

Di-Electric properties

~~Dielectric~~ Dielectric materials are basically insulators which ordinary donot contains any charge carrier for conduction they contain positive & negative charges which are tightly bond together because of that they have very low electric conductivity.

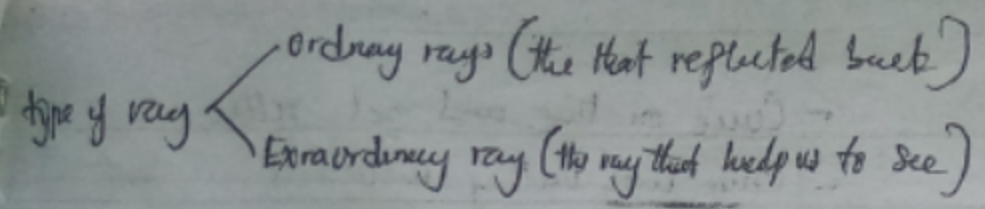
Dielectric materials: Is an electrical insulator that can be polarized by an applied electric field.

A Dielectric is an insulator but not all the insulator is dielectric.

An Insulator: Is a material which does not allows the flow of electric current under the influence of an electric field. While a dielectric is a material with insulation properties which polarizes under the effect of electric field.

Insulators have a low dielectric constant while dielectric have relatively high dielectric constant. Hence, a dielectric can be defined as an insulator that can be polarized. Thus, all dielectric are insulators but all insulators are not dielectric.

A dielectric can hence store stored charges this characteristics make it useful in the form of capacitor.



insulator or good insulator retains heat.
the influence of field dielectrics can conduct e.m.f.
they store charges.
is a good conductor -

dielectric retains heat but they don't give out.

dielectric materials is very low conductivity of

Dies of a dielectric
 Dies - solid
 Dielectric constant
 Dielectric material
 Dielectric loss
 Dielectric strength
 Dielectric absorption
 Dielectric relaxation
 Dielectric permittivity
 Dielectric conductivity
 Dielectric breakdown
 Dielectric polarization
 Dielectric hysteresis
 Dielectric aging
 Dielectric loss factor
 Dielectric loss tangent
 Dielectric loss angle
 Dielectric loss coefficient
 Dielectric loss index
 Dielectric loss ratio
 Dielectric loss factor
 Dielectric loss tangent
 Dielectric loss angle
 Dielectric loss coefficient
 Dielectric loss index
 Dielectric loss ratio

ELECTRIC CONSTANT: Is a measure of
 a material's response to an electric field
 and how much separation of charge

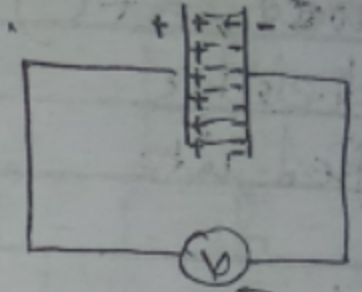
ELECTRIC FUNCTION: will describe the
 standardization or assumptions properties of material
 is a function of wavelength as materials
 respond differently depend on the energy of
 light.

Capacitor: Is an electrical device for
 storing charge. It has the symbol (+ || -)

measured by Anometer or multimeter A

It is (Farad)

Capacitor always set up if (inflow) i.e.



"The capacitor is fully charge
 when it reaches (5v) then the
 source will become zero."

What capacitor does?

- 1) It store charges
- 2) It prevent sparking at the point
- 3) A capacitor smoothing and rectifier (i.e. to reduce the noise) and to rectifier the A.C

Discharging of a capacitor (methods are)

- ① By Rapid method: Is when the source remove and
 join two head of the wire together.
- ② By gradual method: Is when the system is connected
 to the resistor then gradual the will move out.

Di-electric: Is a material in which all electrons
 are tightly bound to the nucleus of an atom i.e.
 no free electron to carry current.

Hence, the electrical conductivity of
 dielectric materials is very low towards

total dielectric material is zero.

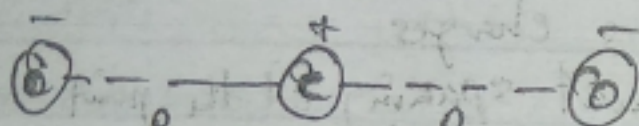
of dielectric materials (paper, air, wood, mica, etc)

molecules in dielectric

are classified into

* Polar

* Non-polar



$$\vec{P} = \vec{P}_1 - \vec{P}_2 = 0$$

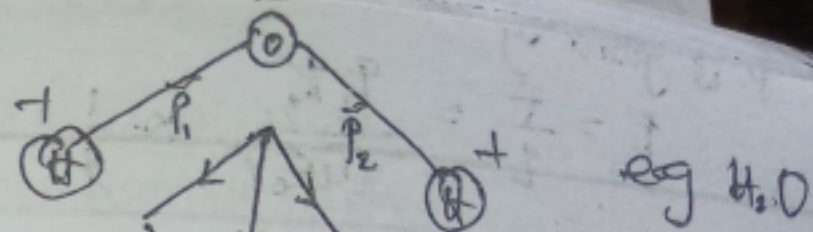
eg C = carbon O = oxygen (twice) = CO₂

non-polar molecule is one in which the center of gravity of the +ve charges (protons) coincides with the center of the gravity of the -ve (electrons)

eg non-polar molecule has symmetrical structure & zero electric dipole moment.

