

Magnetic Materials

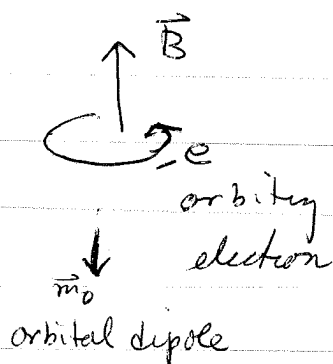
Circulating currents on atomic scale give rise to local magnetic dipole moments, which create local magnetic fields in the material.

Sources of circulating atomic currents:

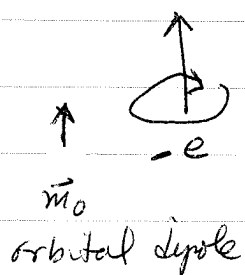
- 1) intrinsic angular momentum of electrons, i.e. "electron spin" - can add up and give a net angular momentum to atom
- 2) orbital angular momentum of electrons - can add up to give net angular momentum of atom.

(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

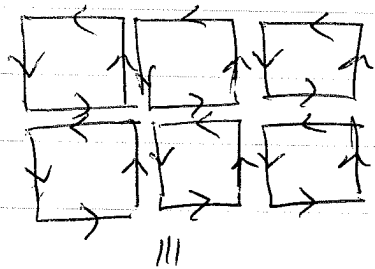
No matter which way electron orbits with respect to $\vec{B}$, result is a decrease in magnetic moment, so $\Delta \vec{m} \propto -\vec{B}$. That $\Delta \vec{m}$ is opposite to $\vec{B}$ is called diamagnetism.

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

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

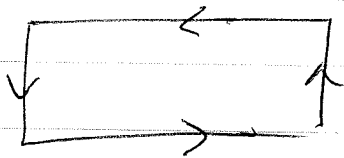
then net effect is to have a current flowing around the system. This current gives rise to magnetic fields

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



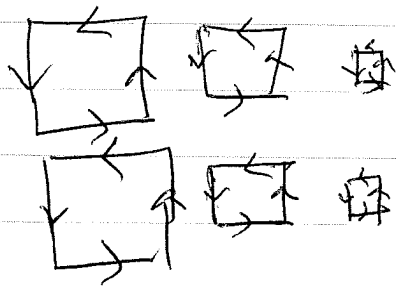
in interior, currents in opposite directions cancel also $\vec{j} = 0$ inside

①
 $\vec{B}$ out of page



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

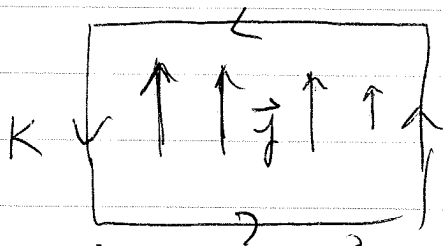
If $\vec{B}$ is not uniform, then $\vec{M}$ is not uniform
Can create finite current density $\vec{j}$ in interior, as well as surface currents



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

$\vec{B}$ strong

$\vec{B}$ weak



$\vec{M}_{\text{strong}}$

$\vec{M}_{\text{weak}}$

$\vec{B}$ 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

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

$$\begin{aligned} \langle \vec{j}_n(\vec{r}, t) \rangle &= \sum_{i \in n} q_i (\vec{v}_n + \vec{v}_{ni}) \langle \delta(\vec{r} - \vec{r}_n(t) - \vec{r}_{ni}(t)) \rangle \\ &= \sum_{i \in n} q_i (\vec{v}_n + \vec{v}_{ni}) f(\vec{r} - \vec{r}_n(t) - \vec{r}_{ni}(t)) \end{aligned}$$

$\vec{v}_n = \frac{d\vec{r}_n}{dt}$ $\vec{v}_{ni} = \frac{d\vec{r}_{ni}}{dt}$ position of CM of molec n position of charge i wrt CM

as with $\langle j_0 \rangle$, 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

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

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

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

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

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

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

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

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

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

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

S_0

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

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

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

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

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

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

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

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

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

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

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

$\uparrow$

although this is not zero,

it is a quadrupole term of the same order as the terms we dropped when we truncated

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

$$So \quad \int d^3r \quad \vec{r} \vec{j} \simeq - \int d^3r \quad \vec{j} \vec{r} \quad \text{ignoring the quadrupole term}$$

$$= \frac{1}{2} \int d^3r \quad [\vec{r} \vec{j} - \vec{j} \vec{r}]$$

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

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

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

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

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

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

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

is magnetic dipole moment of molecule n

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

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

Adding all the pieces

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

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

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

$\uparrow$ current of free charges $\uparrow$ 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

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

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

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

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

when atoms have intrinsic magnetic moments due to electron spin, we can add these to $\vec{M}$ in obvious way

when molecules are neutral, $g_n = 0$, the "bound current" is given by

$$\vec{j}_{\text{bound}} = \sum_n \langle \vec{j}_n \rangle = c \vec{\nabla} \times \vec{M} + \frac{\partial \vec{P}}{\partial t}$$

Note that the $\frac{\partial \vec{P}}{\partial t}$ term is crucial to give conservation of bound charge

$$\begin{aligned} \vec{\nabla} \cdot \vec{j}_{\text{bound}} &= c \vec{\nabla} \cdot (\vec{\nabla} \times \vec{M}) + \vec{\nabla} \cdot \frac{\partial \vec{P}}{\partial t} \\ &= 0 + \frac{\partial}{\partial t} (\vec{\nabla} \cdot \vec{P}) \end{aligned}$$

$$= -\frac{\partial \rho_{\text{bound}}}{\partial t} \quad \text{where } \rho_{\text{bound}} = -\vec{\nabla} \cdot \vec{P} \text{ is bound charge density}$$

$$\text{so } \boxed{\vec{\nabla} \cdot \vec{j}_{\text{bound}} + \frac{\partial \rho_{\text{bound}}}{\partial t} = 0}$$

ad bound charge is conserved.

