

P112A:Electromagnetic Theory

Electric Fields in Matter (Chapter 4)

Electrical properties of solid Materials

- Charges in conductors and insulators (or dielectrics)
 - Conductors: Substances with an unlimited supply of electrons which are free to move throughout the material (but not off it)
 - Dielectrics: all charges are attached to specific atoms or molecules. All they can do is slosh around a bit within the atom or molecule

Dielectrics

- While the motion is limited, the cumulative effects account for the observed changes in E-fields within materials
- There are two principle ways for external electric fields to distort the charge distribution of an atom in a dielectric material: **stretching** and **rotating**
 - While neutral atoms do not typically have a dipole moment, many molecules do.
 - However, most materials with dipole molecules are randomly oriented so no net dipole moment in the absence of an external E-field.

Inducing a dipole by stretching

- If a neutral atom is placed in an external E field, the positive nucleus is pushed away, while the surrounding electron cloud is pulled in opposite direction.
- The displaced + and - charges set up an E -field in the direction opposite to the external field. The larger the displacement, the larger the induced E -field. Equilibrium is reached when the force from the external E is equal and opposite to the force from the induced E field.
- The displaced charge forms a dipole, that is described by a dipole moment, \mathbf{p} . For simple atoms, \mathbf{p} is same direction as E , and the magnitude of the effect is proportional to E : $\mathbf{p} = \alpha \mathbf{E}$, where α is called the atomic polarizability.

Example 4.1: Model of polarized atom

- Assume electrons surrounding an atom uniformly fill a sphere of radius a . In the presence of an external \mathbf{E} pointing to the right, the nucleus shifted is shifted right, and electron cloud to the left. The shift is small compared to “ a ” so the cloud remains spherical. The shift from the center of the electron cloud is called “ d ”. Net force on nuclear charge is 0, so the value of d is determined when \mathbf{E}_e from electron cloud acting on nuclear charge is the same as external \mathbf{E} .

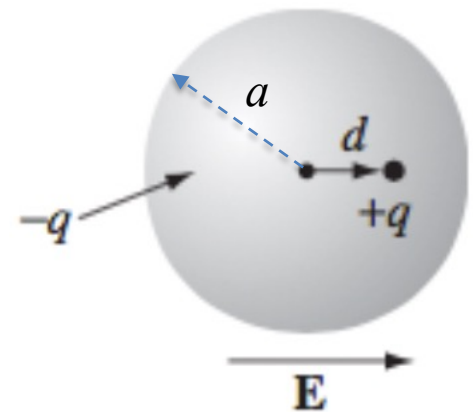
Use Gauss's Law to compute \mathbf{E}_e from uniform density of electric charge as a function of radius d :

$$Q_{enc} = \frac{4}{3}\pi d^3 \rho \quad |E_e| = \frac{1}{\epsilon_0} \cdot \frac{Q_{enc}}{4\pi d^2} \quad \rho = -q/(4/3\pi a^3)$$

$$|\vec{E}_e| = \frac{\rho d}{3\epsilon_0} = \frac{qd}{4\pi\epsilon_0 a^3}; \quad p = qd$$

$$E = |\vec{E}_e| = \frac{qd}{4\pi\epsilon_0 a^3} = \frac{p}{4\pi\epsilon_0 a^3}$$

$$p = (4\pi\epsilon_0 a^3) E = [3\epsilon_0 (\text{volume})] E$$



$p = \alpha E$, atomic polarizability $\alpha = 3\epsilon_0 \cdot \text{Volume}$

Model of polarized atom

- Model predicts magnitude of α is proportional to Volume of atom and ϵ_0 . Lets test it.
- Consider hydrogen: $r \sim 0.5 \times 10^{-10} \text{ m}$
- $\text{Vol} \sim 4(0.5 \times 10^{-10} \text{ m})^3 = 0.5 \times 10^{-30} \text{ m}^3$.
- $\alpha/\epsilon_0 = 3 * \text{Vol} = 1.5 \times 10^{-30} \text{ m}^3$ (predicted)
- $\alpha/\epsilon_0 = 4\pi(0.67) \times 10^{-30} = 8 \times 10^{-30} \text{ m}^3$ (measured)

