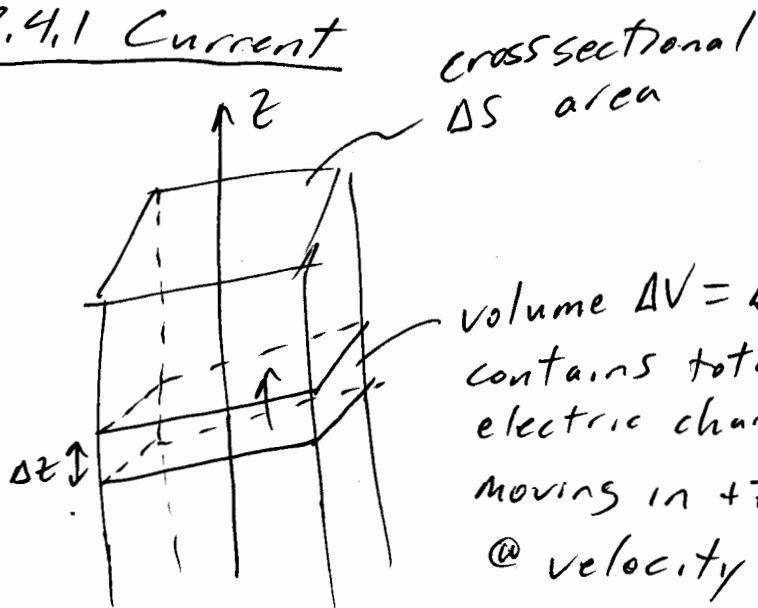


2.4 Current, Conductors, & Conductivity

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→ conductors are materials where there are charges (electrons) free to move when an electric field is applied.

2.4.1 Current



$$\Delta I = \lim_{\Delta t \rightarrow 0} \frac{\Delta Q_e}{\Delta t} = \lim_{\Delta t \rightarrow 0} q_v \Delta S \frac{\Delta z}{\Delta t} = q_v v_z \Delta S \quad (A)$$

Define current density magnitude as

$$J_z = \lim_{\Delta S \rightarrow 0} \frac{\Delta I}{\Delta S} = q_v v_z \quad (A/m^2)$$

In vector form

$$\vec{J} = q_v \vec{v} \equiv \text{Convection current density}$$

This can be broken into positive & negative charges/parts

$$\vec{J} = q_v^+ \vec{v}^+ + q_v^- \vec{v}^-$$

2.4.2 Conductors

- Materials where valence electrons are not tightly bound ⇒ 'free' electrons
- At rest, the free electrons move in random directions ⇒ $I_{net} = 0$ through any surface

→ Excess free charges introduced into a conductor migrate/flow to surface due to repulsive Coulomb forces. The volume charge density inside the conductor is governed by

$$\rho_v(t) = \rho_{v0} e^{-t/\tau_r}$$

where $\tau_r = \epsilon/\sigma$ (s) ≡ relaxation time constant

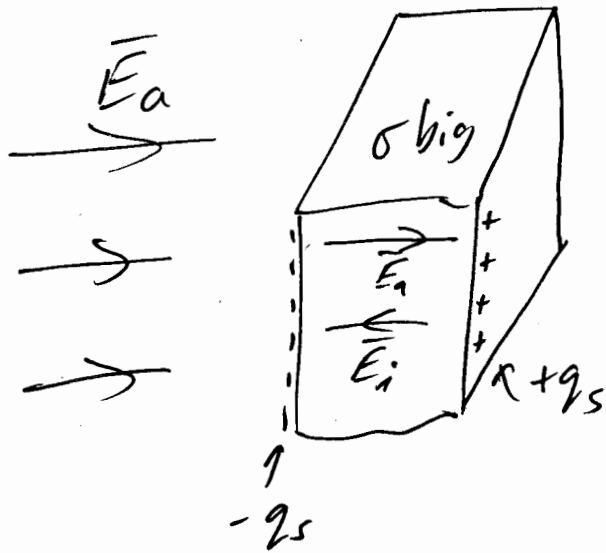
σ ≡ conductivity (S/m) ← Next section

Other properties

- For perfect electrical conductors (PEC) and very good conductors (σ very large)

$$\Delta V_{inside} = 0, \quad \vec{E}_{t,inside} = \vec{D}_{t,inside} = 0$$

Why?



