

Magnetic Materials

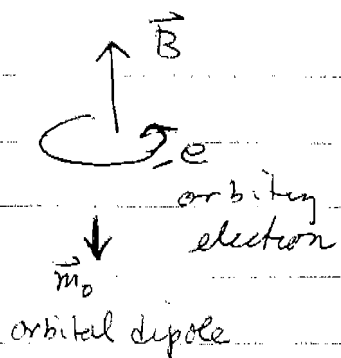
Circulating currents on atomic scale give rise to local magnetic dipole moments, which create local magnetic fields in the material.

Sources of circulating atomic currents:

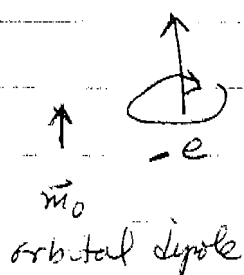
- 1) intrinsic angular momentum of electrons, i.e. "electron spin" - can add up and give a net angular momentum to atom
- 2) orbital angular momentum of electrons - can add up to give net angular momentum of atom.

(1) + (2) $\Rightarrow$ atoms can have a net magnetic dipole moment. When $\vec{B} = 0$, these atomic moments are generally in random orientations ^{and average to zero} (exception is a ferromagnet where moments can align even if $\vec{B} = 0$)
When apply $\vec{B} \neq 0$, the moments tend to align parallel to $\vec{B}$ giving a net magnetization density $\vec{M} \propto \vec{B}$. This is a paramagnetic effect.

But there is also a diamagnetic effect from orbital angular momentum (exists even if total angular momentum of electrons is zero, i.e. exists for atoms with zero net dipole moment)



← applying $\vec{B}$ to orbiting electron speeds up its orbital velocity. Increased angular momentum of negatively charged electron gives change in dipole moment $\Delta \vec{m} \propto -\vec{B}$



← applying $\vec{B}$ to orbiting electron slows down its orbital velocity. Net result is again that $\Delta \vec{m} \propto -\vec{B}$

see Griffiths
chpt 6 + prob
7.17 2nd ed
for details

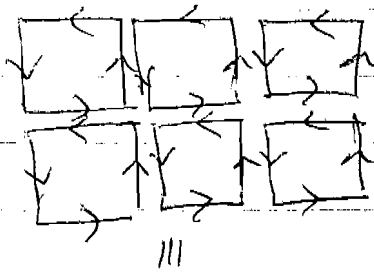
No matter which way electron orbits with respect to $\vec{B}$, result is a decrease in magnetic moment, so $\Delta \vec{m} \propto -\vec{B}$.
That $\Delta \vec{m}$ is opposite to $\vec{B}$ is called diamagnetism

Model atomic magnetic moments as small current loops. When loops get oriented, i.e. there is non zero average magnetization density

$$\vec{M}(\vec{r}) = \sum_i \vec{m}_i \delta(\vec{r} - \vec{r}_i)$$

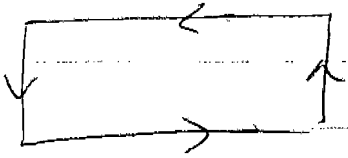
then net effect is to have a current flowing around the system. This current gives rise to magnetic fields

aligned atomic moments in a uniform applied $\vec{B}$



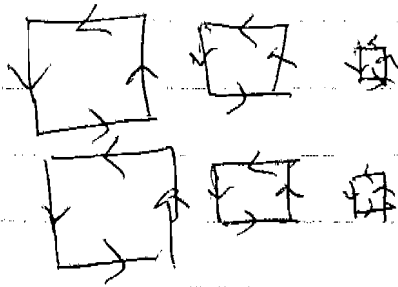
in interior, currents in opposite directions cancel also $\vec{j} = 0$ inside

①
 $\vec{B}$ out of page



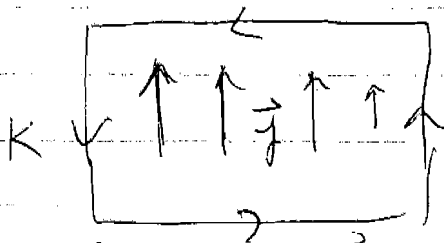
but is net circulation of current around boundary of material
 $\Rightarrow$ surface current $\vec{K}_{\text{bound}}$

If $\vec{B}$ is not uniform, then $\vec{M}$ is not uniform
Can create finite current density $\vec{j}$ in interior, as well as surface currents



Now currents in interior do not cancel, Net current $\vec{j}_{\text{bound}}$ in interior

$\vec{B}$ strong $\vec{B}$ weak



$\vec{M}$ strong

$\vec{M}$ weak

$\vec{B}$ out of page $\Rightarrow \vec{M}$ out of page
 ~~$\vec{M}$ varies along page~~
 $\vec{M}$ varies in direction $\perp$ direction of $\vec{B}$
 $\Rightarrow \vec{\nabla} \times \vec{M} \neq 0$ gives $\vec{j}_{\text{bound}}$

