

FoP 3B Part II

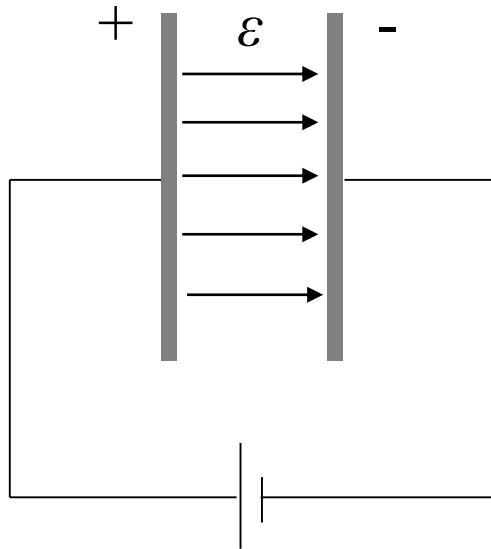
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Room 151

Lecture 10: Polarisation in Dielectrics

Dielectrics in capacitors

Capacitance is defined by:



$$C = \frac{Q}{V} = \epsilon_o \epsilon_r \frac{A}{d}$$

capacitor plate area

dielectric constant

plate separation

Material	Dielectric Constant	Material	Dielectric Constant
Vacuum	1	Benzene	2.28
Helium	1.000065	Diamond	5.7
Neon	1.00013	Salt	5.9
Hydrogen	1.00025	Silicon	11.8
Argon	1.00052	Methanol	33.0
Air (dry)	1.00054	Water	80.1
Nitrogen	1.00055	Ice (-30° C)	99
Water vapor (100° C)	1.00587	KTaNbO₃ (0° C)	34,000

BaTiO_3 } Ferroelectric
 PbTiO_3 } 'perovskites'

- Experimentally it is found that inserting a dielectric in a capacitor causes a smaller potential drop across the plates (charge is however conserved).
- This is due to *polarisation* of the dielectric.

Aim of today's lecture

Definition: Dielectrics are insulator materials that can be polarised by an applied electric field.

► Develop the electrostatic framework used for describing dielectric media

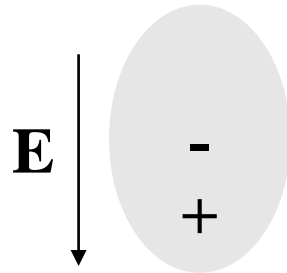
Key concepts:

- Polarisation in dielectrics
- Microscopic vs macroscopic electric fields

Definition of polarisation

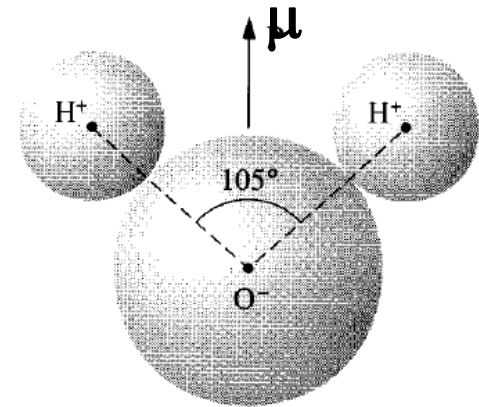


Unpolarised atom
Electric field OFF

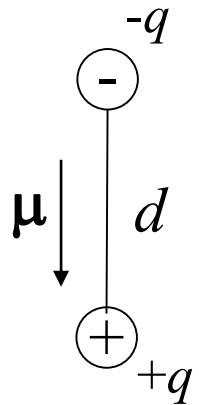


Polarised atom
Electric field ON

More 'complex' polarisations:



Water molecule: *permanent*
electric dipole



Electric dipole moment defined by:

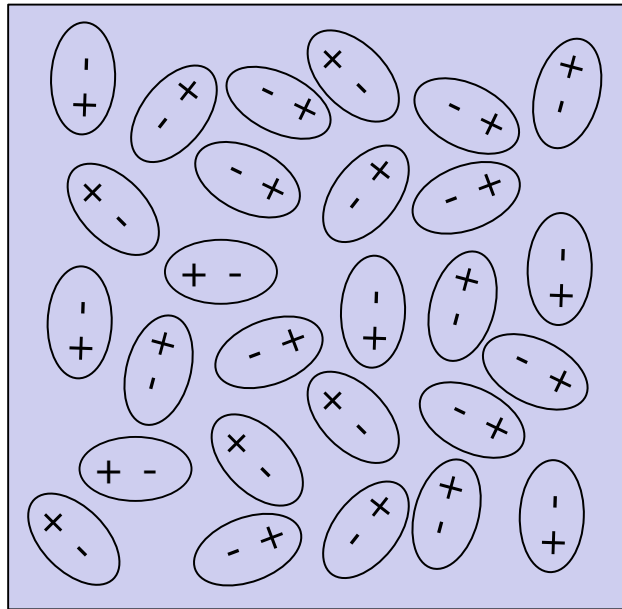
$$|\mu| = qd$$

The direction of μ is from negative
to positive charge.

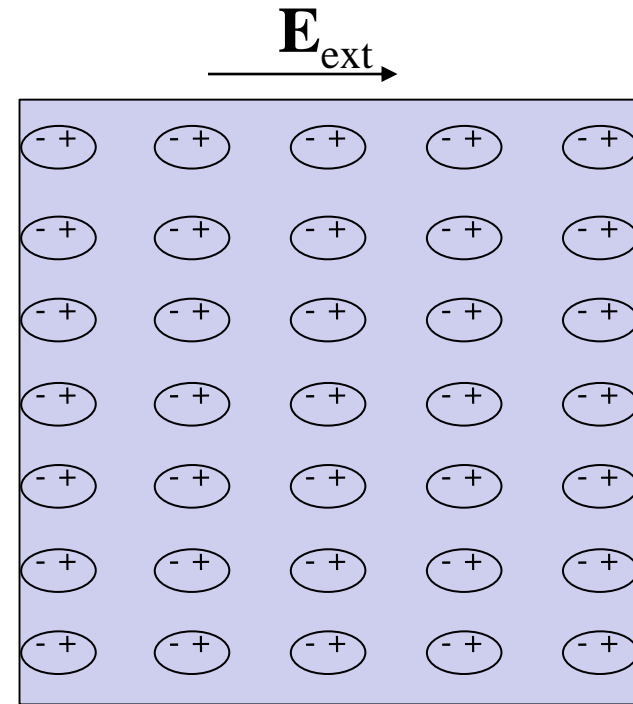
Torque due to electric field: $\tau = \mu \times \mathbf{E}$

Potential energy due to electric field: $U = - \mu \cdot \mathbf{E}$

Dielectric media in an electric field (e.g. capacitor)



Electric field OFF

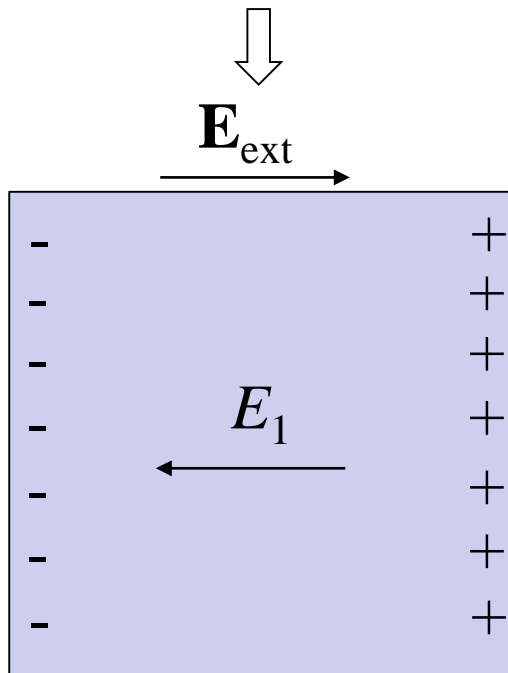
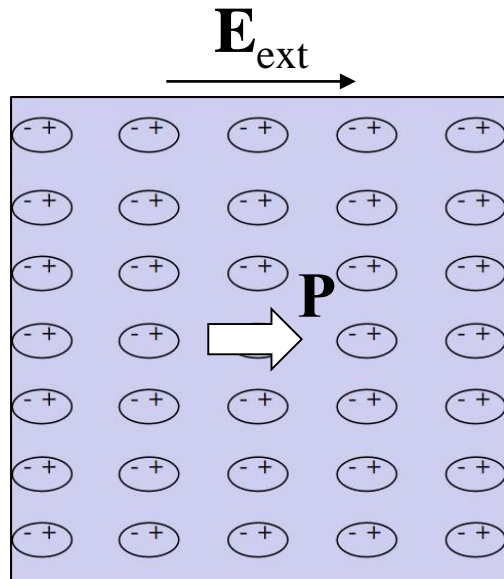