Examples of Non-polar

- 1. O₂
- 2. Oxygen (O₂)
- 3. Nitrogen (N₂)
- 4. Hydrogen (H₂)
- 5. Benzene (C₆H₆)



$$\vec{P} = \vec{P}_1 - \vec{P}_2$$

A polar molecule is one in which the center of gravity of the +ve charges (protons) is separated from the center of gravity of -ve charges (electrons) by a finite distance. i.e. distance that you can not control (tend to infinity). A polar molecule has an intrinsic dipole moment or intrinsic permanent dipole moment.

Example

1. H₂O
2. Carbon monoxide (CO)
3. HCl
4. N₂O

Total dipole moment

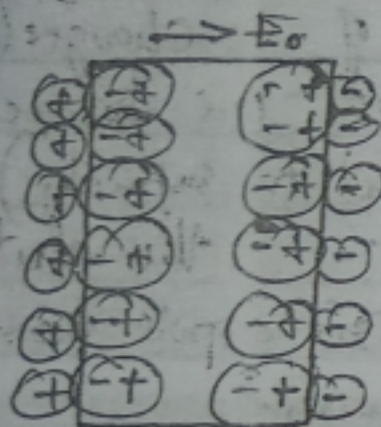
$$\vec{P} = \sum q_n \vec{r}_n$$

r_n is the position vector of the charge (q_n)
The electric field at the point $P(r)$ from a dipole moment

$$E = \frac{F}{Z} = \frac{q_1 q_2}{4\pi\epsilon_0 r^2} \propto \frac{1}{r^2}$$

$$E = \frac{F}{Z} = \frac{q_1}{4\pi\epsilon_0 r^2} \quad \text{--- (2)}$$

Polarization



When a dielectric is placed in an external electric field (E_0) the $+$ and $-$ charges are displaced from their equilibrium position by very small distances throughout the volume of a dielectric.

This results in a formation of a large no. of dipoles each having some dipole moment in the direction of the field. The material ~~is~~ is said to be polarized. The polarization (\hat{P}) is defined as the dipole moment per volume average over the volume

of the cell.

The effect of polarization is to reduce the effect of the external field (E_0). Hence, the magnitude of the resultant field is less than the applied field. [$E < E_0$]. This implies that:

$$E = E_0 + E_1$$

where E_1 is called depolarization field. For within the body it tends to oppose the applied field.

For ordinary electric fields the polarization field is proportional to macroscopic field (E), i.e. $\hat{P} \propto E$.

By removing the proportionality

$$\hat{P} = \epsilon_0 \lambda_e E \quad \text{--- (3)}$$

E is the macroscopic field. λ_e is the electric susceptibility. The electric displacement vector (D) for an isotropic or cubic medium is defined as:

$$D = \epsilon_0 \epsilon_r E = \epsilon_0 E + \hat{P} \quad \text{--- (4)}$$

ϵ_r is the relative permittivity. ϵ_r is related to dielectric susceptibility as $\epsilon_r = 1 + \lambda_e$ --- (5)

- avoid any form of distraction

part of primary interest in the wave theory
field vectors (E and D), and \hat{p} , χ , ϵ_r
these interest we must go to macroscopic ^{electric field}

Macroscopic electric field.

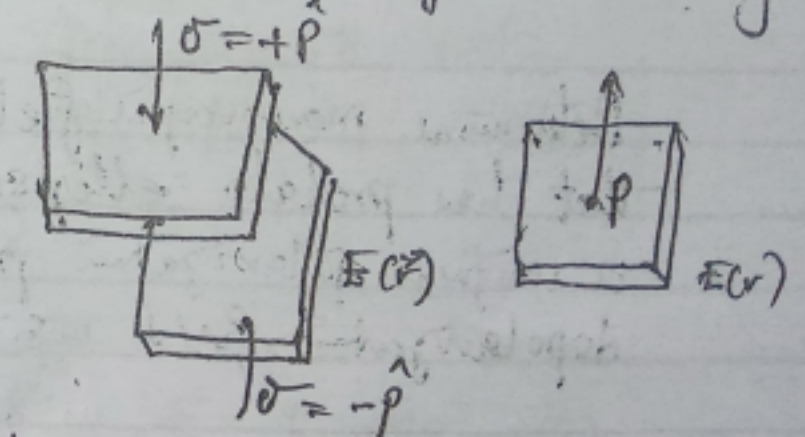
The contribution to electric field inside a body
applied electric field. It's defined as E_0
 E_0 = field produced by forces external to the body
Contribution to the electric field is the sum of
of all charges that constitute the body.
body is neutral the contribution to the
field may be expressed as