Average current

$$\langle \vec{j}_0 \rangle = \underbrace{\left\langle \sum_{i \in \text{free}} q_i \vec{v}_i \delta(\vec{r} - \vec{r}_i) \right\rangle}_{\substack{\uparrow \\ \text{current from free} \\ \text{charges}}} + \sum_n \underbrace{\langle \vec{j}_n \rangle}_{\substack{\uparrow \\ \text{current from} \\ \text{molecule } n \text{ of} \\ \text{the dielectric}}}$$

$$\begin{aligned} \langle \vec{j}_n(\vec{r}, t) \rangle &= \sum_{i \in n} q_i (\vec{v}_n + \vec{v}_{ni}) \langle \delta(\vec{r} - \vec{r}_n(t) - \vec{r}_{ni}(t)) \rangle \\ &= \sum_{i \in n} q_i \underbrace{(\vec{v}_n + \vec{v}_{ni})}_{\substack{\uparrow \quad \uparrow \\ \vec{v}_n = \frac{d\vec{r}_n}{dt} \quad \vec{v}_{ni} = \frac{d\vec{r}_{ni}}{dt}}} f(\underbrace{\vec{r} - \vec{r}_n(t) - \vec{r}_{ni}(t)}_{\substack{\uparrow \\ \text{position of} \\ \text{charge } i \\ \text{wrt CM}}}) \end{aligned}$$

as with $\langle \rho_0 \rangle$, we can expand in $\vec{r}_{ni}$

$$\begin{aligned} \langle \vec{j}_n \rangle &= \sum_{i \in n} q_i (\vec{v}_n + \vec{v}_{ni}) \left\{ f(\vec{r} - \vec{r}_n) - \vec{r}_{ni} \cdot \vec{\nabla} f(\vec{r} - \vec{r}_n) \right. \\ &\quad \left. + \frac{1}{2} \sum_{\alpha\beta} (r_{ni})_\alpha (r_{ni})_\beta \frac{\partial^2 f(\vec{r} - \vec{r}_n)}{\partial r_\alpha \partial r_\beta} + \dots \right\} \end{aligned}$$

we will keep only the first two terms in the expansion

The various terms we have to consider are

$$\textcircled{1} \quad \sum_{i \in n} q_i \vec{r}_n \delta(\vec{r} - \vec{r}_n)$$

$$\textcircled{2} \quad \sum_{i \in n} q_i \vec{r}_{ni} \delta(\vec{r} - \vec{r}_n)$$

$$\textcircled{3} \quad - \sum_{i \in n} q_i \vec{r}_n [\vec{r}_{ni} \cdot \vec{\nabla} f(\vec{r} - \vec{r}_n)]$$

$$\textcircled{4} \quad - \sum_{i \in n} q_i \vec{r}_{ni} [\vec{r}_{ni} \cdot \vec{\nabla} f(\vec{r} - \vec{r}_n)]$$

$$\textcircled{1} = \vec{r}_n f(\vec{r} - \vec{r}_n) \sum_{i \in n} q_i = q_n \vec{r}_n f(\vec{r} - \vec{r}_n) \\ = \langle q_n \vec{r}_n \delta(\vec{r} - \vec{r}_n) \rangle$$

this is just current of molecule as if it were a point charge q_n . For a neutral molecule $q_n = 0$ and this term vanishes.

$$\textcircled{2} \quad \text{Note: } \frac{\partial}{\partial t} \langle \vec{p}_n \delta(\vec{r} - \vec{r}_n) \rangle = \frac{\partial}{\partial t} \left(\sum_{i \in n} q_i \vec{r}_{ni} f(\vec{r} - \vec{r}_n) \right) \\ = \sum_{i \in n} q_i \vec{v}_{ni} f(\vec{r} - \vec{r}_n) \\ + \sum_{i \in n} q_i \vec{r}_{ni} [-\vec{\nabla} f(\vec{r} - \vec{r}_n) \cdot \vec{v}_n]$$

$$\text{So for } \textcircled{2}, \quad \sum_{i \in n} q_i \vec{v}_{ni} f(\vec{r} - \vec{r}_n)$$

$$= \frac{\partial}{\partial t} \langle \vec{p}_n \delta(\vec{r} - \vec{r}_n) \rangle$$

$$+ [\vec{v}_n \cdot \vec{\nabla} f(\vec{r} - \vec{r}_n)] \vec{p}_n$$

So

$$\textcircled{2} = \sum_{i \in n} q_i \vec{r}_{ni} f(\vec{r} - \vec{r}_n) = \frac{\partial}{\partial t} \langle \vec{p}_n \delta(\vec{r} - \vec{r}_n) \rangle + (\vec{v}_n \cdot \vec{\nabla}) \langle \vec{p}_n \delta(\vec{r} - \vec{r}_n) \rangle$$