Not too bad for 7D, 7E physics model

H	He	Li	Be	C	Ne	Na	Ar	K	Cs
0.667	0.205	24.3	5.60	1.67	0.396	24.1	1.64	43.4	59.4

TABLE 4.1 Atomic Polarizabilities ($\alpha/4\pi\epsilon_0$, in units of 10^{-30} m^3). *Data from: Handbook of Chemistry and Physics*, 91st ed. (Boca Raton: CRC Press, 2010).

Values, for the most part, increase with volume of atom

Is model reasonable? How much displacement are we talking about?

- Suppose hydrogen is placed between 2 metal plates, $1\text{mm}=10^{-3}\text{m}$ apart, connected to a 1 kV power supply. What is d , the separation between + nucleus and center of electron cloud?

$$|\mathbf{E}_{\text{ext}}| = V/d = 10^3/10^{-3} = 10^6 \text{ V/m}$$

$$|\mathbf{p}| = dq = \alpha |\mathbf{E}_{\text{ext}}|$$

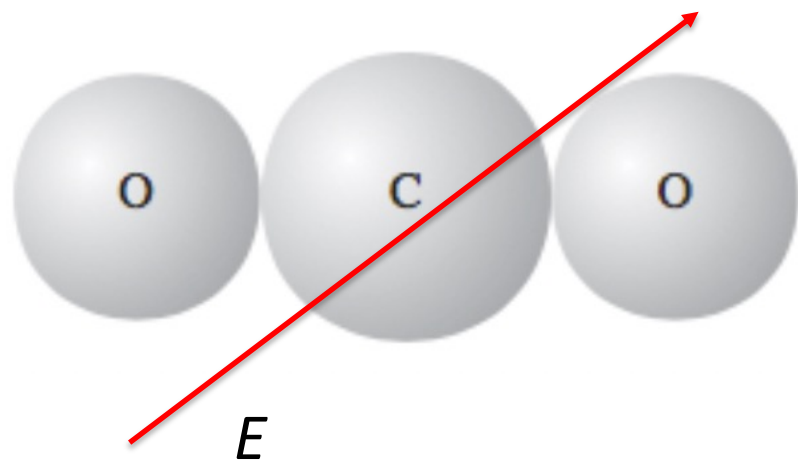
$$d = \alpha |\mathbf{E}_{\text{ext}}| / q = 4\pi\epsilon_0 (0.667 \times 10^{-30}) [10^6] / 1.6 \times 10^{-19} \text{C} = 12(2/3 \times 10^{-30})(9 \times 10^{-12}) [10^6] / 2 \times 10^{-19} = 36 \times 10^{-17} \text{ m}$$

$d = 0.4 \times 10^{-15} \text{ m}$, which is 10^{-5} of the atomic radius $\sim 10^{-10} \text{ m}$, so displacements are really tiny for most lab situations. Seems reasonable to assume the electron cloud does not change much \rightarrow our model is reasonable

Polarizability of Molecules

- For molecules, frequently, they polarize more readily in one direction than another.
- For example, CO₂, polarizability is larger along the axis than perpendicular to it :

$$\mathbf{p} = \alpha_{\perp} \mathbf{E}_{\perp} + \alpha_{\parallel} \mathbf{E}_{\parallel}.$$



The direction of \mathbf{p} may not be the same direction as the External E field

Asymmetric Molecules

- CO₂ is linear, and relatively symmetric. But most molecules are not. A more complicated geometry requires:

$$\left. \begin{aligned} p_x &= \alpha_{xx} E_x + \alpha_{xy} E_y + \alpha_{xz} E_z \\ p_y &= \alpha_{yx} E_x + \alpha_{yy} E_y + \alpha_{yz} E_z \\ p_z &= \alpha_{zx} E_x + \alpha_{zy} E_y + \alpha_{zz} E_z \end{aligned} \right\}$$