A macroscopic field is the volume
average over a region large compared
atomic dimensions, so the defined
the field $E(r)$ as the average field
in the volume of the crystal cell
at contains the lattice point (r_0)

$$E(r_0) = \frac{1}{\epsilon_0} \int dV e(r) \quad \text{--- (2)}$$

To find the contribution
Here $e(r)$ is the microscopic electric field at the point (r)
where E is the macroscopic electric field.
To find the contribution of the polarization to the
microscopic field we can sum over all the dipoles
in a specimen. Applying electrostatic theorem
the macroscopic field caused by a uniform
polarization (P) is equal to the electric field
in vacuum of a fictitious surface
charge density.

$\sigma = \hat{n} \cdot \hat{P}$ on the surface on the body.
Unit normal to the surface.

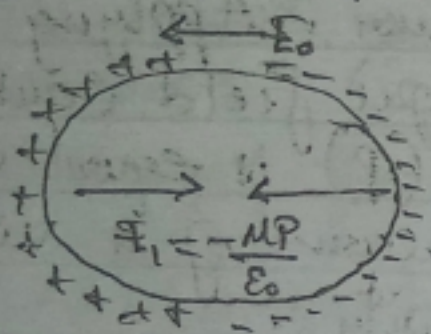


Then, we now have
 $\sigma = \hat{n} \cdot P$ on the surface of the slab.
 $\sigma = \hat{n} \cdot P = P$ per unit area
 $\sigma = \hat{n} \cdot P = -P$ the lower boundary bears $-P$
By Gauss's law E_i due to the charge σ given by
 $E_i = \frac{P}{\epsilon_0}$

lecturer material
 here inside a dielectric
 $\epsilon_0 + E_1 = E_0 - \frac{P}{\epsilon_0}$
 is a unit vector normal to the plane of
 ab.

= field of the surface charge density (σ) on the bounding

Depolarization field



$$E = E_0 + E_1$$

Determine macroscopic field within the dielectric that has prolate ellipsoidal form. In ellipsoid a uniform polarization produces a uniform depolarization field inside the body.

E_1 is a field due to polarization. The polarization \hat{p} produces a surface distribution of charge whose polarity gives rise to a field within the dielectric that's directly opposite to the applied field (E_0) in the figure above.

Is called the depolarization field (E_1)
 Let P_x, P_y, P_z be the components of the polarization

$$E_{1x} = - \frac{N_x P_x}{\epsilon_0}$$

$$E_{1y} = - \frac{N_y P_y}{\epsilon_0}$$

$$E_{1z} = - \frac{N_z P_z}{\epsilon_0}$$

We now have depolarization factors:

N_x, N_y, N_z depolarization factors.

Their value depends on the ratio

N_x, N_y, N_z have +ve values

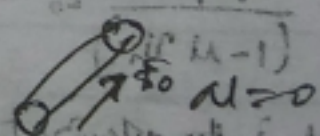
$$N_x + N_y + N_z = 1$$

$$E_1 = - \frac{N P}{\epsilon_0}$$

macroscopic field, $E = E_0 - \frac{N P}{\epsilon_0}$



$$N = \frac{1}{3}$$



$$N = 0$$

$N = 1$ is the geometrical symmetrical polarization

Shape	Axis	γ
Sphere	any	$\frac{1}{3}$
Thin slab	in plane	0
Thin slab	normal	1
long Circular Cylinder	longitudinal	0
long Circular Cylinder	Transverse	$\frac{1}{2}$

* \hat{P} = polarizability
 P = dipole moment

Polarization (\hat{P}) is proportional to the macroscopic \vec{E} and is given by

$$\hat{P} = \epsilon_0 \chi_e \vec{E} \quad (1)$$

In isotropic medium, the displacement vector \vec{D} is

$$\vec{D} = \epsilon_0 \epsilon_r \vec{E} = \epsilon_0 \vec{E} + \vec{P} \quad (2)$$

we can now say

$$\epsilon_r = 1 + \chi_e \quad (3)$$

$$\hat{P} = \frac{\epsilon_0 \chi_e}{(1 - \chi_e)} \vec{E}_0 \quad (4)$$

where χ_e is the polarizability factor.
 The value of polarizability (\hat{P}) depends on the local polarization field

come on the
 come with uniting materials

(2)

The local field in an atom
 The local electric field denoted by (E_{loc}) that act at the site of an atom in a crystal is not the same as the macroscopic average field (\vec{E})
 To understand this let's consider the local field at the site with a cubic arrangement in a crystal of a spherical shape.