Electric field ON

-Torque on permanent dipoles rotate them to the minimum energy configuration.

-Define polarisation \mathbf{P} as dipole moment per unit volume, i.e. $\mathbf{P} = N\boldsymbol{\mu}$, where N is the number density of dipoles.

Macroscopic electric field



Electric field due to a single dipole is given by*:

$$\mathbf{E}(\mathbf{r}) = \frac{3(\boldsymbol{\mu} \cdot \mathbf{r})\mathbf{r} - r^2\boldsymbol{\mu}}{4\pi\epsilon_0 r^5}$$

For *uniform* polarisation \mathbf{P} the collective effect of all dipoles can be modelled by a surface charge density (σ) *:

$$\sigma = \mathbf{P} \cdot \hat{\mathbf{n}} \quad \leftarrow \text{Unit surface normal vector}$$

A depolarisation field is therefore present:

$$E_1 = -\frac{\sigma}{\epsilon_0} \quad (\text{Gauss' law})$$

Internal field is $(E_{\text{ext}} - |E_1|)$.

* See for example Griffiths, *Introduction to Electrodynamics*, Chapters 3, 4 (non-examinable)

Macroscopic vs microscopic fields

E_1 is a *macroscopic* electric field ‘smoothed’ over many dipoles. The local *microscopic* field at an individual dipole can however differ significantly from E_1 .

The polarisability (α) of a single dipole is defined as:

$$\mu = \alpha E_{\text{local}}$$

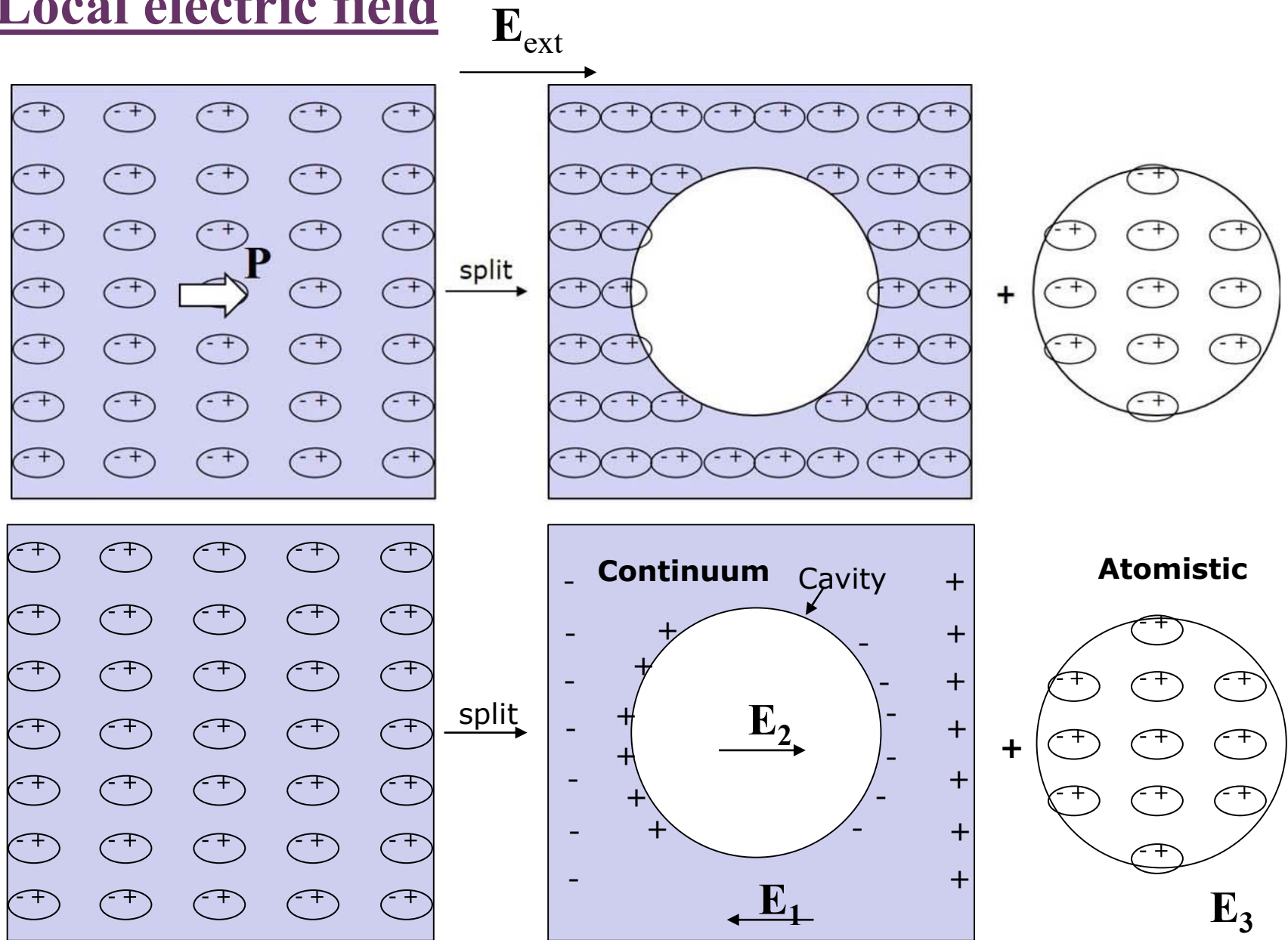
Electric dipole moment Local microscopic electric field

Q: What is the local electric field at a dipole?

Q: How do you calculate polarisability?

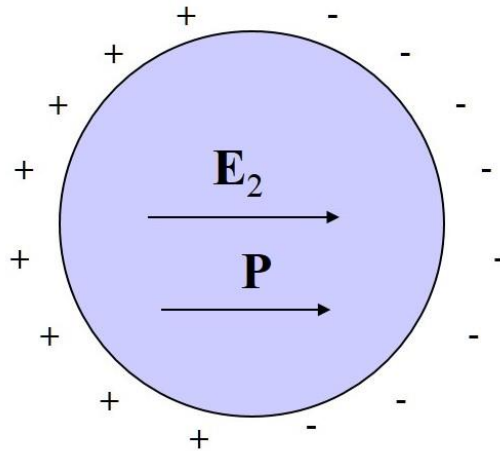
(Clausius-Mossotti relation- next lecture)

Local electric field



$$\text{Local electric field } \mathbf{E}_{\text{local}} = (\mathbf{E}_{\text{ext}} + \mathbf{E}_1) + \mathbf{E}_2 + \mathbf{E}_3$$

Lorentz (E_2) and atomistic (E_3) fields

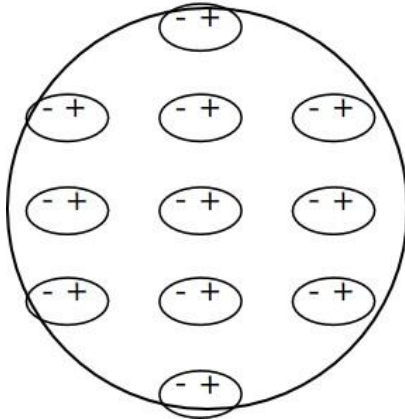


Lorentz field is given by*:

$$\mathbf{E}_2 = \frac{\mathbf{P}}{3\epsilon_0}$$

Atomistic field is given by:

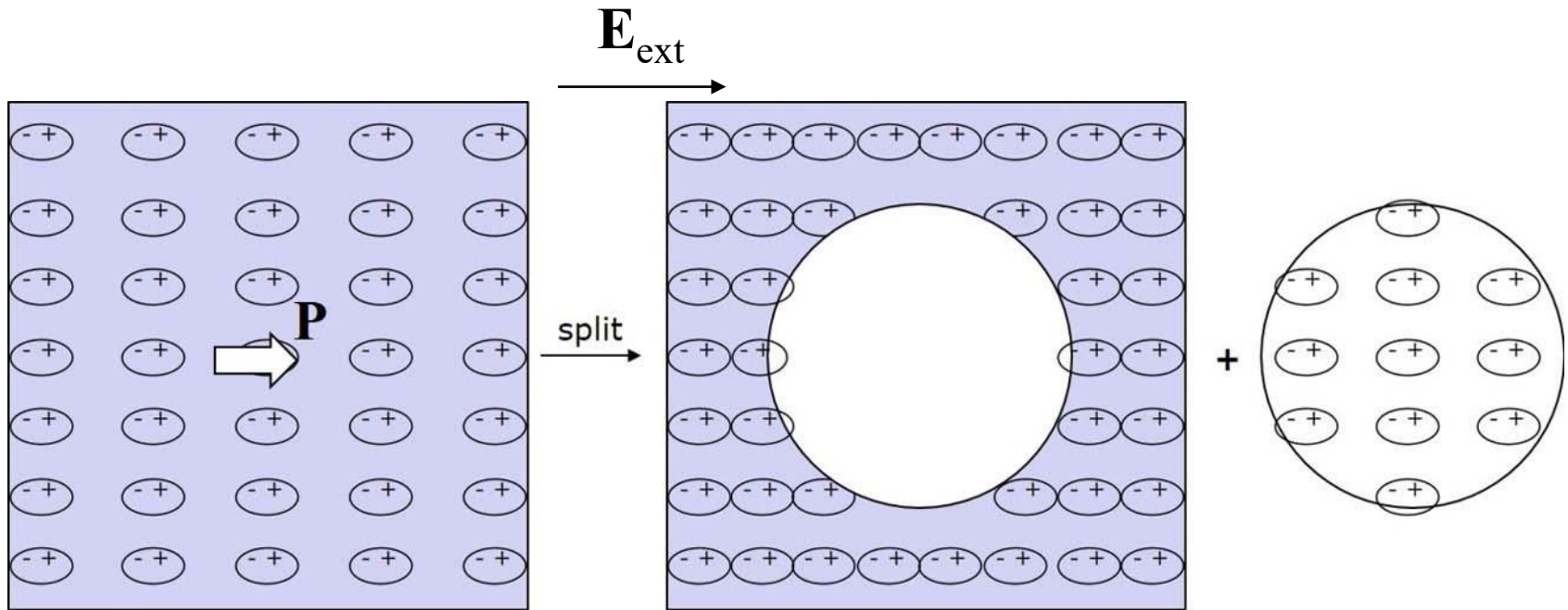
$$\begin{aligned} \mathbf{E}_3 &= \sum_{\text{dipoles } i} \frac{3(\boldsymbol{\mu}_i \cdot \mathbf{r}_i)\mathbf{r}_i - r_i^2 \boldsymbol{\mu}_i}{4\pi\epsilon_0 r_i^5} \\ &= \mu \sum_i \frac{3x_i^2 - r_i^2}{4\pi\epsilon_0 r_i^5} = 0 \end{aligned}$$



For polarisation along x and *cubic* crystal.

\therefore only the Lorentz field alters the microscopic field from the macroscopic field.

Local electric field



$$\text{Local electric field } \mathbf{E}_{\text{local}} = (\mathbf{E}_{\text{ext}} + \mathbf{E}_1) + \mathbf{E}_2 + \mathbf{E}_3$$

$E_1 = -\sigma/\epsilon_0$, $E_2 = P/3\epsilon_0$, $E_3 = 0$ gives:

$$\mathbf{E}_{\text{local}} = \mathbf{E}_{\text{ext}} - \frac{\sigma}{\epsilon_0} + \frac{P}{3\epsilon_0}$$