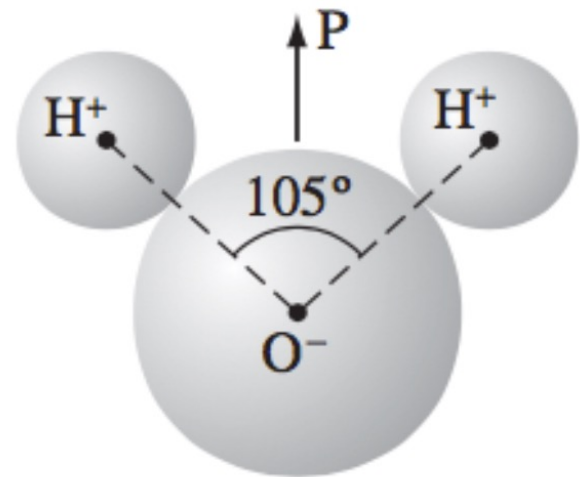
*Can be written in terms of a
polarizability matrix, or Tensor.
Here \mathbf{p} is induced by external \mathbf{E}*

*One needs to measure 9 numbers, but the numbers vary with choice of axis. If it is possible to predict direction of “**principle axis**”, then only need to measure 3 numbers because all off-diagonal terms vanish.*

Polar molecules

- A neutral atom has no dipole moment to start with, p was induced by the applied E field. However, some molecules have a built-in, permanent dipole moment.

Water molecule has $|p|=6 \times 10^{-30} \text{ C}\cdot\text{m}$, which is atypically large. In a liquid or gas, the orientations are random so the net dipole moment is 0. Not so obvious that solid should have no net dipole moment.



Torque on permanent dipole moment

- In a uniform field, the force on the positive charge, $F_+ = qE$, cancels the force on the negative side. However, there will be a torque, \mathbf{N} . (τ in 7D)

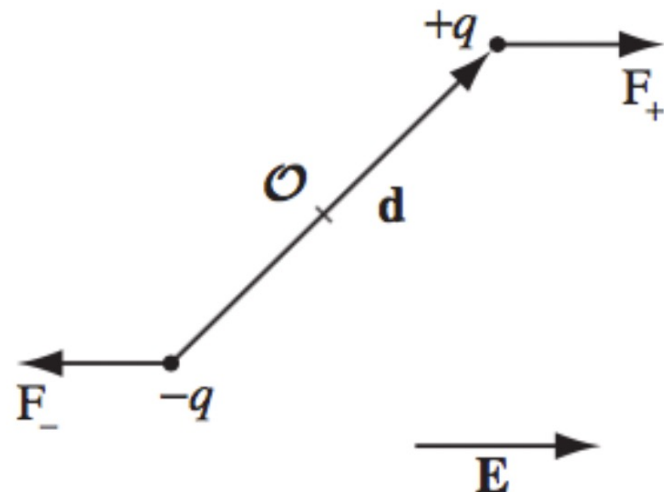
In uniform external E field, net force = 0

Torque:

$$\vec{N} = (\vec{r}_+ \times \vec{F}_+) + (\vec{r}_- \times \vec{F}_-)$$

$$\vec{N} = \left(\left[\frac{\vec{d}}{2} \right] \times q\vec{E} \right) + \left(\left[\frac{-\vec{d}}{2} \right] \times [-q\vec{E}] \right)$$

$$\vec{N} = q\vec{d} \times \vec{E} = \vec{p} \times \vec{E}$$



Non-uniform external E Field

- In non-uniform external \vec{E} , the forces do not necessarily cancel. Of course, \vec{E} must change rather abruptly over the distance of a molecule for this to be significant. So this is usually not considered in dielectrics.
- Nevertheless, the magnitude of the force is worth thinking about

$$\vec{F} = \vec{F}_+ - \vec{F}_- = q(\vec{E}_+ - \vec{E}_-) = q\Delta\vec{E} = q(\Delta E_x \hat{x} + \Delta E_y \hat{y} + \Delta E_z \hat{z})$$