$$\vec{E} = \vec{E}_0 + \vec{E}_s, \quad \vec{E}_s = \vec{E}_0 - \frac{1}{3\epsilon_0} \vec{P} \quad (5)$$

Consider the field that act on the atom at the center of the sphere, let all the dipoles be parallel to the z-axis and have magnitude (p) , then the component of the field along the z-axis at the center due to other dipoles is given as

$$E_{spher} = \frac{P}{4\pi\epsilon_0} \sum \frac{2z_i^2 - r_i^2}{r_i^5} \\
= \frac{P}{4\pi\epsilon_0} \sum \frac{2z_i^2 - x_i^2 - y_i^2}{r_i^5} \quad (6)$$

Because of the symmetry of the lattice and the sphere the x, y, z direction dimensions are the same & equivalent

$$\sum \frac{z_i^2}{r_i^5} = \sum \frac{x_i^2}{r_i^5} = \sum \frac{y_i^2}{r_i^5} \quad (7)$$

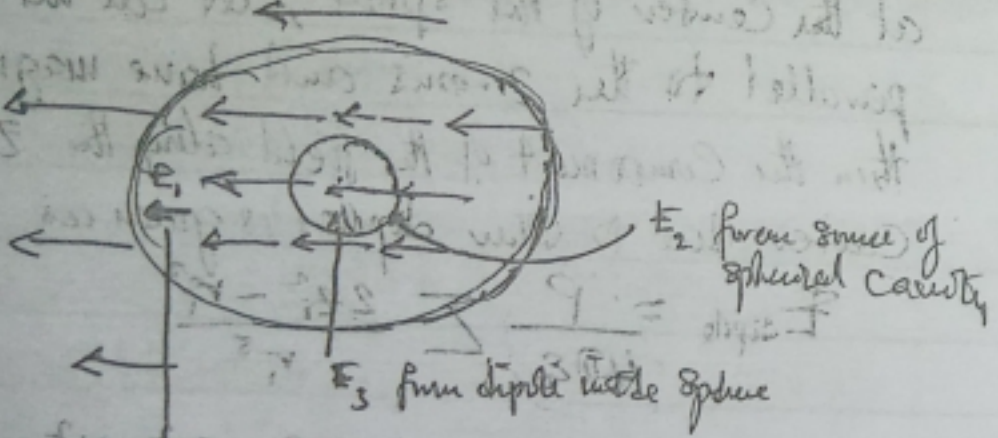
then, the $E_{spher} = 0$

lecture main
 $\vec{E}_{loc} = \vec{E}_0$ outside local cell

atom in a cubic environment in a spherical cavity

in a general lattice site, i.e. not necessary of cubic symmetry, the local field at an atom is the sum of the applied electric field given as (\vec{E}_0) , from external sources, and the field from the dipole within the specimen, this is given as:

$$\vec{E}_{loc} = \vec{E}_0 + \vec{E}_1 + \vec{E}_2 + \vec{E}_3 \quad (8)$$

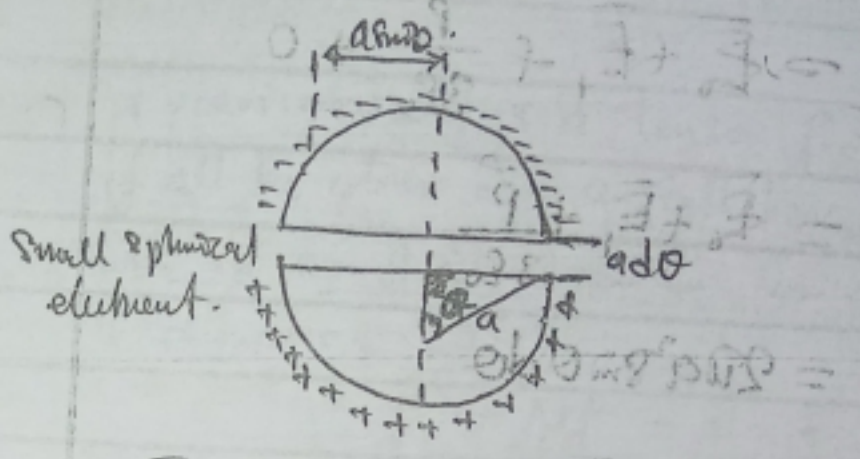


from outer boundary $(E_1) \sum \frac{q}{3\pi r^2} =$
 from fixed charges external to the body charges.

field from the surface charge density on the outer boundary
 the specimen is removed
 when Lorentz cavity field (a field from polarized dipoles inside a spherical cavity cut out of the specimen) hence atom as a center.