2nd term is $\sum_{\alpha} v_{n\alpha} \frac{\partial}{\partial r_{\alpha}} \langle \vec{p}_n \delta(\vec{r} - \vec{r}_n) \rangle$

$$\textcircled{3} = -\vec{v}_n \left(\sum_{i \in n} q_i \vec{r}_{ni} \right) \cdot \vec{\nabla} f(\vec{r} - \vec{r}_n) = -\vec{v}_n (\vec{p}_n \cdot \vec{\nabla} f(\vec{r} - \vec{r}_n))$$

$$= -\vec{v}_n \cdot \vec{\nabla} \langle \vec{p}_n \delta(\vec{r} - \vec{r}_n) \rangle = \sum_{\alpha} \vec{v}_n \frac{\partial}{\partial r_{\alpha}} \langle p_{n\alpha} \delta(\vec{r} - \vec{r}_n) \rangle$$

$$\textcircled{4} = -\vec{\nabla} f(\vec{r} - \vec{r}_n) \cdot \sum_{i \in n} q_i \vec{r}_{ni} \vec{v}_{ni}$$

We have seen the tensor $\sum_{i \in n} q_i \vec{r}_{ni} \vec{v}_{ni}$ before when we considered the magnetic dipole moment

$$\sum_{i \in n} q_i \vec{r}_{ni} \vec{v}_{ni} = \int d^3r \vec{r} \vec{j} \quad \text{where } \vec{j}(\vec{r}) \equiv \sum_{i \in n} q_i \vec{v}_{ni} \delta(\vec{r} - \vec{r}_{ni})$$

is current density with respect to center of mass of molecule

We had $\int d^3r \vec{r} \vec{j} = -\int d^3r \vec{j} \vec{r} - \int d^3r (\vec{\nabla} \cdot \vec{j}) \vec{r} \vec{r}$

↑
in statics, $\vec{\nabla} \cdot \vec{j} = 0$

in general $\vec{\nabla} \cdot \vec{j} = -\frac{\partial \rho}{\partial t}$

$$\int d^3r \vec{r} \vec{j} = -\int d^3r \vec{j} \vec{r} + \int d^3r \frac{\partial \rho}{\partial t} \vec{r} \vec{r}$$

$$= -\int d^3r \vec{j} \vec{r} + \frac{\partial}{\partial t} \left[\int d^3r \rho \vec{r} \vec{r} \right]$$

↑
although this is not zero,

it is a quadrupole term of the same order as the term we dropped when we truncated expansion to linear order.

$$\sim O\left(\frac{a_0}{L}\right)^2$$

$$S_0 \quad \int d^3r \quad \vec{r} \vec{f} \approx - \int d^3r \vec{f} \vec{r} \quad \text{ignoring the quadrupole term}$$

$$= \frac{1}{2} \int d^3r \left[\vec{r} \vec{f} - \vec{f} \vec{r} \right]$$

$$\sum_{i \in n} g_i \vec{r}_{ni} \vec{v}_{ni} = \frac{1}{2} \sum_{i \in n} g_i \left[\vec{r}_{ni} \vec{v}_{ni} - \vec{v}_{ni} \vec{r}_{ni} \right]$$

$$- \vec{\nabla} f(\vec{r} - \vec{r}_n) \cdot \sum_{i \in n} g_i \vec{r}_{ni} \vec{v}_{ni} = - \vec{\nabla} f(\vec{r} - \vec{r}_n) \cdot \frac{1}{2} \sum_{i \in n} g_i \left[\vec{r}_{ni} \vec{v}_{ni} - \vec{v}_{ni} \vec{r}_{ni} \right]$$

$$= -\frac{1}{2} \sum_{i \in n} g_i \left[(\vec{\nabla} f \cdot \vec{r}_{ni}) \vec{v}_{ni} - (\vec{\nabla} f \cdot \vec{v}_{ni}) \vec{r}_{ni} \right]$$

$$= -\frac{1}{2} \sum_{i \in n} g_i \vec{\nabla} f \times (\vec{v}_{ni} \times \vec{r}_{ni}) \quad \text{triple product rule}$$

$$= \vec{\nabla} f(\vec{r} - \vec{r}_n) \times \frac{1}{2} \sum_{i \in n} \vec{r}_{ni} \times \vec{v}_{ni} g_i$$

$$= \vec{\nabla} f(\vec{r} - \vec{r}_n) \times \frac{1}{2} \int d^3r \vec{r} \times \vec{f}$$

$$= \vec{\nabla} f(\vec{r} - \vec{r}_n) \times c \vec{m}_n \quad \text{where} \quad \vec{m}_n = \frac{1}{2c} \sum_{i \in n} \vec{r}_{ni} \times \vec{v}_{ni} g_i$$

is magnetic dipole moment of molecule n

$$= \vec{\nabla} \times \int d^3r f(\vec{r} - \vec{r}_n) c \vec{m}_n$$

$$= \vec{\nabla} \times \langle c \vec{m}_n \delta(\vec{r} - \vec{r}_n) \rangle$$