$$(1.34) \quad dT = \left(\frac{\partial T}{\partial x}\right)dx + \left(\frac{\partial T}{\partial y}\right)dy + \left(\frac{\partial T}{\partial z}\right)dz$$

$$(1.35) \quad dT = \left[\left(\frac{\partial T}{\partial x}\right)\hat{x} + \left(\frac{\partial T}{\partial y}\right)\hat{y} + \left(\frac{\partial T}{\partial z}\right)\hat{z} \right] \cdot (dx\hat{x} + dy\hat{y} + dz\hat{z})$$

$$dT = \nabla T \cdot d\vec{\ell} \Rightarrow \Delta T = \nabla T \cdot \Delta\vec{\ell} \quad \text{Change from differential distance to finite difference}$$

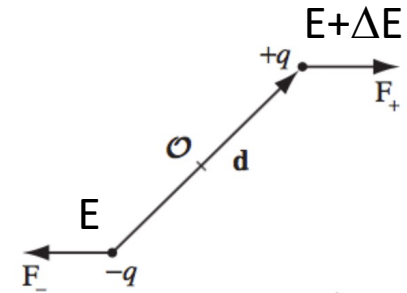
$$\Delta E_x = (\nabla E_x) \cdot \vec{d} = \vec{d} \cdot (\nabla E_x) \quad \text{Dot products commute}$$

$$\Delta E_y = \vec{d} \cdot (\nabla E_y); \quad \Delta E_z = \vec{d} \cdot (\nabla E_z)$$

$$\vec{F} = q(\Delta E_x \hat{x} + \Delta E_y \hat{y} + \Delta E_z \hat{z}) = q \left([\vec{d} \cdot (\nabla E_x)] \hat{x} + [\vec{d} \cdot (\nabla E_y)] \hat{y} + [\vec{d} \cdot (\nabla E_z)] \hat{z} \right)$$

$$\vec{F} = q(\vec{d} \cdot \nabla) \vec{E} \quad \text{Gives same thing as previous eqn, check with Cartesian coordinates}$$

$$\vec{F} = (\vec{p} \cdot \nabla) \vec{E}$$



Dipole moment per volume, \mathbf{P}

- To summarize. If dielectric material placed in external \mathbf{E} field.
 - Neutral atoms: Induce dipoles, pointing in same direction as \mathbf{E} (stretching)
 - Polar molecules: experience a torque, tending to line up dipole moment with \mathbf{E} (rotating)
- So a lot of dipoles in material point along direction of field, so material becomes polarized.
- Dipole moment per unit volume, \mathbf{P} , is called the polarization.
- Lets no longer worry about what causes \mathbf{P} (rotating, stretching or some combination), but lets examine the \mathbf{E} created by a chunk of polarized material.
- Eventually we will want to combine the \mathbf{E} created by \mathbf{P} AND the original \mathbf{E} that created \mathbf{P} in the first place.

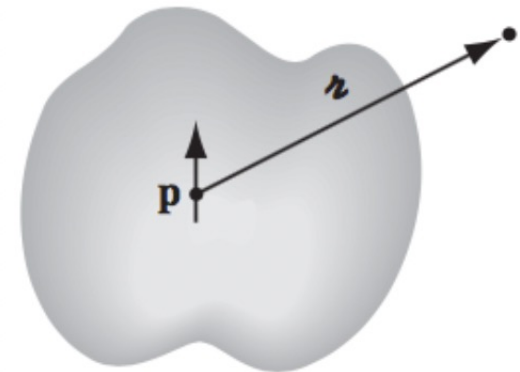
Field of a Polarized Object

- Suppose we have a piece of polarized material with polarization \mathbf{P} . What is the field produced by this object?
- For a single dipole chunk, $d\mathbf{p}$, within a material characterized by \mathbf{P} , the potential is