Come on
 Come with writing materials
Multipole E_{pol} is dependent on the geometrical shape
 Dipole local field at center of hollow sphere

$$E_1 = -\frac{1}{3\epsilon_0} P \quad (9)$$



The Cavity. $E + P = E_{tot}$

The surface charge density on the surface of cavity = $-p \cos \theta$
 The area of the ring shape element $abd = 2\pi a^2 \sin \theta d\theta$
 The charge on the ring ~~shape~~ element $q = p \cos \theta \cdot 2\pi a^2 \sin \theta d\theta$
 Thus, the electric field at the centre of cavity or radius a is given as:

$$\vec{E}_2 = \int \frac{1}{4\pi \epsilon_0} (a^2) (2\pi a^2 \sin \theta d\theta) (p \cos \theta) \cos \theta = \frac{P}{3\epsilon_0} \quad (10)$$

eqn (10) is the one of the field in a polarized sphere.

(E_3) for a reference site with cubic symmetry in sphere ($E_3 = 0$) It depends on the structure of the

$$E = \alpha E_0$$

atoms are replaced by point dipoles
 is each other than the total local field
 eqn (1) & (2)

$$E_{loc} = \bar{E}_0 + \bar{E}_1 + \frac{P}{3\epsilon_0} + 0$$

$$= \bar{E}_0 + \bar{E}_1 + \frac{P}{3\epsilon_0}$$

$$= 2Na^2 \epsilon_0 \omega_0^2 d_0$$

But $\bar{E} = \bar{E}_0 + \bar{E}_1$

$$\bar{E}_{loc} = \bar{E} + \frac{P}{3\epsilon_0}$$

we did you generate field?
 yet we have field?
 what are the steps?

Polarizability α of an atom is defined as the
 dipole moment induced with local field of an atom

$$P = \alpha E_{loc} \quad (1)$$

Explain
Question

Polarizability is an atomic property but dielectric
 constant is the macroscopic property dependent on atoms
 within the crystal.

Distinguished between polarizability & dielectric
 constant.

Polarizability is a tensor for non spherical atoms
 if all the atoms in a crystal have the same polarizability
 with n no of atom per unit volume. The polarization can
 be expressed as:

$\bar{P} = NP = N\alpha \bar{E}_{loc}$ (2)
 For n numbers of atom in a crystal the polarization is
 expressed as:

$$\bar{P} = \sum_i N_i p_i = \sum_i N_i \alpha_i \bar{E}_{loc} \quad (3)$$

where N_i is the concentration =
 \bar{E}_{loc} is the local field at the site
 p_i is the dipole at the i th atom
 α_i is the polarizability at the i th atom.

Substituting eqn (1) & (2)

$$\bar{E}_{loc} = \bar{E} + \frac{\bar{P}}{3\epsilon_0}$$

$$P = (\sum N_i \alpha_i) \left[\bar{E} + \frac{P}{3\epsilon_0} \right] = \frac{1-n}{1+n} P$$

$$\frac{\alpha_i}{3\epsilon_0} = \frac{P}{\epsilon_0 E}$$

$$= \frac{\sum N_i \alpha_i}{\epsilon_0 \left[1 - \frac{\sum N_i \alpha_i}{3\epsilon_0} \right]}$$

$$\epsilon_r = 1 + \frac{\sum N_i \alpha_i}{\epsilon_0 \left[1 - \frac{\sum N_i \alpha_i}{3\epsilon_0} \right]}$$