① Applied electric field \vec{E}_a causes charges to move $\vec{F} = q\vec{E}_a$ leading to positive & negative surface charge densities

② The newly formed surface charge densities cause/create an induced electric field \vec{E}_i . \vec{E}_i , in opposition to \vec{E}_a will keep getting bigger & bigger until $\vec{E}_t = \vec{E}_i + \vec{E}_a = 0$ and charges stop moving!

③ By $\vec{D} = \epsilon \vec{E} \Rightarrow \underline{\underline{\vec{D}_t = 0}}$

By $\Delta V = - \int_A^B \vec{E} \cdot d\vec{x} \Rightarrow \Delta V = 0$ inside conductor

\Rightarrow all points in conductor @ same potential!

2.4.3 Conductivity

To characterize the flow of charges in a conductor, we'll define a quantity / parameter called conductivity σ

→ Apply an external electric field \vec{E}_a to a conductor. The free electrons which were moving in random directions w/ random velocities now, on average, move at a drift velocity \vec{v}_e in a direction opposite to \vec{E}_a ($\vec{v} = q\vec{E}_a$ w/ q negative)

⇒ Conduction current ←

Define $\vec{v}_e = -\mu_e \vec{E}$ where $\mu_e \equiv$ electron mobility $\left(\frac{m^2}{V \cdot s}\right)$

Use \vec{v}_e in $\vec{J} = q_n \vec{v}$ to get

$$\vec{J} = -q_{ve} \mu_e \vec{E} = \sigma_s \vec{E}$$

Where the static conductivity $\equiv \sigma_s = -q_{ve} \mu_e \left(\frac{S}{m}\right)$

Older texts resistivity = $\rho = \frac{1}{\sigma}$ ($\Omega \cdot m$)

⇒ See Table 2-3 ←

TABLE 2-3 Typical conductivities of insulators, semiconductors, and conductors

Material	Class	Conductivity σ (S/m)
Fused quartz	Insulator	$\sim 10^{-17}$
Ceresin wax	Insulator	$\sim 10^{-17}$
Sulfur	Insulator	$\sim 10^{-15}$
Mica	Insulator	$\sim 10^{-15}$
Paraffin	Insulator	$\sim 10^{-15}$
Hard rubber	Insulator	$\sim 10^{-15}$
Porcelain	Insulator	$\sim 10^{-14}$
Glass	Insulator	$\sim 10^{-12}$
Bakelite	Insulator	$\sim 10^{-9}$
Distilled water	Insulator	$\sim 10^{-4}$
Gallium arsenide (GaAs)*	Semiconductor	$\sim 2.38 \times 10^{-7}$
Fused silica*	Semiconductor	$\sim 2.1 \times 10^{-4}$
Cross-linked polystyrene (unreinforced)*	Semiconductor	$\sim 3.7 \times 10^{-4}$
Beryllium Oxide (BeO)*	Semiconductor	$\sim 3.9 \times 10^{-4}$
Intrinsic silicon	Semiconductor	$\sim 4.39 \times 10^{-4}$
Sapphire*	Semiconductor	$\sim 5.5 \times 10^{-4}$
Glass-reinforced Teflon (microfiber)*	Semiconductor	$\sim 7.8 \times 10^{-4}$
Teflon quartz (woven)*	Semiconductor	$\sim 8.2 \times 10^{-4}$
Dry soil	Semiconductor	$\sim 10^{-4} - 10^{-3}$
Ferrite(Fe ₂ O ₃)*	Semiconductor	$\sim 1.3 \times 10^{-3}$
Glass-reinforced Polystyrene*	Semiconductor	$\sim 1.45 \times 10^{-3}$
Polyphenylene oxide (PPO)*	Semiconductor	$\sim 2.27 \times 10^{-3}$
Glass-reinforced Teflon (woven)*	Semiconductor	$\sim 2.43 \times 10^{-3}$
Plexiglas*	Semiconductor	$\sim 5.1 \times 10^{-3}$
Wet soil	Semiconductor	$\sim 10^{-3} - 10^{-2}$
Fresh water	Semiconductor	$\sim 10^{-2}$
Human and animal tissue	Semiconductor	$\sim 0.2 - 0.7$
Intrinsic germanium	Semiconductor	~ 2.227
Seawater	Semiconductor	~ 4
Tellurium	Conductor	$\sim 5 \times 10^{-2}$
Carbon	Conductor	$\sim 3 \times 10^{-4}$
Graphite	Conductor	$\sim 3 \times 10^4$
Cast iron	Conductor	$\sim 10^6$
Mercury	Conductor	10^6
Nichrome	Conductor	10^6
Silicon steel	Conductor	2×10^6
German silver	Conductor	2×10^6
Lead	Conductor	5×10^6
Tin	Conductor	9×10^6
Iron	Conductor	1.03×10^7
Nickel	Conductor	1.45×10^7
Zinc	Conductor	1.7×10^7
Tungsten	Conductor	1.83×10^7
Brass	Conductor	2.56×10^7
Aluminum	Conductor	3.96×10^7
Gold	Conductor	4.1×10^7
Copper	Conductor	5.76×10^7
Silver	Conductor	6.1×10^7