Adding all the pieces

$$\begin{aligned}
 \langle \vec{f}_n \rangle &= \underbrace{\langle g_n \vec{v}_n \delta(\vec{r} - \vec{r}_n) \rangle}_{(1)} + c \underbrace{\vec{\nabla} \times \langle \vec{m}_n \delta(\vec{r} - \vec{r}_n) \rangle}_{(4)} \\
 &+ \underbrace{\frac{\partial}{\partial t} \langle \vec{p}_n \delta(\vec{r} - \vec{r}_n) \rangle}_{(2)} + \underbrace{(\vec{v}_n \cdot \vec{\nabla}) \langle \vec{p}_n \delta(\vec{r} - \vec{r}_n) \rangle}_{(2)} \\
 &\quad - \underbrace{\vec{v}_n \cdot \vec{\nabla} \langle \vec{p}_n \delta(\vec{r} - \vec{r}_n) \rangle}_{(3)}
 \end{aligned}$$

Define $\vec{M}(\vec{r}) \equiv \sum_n \langle \vec{m}_n \delta(\vec{r} - \vec{r}_n) \rangle$ average magnetization density

$\vec{P}(\vec{r}) \equiv \sum_n \langle \vec{p}_n \delta(\vec{r} - \vec{r}_n) \rangle$ polarization density, as before

$$\begin{aligned}
 \sum_n \langle \vec{f}_n \rangle &= \sum_n \langle g_n \vec{v}_n \delta(\vec{r} - \vec{r}_n) \rangle + c \vec{\nabla} \times \vec{M} + \frac{\partial \vec{P}}{\partial t} \\
 &+ \sum_n \left[(\vec{v}_n \cdot \vec{\nabla}) \langle \vec{p}_n \delta(\vec{r} - \vec{r}_n) \rangle - \vec{v}_n \cdot \vec{\nabla} \langle \vec{p}_n \delta(\vec{r} - \vec{r}_n) \rangle \right]
 \end{aligned}$$

see Jackson (6.96) for additional electric quadrupole terms

The last term on the right hand side is usually small and ignored. This is because the molecular velocities $\vec{v}_n$ are usually small, and randomly oriented, so that they average to zero. (see Jackson (6.100) for case of net translation of dielectric, $\vec{v}_n = \text{const}$ all n)

Define macroscopic current density

$$\vec{j}(\vec{r}, t) = \left\langle \sum_{i \in \text{free}} q_i \vec{v}_i \delta(\vec{r} - \vec{r}_i) \right\rangle + \left\langle \sum_n q_n \vec{v}_n \delta(\vec{r} - \vec{r}_n) \right\rangle$$

$\uparrow$ $\uparrow$
 current of free charges current of molecular drifting
if molecules are charged

Then $\langle \vec{j}_0 \rangle = \vec{j} + c \vec{\nabla} \times \vec{M} + \frac{\partial \vec{P}}{\partial t}$

Ampere's Law becomes upon averaging

$$\vec{\nabla} \times \vec{B} = \frac{4\pi}{c} \langle \vec{j}_0 \rangle + \frac{1}{c} \frac{\partial \vec{E}}{\partial t}$$

$$= \frac{4\pi}{c} \vec{j} + 4\pi \vec{\nabla} \times \vec{M} + \frac{4\pi}{c} \frac{\partial \vec{P}}{\partial t} + \frac{1}{c} \frac{\partial \vec{E}}{\partial t}$$

$$\vec{\nabla} \times (\vec{B} - 4\pi \vec{M}) = \frac{4\pi}{c} \vec{j} + \frac{1}{c} \frac{\partial}{\partial t} (\vec{E} + 4\pi \vec{P})$$

define $\boxed{\vec{H} \equiv \vec{B} - 4\pi \vec{M}}$ to get

$$\boxed{\vec{\nabla} \times \vec{H} = \frac{4\pi}{c} \vec{j} + \frac{1}{c} \frac{\partial \vec{D}}{\partial t}}$$

$$\boxed{\vec{D} = \vec{E} + 4\pi \vec{P}} \text{ as before}$$

official nomenclature: $\vec{B}$ is the magnetic induction

$\vec{H}$ is the magnetic field

common usage: both ~~H~~ and $\vec{B}$ are called magnetic field