$$\frac{1}{r+2} = \frac{1}{3\epsilon_0} \sum N_i \alpha_i$$

Classical + Lorentz + Mossott's Relation

$$\frac{n^2 - 1}{n^2 + 2} = \frac{4\pi N}{3\epsilon_0} \sum f_i \alpha_i$$

Let's polarizability have the below form

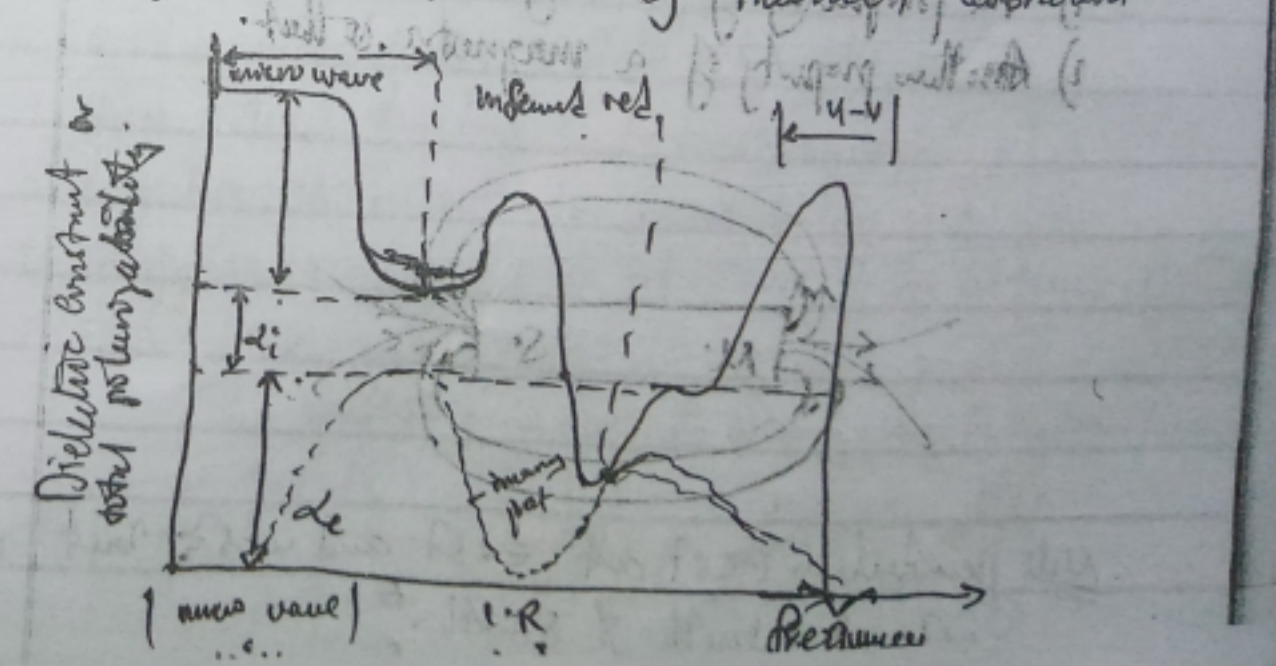
$$\alpha = \alpha_e + \alpha_i + \alpha_d$$

α_e electronic polarizability
 α_i ionic
 α_d - dipolar

① Electronic polarizability (α_e): It involves the displacement of electrons in an atom relative to the nucleus under the influence of electric field.

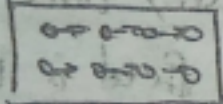
② The ionic polarization: The movement of dipole by relative movement by (+ve) and (-ve) ions, in ionic solid under the influence of an electric field is known as ionic polarizability (α_i)

③ Dipole Orientation or dipolar polarizability (α_d): This disorder to align all permanent dipoles with the apply field is frustrated by thermal vibrations.



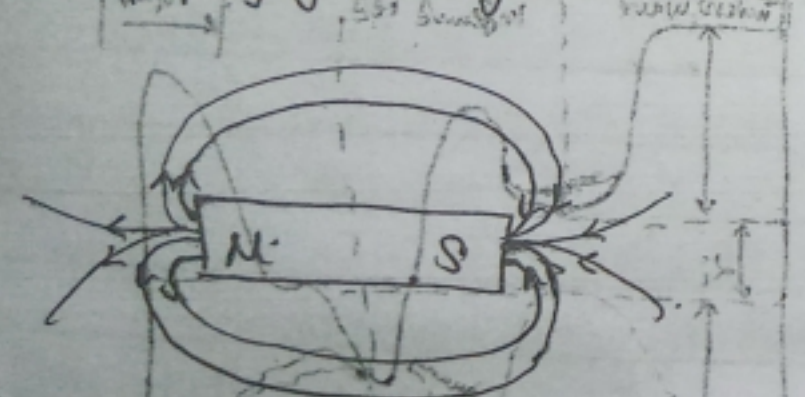
ability of a material to be induced influence of a magnetic field known as magnetism.

~~Further writing that~~
 magnetic moments are induced by magnetic field. Every material is a matter and it has molecules. The molecules in magnetic are called domains.



Is that materials that has a field?

The property of a magnet is South & North pole. Another property of a magnet is that...



Always rest at east and west but magnet...

- 1) Ferromagnetic (ready material) materials
- 2) Paramagnetic material
- 3) Diamagnetic materials.

Ferromagnetic material: They have what we called relative permeability (ϵ_r). They are materials that easily attracted.

- Examples of ferromagnetic materials:
- i) Nickel
 - ii) Cobalt
 - iii) Iron

- Characteristics of ferromagnetic materials:
- 1) Relative permeability is greater than one i.e. $\epsilon_r > 1$
 - 2) They are dependent on a field strength.
 - 3) They attract the line of force strongly.
 - 4) They have high magnetic susceptibility.
 - 5) Magnetic properties exist even in the absence of applied magnetic field.

The spinning... they are parallel to each other.

And arises out of electron structure
 about

Diamagnetic materials. The atoms and the molecules of diamagnetic material has zero magnetic field in a absence of an external field.

With application of the external field a net dipole moment opposing the field (Lenz law) is induced. Diamagnetic material arises from Lenz law.