$$dV(r) = \frac{1}{4\pi\epsilon_0} \frac{(d\vec{p}) \cdot \hat{R}}{R^2}$$

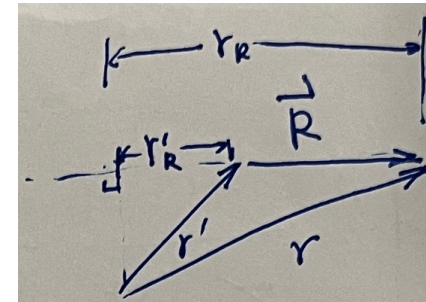
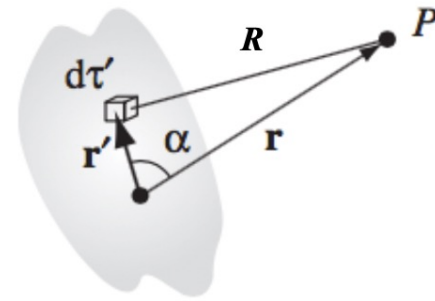
$$\text{where } d\vec{p} = \vec{P} d\tau$$

$$V(r) = \frac{1}{4\pi\epsilon_0} \int \frac{(d\vec{p}) \cdot \hat{R}}{R^2} = \frac{1}{4\pi\epsilon_0} \int \frac{(\vec{P}(\vec{r}') \cdot \hat{R})}{R^2} d\tau'$$



Field of a Polarized Object

$$\begin{aligned}\nabla'(1/R) &= \frac{\partial\left(\frac{1}{R}\right)}{\partial r'_R} \hat{R} = \frac{\partial\left(\frac{1}{|r-r'|}\right)}{\partial r'_R} \hat{R} = \frac{\partial\left(\frac{1}{r_R-r'_R}\right)}{\partial r'_R} \hat{R} \\ &= -\frac{1}{(r_R-r'_R)^2} \frac{\partial(r_R-r'_R)}{\partial r'_R} \hat{R} = \frac{\hat{R}}{R^2}\end{aligned}$$



$$V(r) = \frac{1}{4\pi\epsilon_0} \int \frac{(\vec{P}(\vec{r}') \cdot \hat{R})}{R^2} d\tau' = \frac{1}{4\pi\epsilon_0} \int (\vec{P}(\vec{r}') \cdot \nabla' \left(\frac{1}{R} \right) d\tau'$$

$$= \frac{1}{4\pi\epsilon_0} \left[\int_{Vol} \nabla' \cdot \left(\frac{\vec{P}}{R} \right) d\tau' - \int_{Vol} \left(\frac{1}{R} \right) (\nabla' \cdot \vec{P}) d\tau' \right]$$

$$= \frac{1}{4\pi\epsilon_0} \left[\int_{surf} \left(\frac{\vec{P}}{R} \right) \cdot d\mathbf{a}' - \int_{Vol} \left(\frac{1}{R} \right) (\nabla' \cdot \vec{P}) d\tau' \right]$$

Product Rule 5

$$(5) \nabla \cdot (f\mathbf{A}) = f(\nabla \cdot \mathbf{A}) + \mathbf{A} \cdot (\nabla f)$$

Divergence Thm

Surface and Volume **Bound Charges**

$$V(r) = \frac{1}{4\pi\epsilon_0} \left[\int_{surf} \left(\frac{\vec{P}}{R} \right) \cdot d\vec{a}' - \int_{Vol} \left(\frac{1}{R} \right) (\nabla' \cdot \vec{P}) d\tau' \right]$$

The first term looks like the potential from the **surface charge density**

$$\sigma_b \equiv \vec{P} \cdot \hat{n}$$

The second term looks like the potential of a **volume charge density**

$$\rho_b \equiv -\nabla \cdot \vec{P}$$

Instead of integrating contributions from all **dipoles** in dielectric, fields can be computed from these **bound charges**:

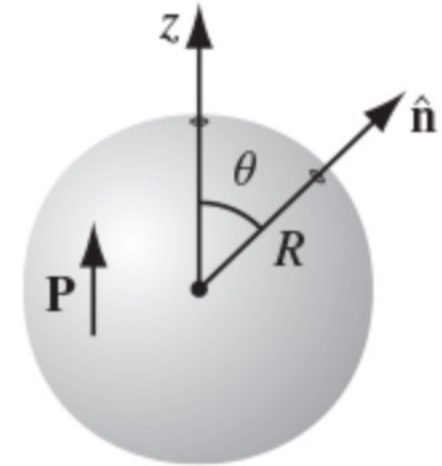
$$V(r) = \frac{1}{4\pi\epsilon_0} \left[\int_{surf} \left(\frac{\sigma_b}{R} \right) da' + \int_{Vol} \left(\frac{\rho_b}{R} \right) d\tau' \right]$$

Example 4.2

Find E field produced by uniformly polarized, \mathbf{P} , dielectric sphere of radius R . Choose z -axis to correspond to direction of polarization. Normal to spherical surface is radial direction, \mathbf{r} .

$$\sigma_b \equiv \vec{P} \cdot \hat{n} = P \hat{z} \cdot \hat{r} = P \cos \theta$$

$$\rho_b \equiv -\nabla \cdot \vec{P} = -\nabla \cdot P \hat{z} = 0$$



Bound charge only on surface, solved in Example 3.9 with variable separation method:

$$\sigma_0(\theta) = k \cos \theta = k P_1(\cos \theta), \quad \xrightarrow{k=P} \quad V(r, \theta) = \begin{cases} \frac{P}{3\epsilon_0} r \cos \theta & \text{for } r \leq R, \\ \frac{P}{3\epsilon_0} \frac{R^3}{r^2} \cos \theta & \text{for } r \geq R. \end{cases}$$

$$V(r, \theta) = \frac{k R^3}{3\epsilon_0} \frac{1}{r^2} \cos \theta \quad (r \geq R).$$

Example 4.2

$$V(r, \theta) = \frac{r P \cos \theta}{3\epsilon_0} \quad r < R$$

$$V_{\text{dip}}(r, \theta) = \frac{\hat{\mathbf{r}} \cdot \mathbf{p}}{4\pi\epsilon_0 r^2} = \frac{p \cos \theta}{4\pi\epsilon_0 r^2}$$

$$V(r, \theta) = \frac{P R^3 \cos \theta}{3\epsilon_0 r^2} \quad r > R$$

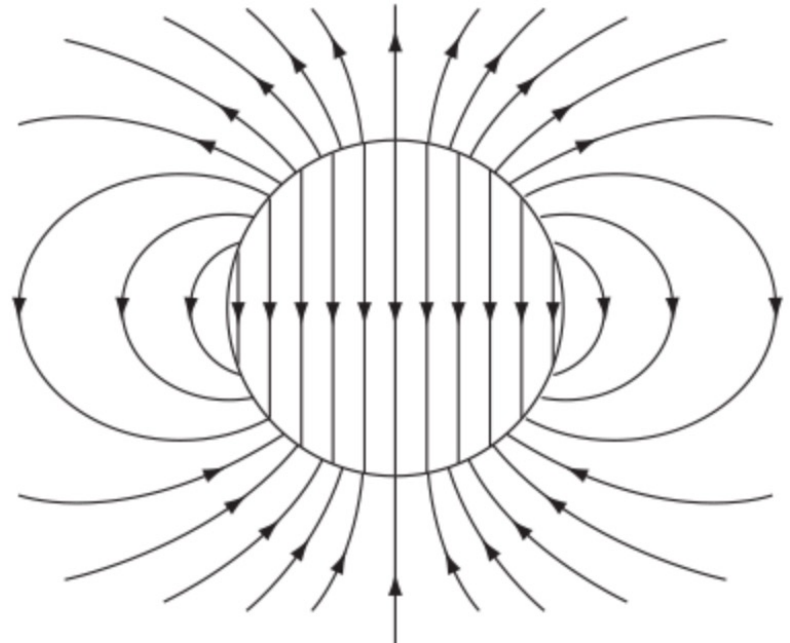
*Uniformly polarized sphere
produces the same potential outside as a
ideal dipole, $p = P \cdot (\text{volume of sphere})$*

