frac{\vec{P}}{3\epsilon_0} + \vec{E}_{near} \quad \text{is correction to macro } \vec{E} \text{ in the dielectric}$$

Size? $\frac{E}{E_0} = 1 + \chi_e$

$$\chi_e = \frac{N \delta_m}{1 - \frac{1}{3} N \delta_m} \sim N \delta_m \text{ if } N \delta_m \text{ is small}$$

Back of the envelope in CGS --

gas: $N \sim \frac{6 \cdot 10^{23}}{22.4 \times 10^3 \text{ cm}^3} \times \left(\frac{1 \text{ cm}}{10^8 \text{ \AA}} \right)^3 \sim 10^{-5} \frac{1}{\text{ \AA}^3}$

$$N \frac{e^2}{m a^2} = N \frac{e^2}{\hbar c} \frac{1}{m c^2} \frac{1}{(\hbar \omega)^2} (h c)^3$$

$$= \frac{10^{-5}}{\text{ \AA}^3} \times \frac{1}{137} \times \frac{1}{5 \times 10^5 \text{ eV}} \times \frac{1}{(\text{eV})^2} \times (2000 \text{ eV} \cdot \text{ \AA})^3$$

$$\frac{e^2}{E_0} = 4\pi \left(\frac{e^2}{4\pi\epsilon_0} \right) = \frac{1}{137} \times 10^{-5-5+9} \sim 10^{-3}$$

$$\frac{E}{E_0} = 1.005 \text{ for air}$$

$N \bar{v} \sim 10^5$ times greater for solids -

$$\frac{E}{E_0} \sim 1 + 1$$

~~$E_{tot} = E + E_{near}$~~ MM-2
 \vec{E}_{near} involves a better calculation. Sometimes

(believe it or not) it is zero! (amorphous materials, gases, Lorentz model - see Jackson). If that is so

we $\vec{P} = N \langle \vec{P}_{mol} \rangle$ $N = \text{density}$
 $\langle \vec{P}_{mol} \rangle \equiv \text{average molecular dipole moment.}$

Define molecular polarizability χ_m via

$$\langle \vec{P}_{mol} \rangle = \epsilon_0 \chi_m \vec{E}_{tot}$$

$$\begin{aligned} \vec{P} &= \epsilon_0 N \chi_m \vec{E}_{tot} = \epsilon_0 N \chi_m (\vec{E} + \vec{E}_L) \\ &= \epsilon_0 N \chi_m \left(\vec{E} + \frac{1}{3\epsilon_0} \vec{P} \right) \end{aligned}$$

solve for \vec{P} :

$$\vec{P} = \epsilon_0 \left[\frac{N \chi_m}{1 - \frac{1}{3} N \chi_m} \right] \vec{E}$$

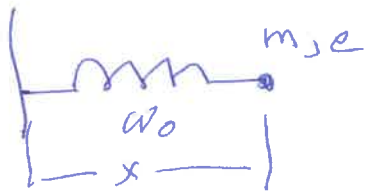
$$\chi_E = \frac{N \chi_m}{1 - \frac{1}{3} N \chi_m} ; \quad \frac{\epsilon}{\epsilon_0} = 1 + \chi_E$$

Inverse is called the Clausius Mossotti formula (1850, 1870)

$$\chi_m = \frac{3}{N} \left[\frac{\frac{\epsilon}{\epsilon_0} - 1}{\frac{\epsilon}{\epsilon_0} + 2} \right] \quad [] \text{ like } \rho_p$$

Kind of odd - micro χ_m in terms of macro ϵ -
 this is from the ~~very~~ early days of atoms - but
 they knew macro facts, trying to get at micro probs

An oscillator model for χ_m ($\vec{P} = \epsilon_0 \chi_m \vec{E}$) MM-3
 electron on a spring!



$$e \vec{E} = m \omega_0^2 \vec{x}$$

$$\vec{P} = e \vec{x} = \frac{e^2}{m \omega_0^2} \vec{E}$$

$$\chi_m = \frac{1}{\epsilon_0} \frac{e^2}{m \omega_0^2} \longrightarrow \frac{1}{\epsilon_0} \sum_i \frac{e_i^2}{m_i (\omega_0^i)^2}$$

~~mkgs~~ mks: drop for cgs ($\frac{1}{\epsilon_0} \leftrightarrow 4\pi$)

Or - at finite temperature

$$H = \frac{p^2}{2m} + \frac{1}{2} m \omega_0^2 x^2 - e E z$$

$$\langle P_z = e \cdot z \rangle_{T=0} = \frac{\int d^3 p \int d^3 x e^{-H/kT} \cdot e z}{\int d^3 p \int d^3 x e^{-H/kT}}$$

$$= \frac{\int dz \cdot e z \cdot e^{-\frac{1}{2} m \omega_0^2 z^2 - e E z}}{\int dz e^{-\frac{1}{2} m \omega_0^2 z^2 - e E z}}$$

$$\langle \vec{P} \rangle = \frac{e^2}{m \omega_0^2} \vec{E} \text{ again}$$

We will use generalizations of this model
 again & again!