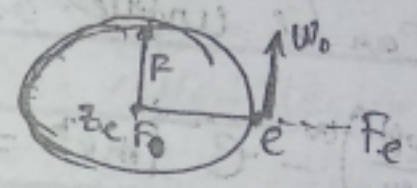
Examples of Diamagnetic materials

- Silver
- Copper
- Bismuth
- Hydrogen
- Graphite
- Water
- Crystal

Characteristics of Diamagnetic material
 In a diamagnetic the relative permeability is slightly less than one. The magnetic susceptibility (χ) is very small and negative. Susceptibility is determined by the electron structure

of the system does not depend on external condition like temperature.

4) Repeat the line of force slightly.



Consider a circular orbital of radius r in which an electron revolves with an angular velocity ω around a nucleus (Ze)

Force of attraction between electron & nucleus plays a roles of centripetal force

$$F_c = m\omega^2 r = \frac{e^2}{r^2} \quad \text{--- (1)}$$

$$\omega = \left(\frac{e^2}{mr^3} \right)^{1/2} \quad \text{--- (2)}$$

In the presence of the magnetic field (H) there is additional force known as Lorentz force.

$$F_2 = -e v \times H \quad \text{--- (3)}$$

$$F_2 = e r \omega H \text{ if } H \text{ is } (1) \text{--- (4)}$$

$\frac{e^2}{r^2} \dots$

$$m\omega^2 = \frac{e^2}{r^2} - e r \omega^2 \quad (1)$$

Equation (1) can be written as

$$\omega^2 + \left(\frac{e\hbar}{m}\right)\omega - \frac{e^2}{mr^2} = 0 \quad (2)$$

thus for ω

$$\omega = \frac{-\frac{e\hbar}{m} \pm \sqrt{\left(\frac{e\hbar}{m}\right)^2 + 4\frac{e^2}{mr^2}}}{2} \quad (3)$$

$$\omega = -\frac{e\hbar}{2m} \pm \sqrt{\left(\frac{e\hbar}{2m}\right)^2 + \frac{e^2}{mr^2}}$$

In eqn (2)

$$\omega = -\frac{e\hbar}{2m} \pm \sqrt{\omega_0^2 + \left(\frac{e\hbar}{2m}\right)^2} \quad (4)$$

If $\frac{e\hbar}{2m} \ll \omega_0$ then $\omega = \omega_0 - \frac{e\hbar}{2m}$

$$\omega = -\frac{e\hbar}{2m} + \omega_0 \quad (5)$$

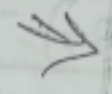
~~$$m\omega^2 = \frac{e^2}{r^2} - e r \omega^2$$~~

Paramagnetic

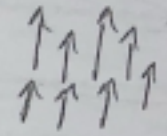
$v(1)$ is the ...

Eqn (2) shows the apply field changes ω_0 to ω (\pm) on ω_0 implies electron with orbital moment parallel to the field or slowed down and by amount $(\frac{e\hbar}{2m})$ & is Larmor theorem and the term $(\frac{e\hbar}{2m})$ are called Larmor frequency.

Larmor

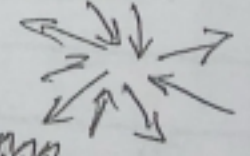


ω is the resonant frequency. The change in frequency gives rise to an additional current and therefore change in orbital magnetic dipole moment or change in magnetization.



PARA MAGNETIC MATERIAL

Before the external magnetic field.



↑↑↑↑ measure of the external magnetic field.

Paramagnetism: This is a result of the orientation of already existing magnetic dipole due to the presence of external magnetic field. In a paramagnetic gas subjected to applied magnetic field it has a Curie law.

$$= \frac{2J+1}{2J} \coth\left(\frac{2J+1}{2J} x\right) - \frac{1}{2J} \coth\left(\frac{x}{2J}\right) \quad (4)$$

is Brillouin function

$$\chi = \frac{gJ\mu_B N}{k_B T} \left(H + \lambda M \right) \quad (5)$$

$H = 0$ in spontaneous magnetization eqn (5)

$$\chi = \frac{gJ\mu_B N}{k_B T} = \frac{gJ\mu_B N \lambda M}{k_B T}$$

$$M(T) = \frac{\chi k_B T}{gJ\mu_B N} \quad (7)$$

As $T \rightarrow 0$, $\chi \rightarrow \infty$ and $B_J(x) \rightarrow 1$
 the magnetic moment then aligns parallel to the field
 and M becomes saturation magnetization $M_s(0)$

eqn 5 becomes

$$M_s(0) = N g J \mu_B \quad (8)$$

eqn (7) divide by eqn (8)