$r \cos(\theta) = z$, when $r \leq R$:

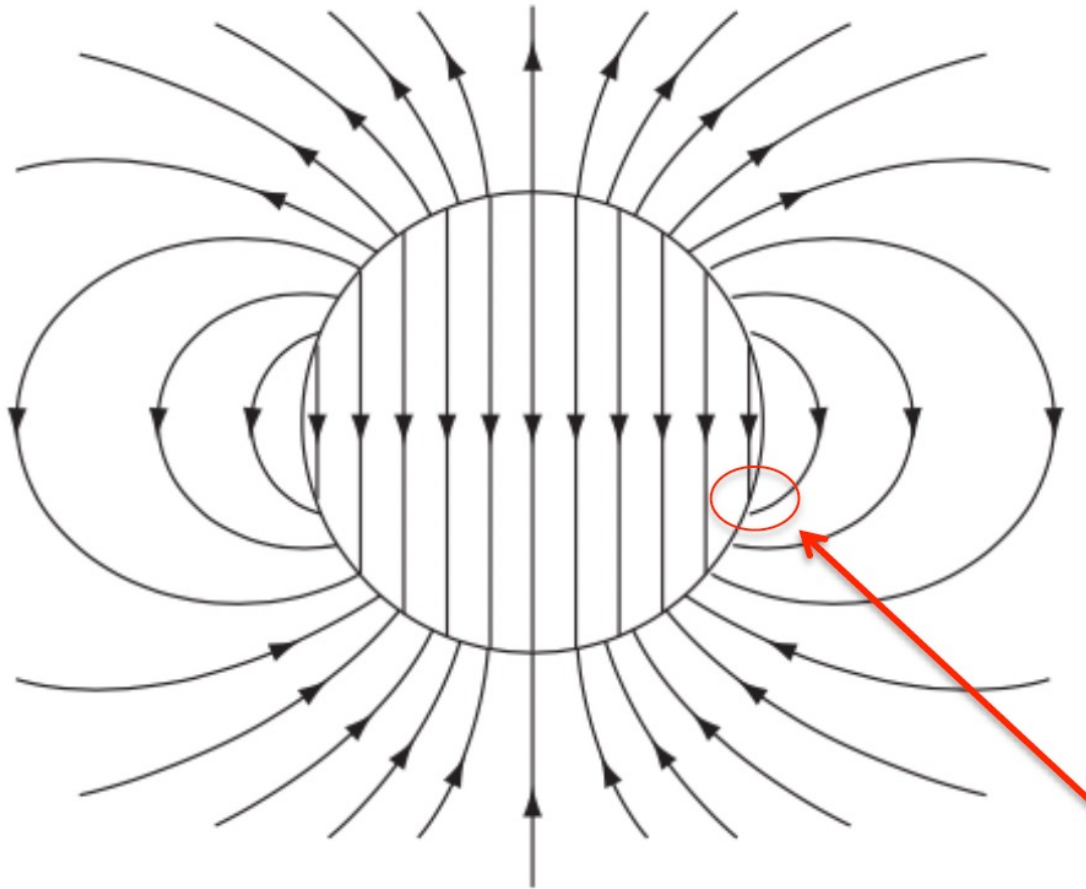
$$V(z) = \frac{r P \cos \theta}{3\epsilon_0} = \frac{P z}{3\epsilon_0}$$

$$\vec{E} = -\nabla V(z) = -\frac{\partial}{\partial z} \left(\frac{P z}{3\epsilon_0} \right) \hat{\mathbf{z}} = -\frac{P}{3\epsilon_0} \hat{\mathbf{z}}$$

*E is uniform within a dielectric
sphere of constant polarization*



Graphic of E field



Field lines are not continuous because of the bound surface charge created by the dipoles (or polarized material)

We learned that there is a discontinuity of the E field

$$\mathbf{E}_{\text{above}} - \mathbf{E}_{\text{below}} = (\sigma/\epsilon_0)\mathbf{n}$$

Since not a conductor, you can have E tangent to spherical surface. But it does have to be continuous across boundary.