*For most semiconductors the conductivities are representative for a frequency of about 10 GHz.

Skip sections 2.5 Semiconductors,

2.6 Superconductors, & 2.7 Metamaterials

2.8 Linear, Homogenous, Isotropic, and Nondispersive Media

Linear - $\epsilon, \mu, \text{ \& } \sigma$ are NOT functions of applied electric or magnetic field strength

examples of nonlinearity

→ dielectric breakdown $\Rightarrow |E|$ where electrons ripped loose from material

→ magnetic saturation $\Rightarrow |H|$ where all magnetic dipoles are aligned

Homogenous - $\epsilon, \mu, \text{ \& } \sigma$ are NOT functions of position/location w/in material; otherwise material is inhomogeneous or nonhomogenous

→ this can be a concern for circuit board manufacturers as well as for TL insulation.

Nondispersive - $\epsilon, \mu, \text{ \& } \sigma$ are NOT functions of frequency (or time)

→ Reality is that all materials are dispersive. However, over limited BW they may be effectively nondispersive.

2.8 cont.

Isotropic - $\epsilon, \mu, \text{ \& } \sigma$ are NOT functions of direction w/in the material

→ Many crystalline materials are anisotropic or nonisotropic. In this case, $\epsilon, \mu, \text{ \& } \sigma$ are usually written as 3×3 tensors instead of a single value. E.g.,

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

$$\begin{bmatrix} D_x \\ D_y \\ D_z \end{bmatrix} = \begin{bmatrix} \epsilon_{xx} & \epsilon_{xy} & \epsilon_{xz} \\ \epsilon_{yx} & \epsilon_{yy} & \epsilon_{yz} \\ \epsilon_{zx} & \epsilon_{zy} & \epsilon_{zz} \end{bmatrix} \begin{bmatrix} E_x \\ E_y \\ E_z \end{bmatrix}$$

⇓

$$D_x = \epsilon_{xx} E_x + \epsilon_{xy} E_y + \epsilon_{xz} E_z$$

$$D_y = \epsilon_{yx} E_x + \epsilon_{yy} E_y + \epsilon_{yz} E_z$$

$$D_z = \epsilon_{zx} E_x + \epsilon_{zy} E_y + \epsilon_{zz} E_z$$

→ Each of the terms in the $\vec{\epsilon}$ tensor can be complex

→ $\epsilon_{xx}, \epsilon_{yy}, \text{ \& } \epsilon_{zz}$ are called principal permittivities