$$\frac{M(T)}{M_s(0)} = \frac{\chi k_B T}{N g^2 J^2 \mu_B^2 \lambda}$$

$$M(T) = N g J \mu_B (x) = M_s(0) B_J(x)$$

Example

$$\frac{M(T)}{M_s(0)} = B_J(x) \quad (10)$$

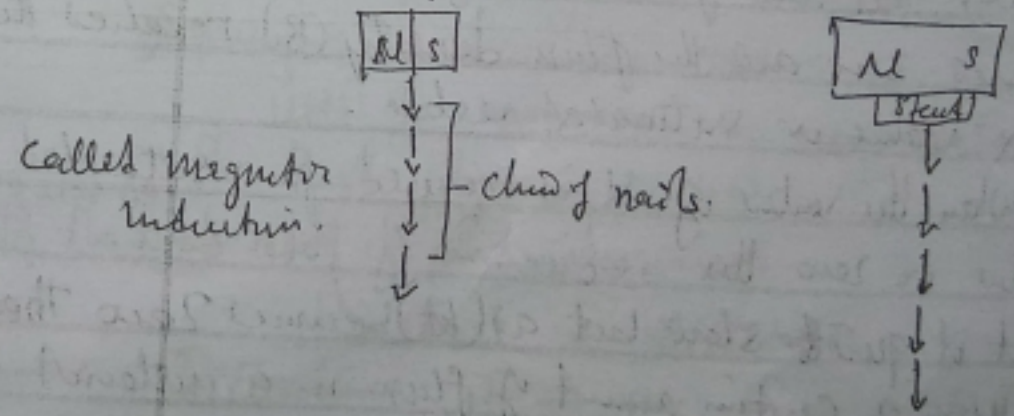
~~What is the value of~~ This is how we obtained it:

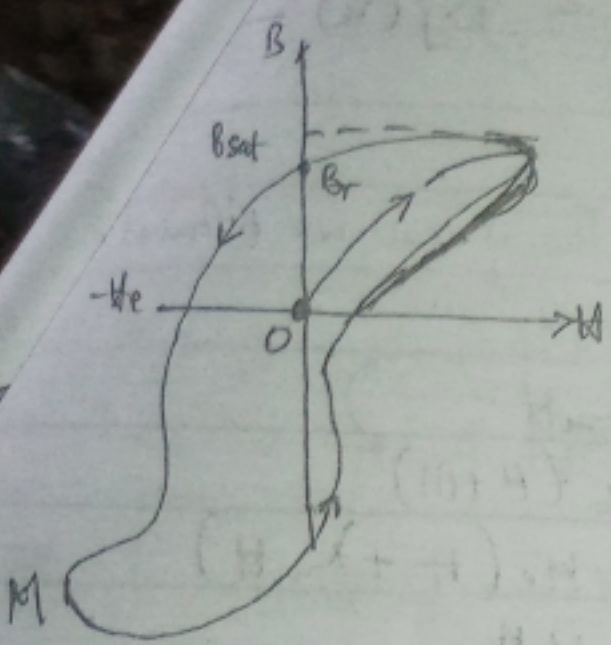
From $B = \mu_0 H$
 $M = \chi_{int} H$
 $B = \mu_0 (H + M)$
 $B = \mu_0 (H + \chi_{int} H)$
 $B = \mu_0 H$
 $\mu = \mu_0 (1 + \chi_m)$

Iron

Susceptibility measures the ease with which

Notes: Iron easily loses its magnetism but steel hardly
 loses its magnetism.





The B-H curve when the temp of ferromagnetic material is below T_c it exhibits normal behavior & follow the ~~extensive~~ ^{well known} hysteresis curve as shown above.

The flux ~~start~~ builds up from the origin (O) at a value of magnetic field (H) in a range μ is nearly proportional to (H) given ratio to constant permeability of the material when H is further increase the rate of increase of (B) falls and eventually becomes zero as the flux density (B) reaches the max value or saturated value.

When the value of H is reduced from saturated value to zero the reduction of flux does not follow ^{the same path} but it quite slow but as (H) becomes zero there remain a certain amount of flux in a material.
 Called remnant flux given as (B_r)

extensive

Below Curie temp (it's raising temp at which a given material ceases to be ^{ferromagnetic} or the falling temp at which it become ferromagnetic).

Ex: \rightarrow All ferromagnetic materials exhibit phenomenon called ~~extensive~~ which is defined as locality of magnetization w induction flux density (B) beside the magnetization force (H) or it's that quality of a substance to which energy is dissipated of the reverse of the magnetism

Magnetic Resonance

Is the absorption of energy emitted by particles (as atomic nuclei w electron) in a static magnetic field when the particles are expose to electromagnetic radiation to a certain frequency

Magnetostriction

1) When a ferromagnetic material magnetized small changes in dimension occurs

uses of magnetostriction

- 1) for high frequency oscillator
- 2) we use it for super sonics

Defects in Metals

Dislocation: it can come in two ways

- a) line dislocation
- ii) screw dislocation

Dis result of deformation

a measure
force if too much on a material it can cause
dislocation.

Dislocation and fracture

Defect in material

There are two types of dislocation.

Line dislocation

1) Screw dislocation

Advantage

1) If one is dead the other crystal will contribute to fill that gap.

Causes - Too much force on a material it cause dislocation.