Since total average charge must be conserved, i.e.

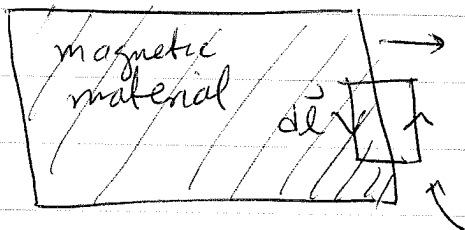
$$\vec{\nabla} \cdot \langle \vec{j}_0 \rangle - \frac{\partial \langle \rho_0 \rangle}{\partial t} = 0, \quad \text{and } \langle \vec{j}_0 \rangle = \underbrace{\vec{j}}_{\text{free current}} + \vec{j}_{\text{bound}}$$

$$\langle \rho_0 \rangle = \underbrace{\rho}_{\text{free charge}} + \rho_{\text{bound}}$$

$$\Rightarrow \boxed{\vec{\nabla} \cdot \vec{j} + \frac{\partial \rho}{\partial t} = 0}$$

free charge is also conserved

At a surface of a magnetic material



$\hat{m}$ outward normal to surface

take $\hat{z} = d\vec{l} \times \hat{m}$ out of page

Amperian loop C bounding surface of area da

$$\begin{aligned} \oint_C da \hat{z} \cdot (\vec{\nabla} \times \vec{M}) &= \int_S da \hat{z} \cdot \vec{J}_{\text{bound}} = da \hat{z} \cdot \vec{J}_{\text{bound}} \\ &= (d\vec{l} \times \hat{m}) \cdot \vec{K}_{\text{bound}} \quad \text{as width of loop} \\ &\quad \rightarrow 0 \\ &= (\hat{m} \times \vec{K}_{\text{bound}}) \cdot d\vec{l} \end{aligned}$$

But by Stokes theorem

$$\oint_S da \hat{z} \cdot (\vec{\nabla} \times \vec{M}) = \oint_C d\vec{l} \cdot \vec{M} = c d\vec{l} \cdot \vec{M} \quad \text{since width} \rightarrow 0$$

and $\vec{M} = 0$ outside

$$\Rightarrow c d\vec{l} \cdot \vec{M} = (\hat{m} \times \vec{K}_{\text{bound}}) \cdot d\vec{l} \quad \text{for any } d\vec{l} \text{ in plane of surface}$$

$$\Rightarrow c \vec{M}_t = \hat{m} \times \vec{K}_{\text{bound}}$$

where $\vec{M}_t$ is component of $\vec{M}$ tangential to the surface (since $\vec{K}_b$ is in plane of surface, $\hat{m} \times \vec{K}$ is also entirely in the plane of the surface)

$$\begin{aligned} \Rightarrow c \hat{m} \times \vec{M}_t &= c \hat{m} \times \vec{M} = \hat{m} \times (\hat{m} \times \vec{K}_{\text{bound}}) \\ &= -\vec{K}_{\text{bound}} \end{aligned}$$

$$\Rightarrow \boxed{\begin{aligned} \vec{K}_{\text{bound}} &= c \vec{M} \times \hat{m} \\ \vec{H}_{\text{bound}} &= c \vec{\nabla} \times \vec{M} \end{aligned}}$$

Total bound charge vanishes

$$Q_{\text{bound}} = \int_V d^3r \rho_{\text{bound}} + \int_S da \sigma_{\text{bound}}$$

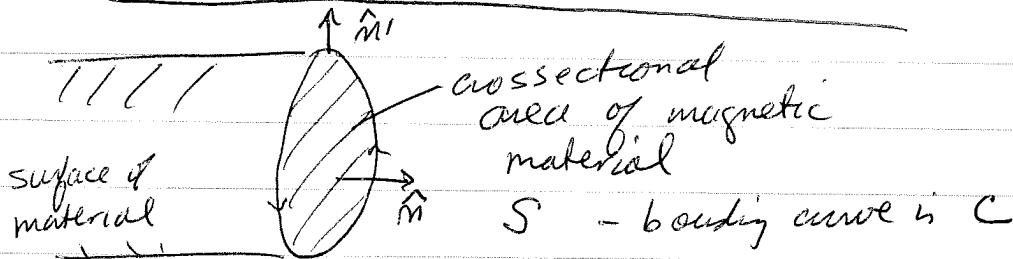
$\uparrow$ vol of dielectric $\leftarrow$ surface of dielectric

$$= \int_V d^3r -\vec{\nabla} \cdot \vec{P} + \int_S da \hat{n} \cdot \vec{P}$$

but by Gauss theorem $\int_V d^3r \vec{\nabla} \cdot \vec{P} = \int_S da \hat{n} \cdot \vec{P}$

so $Q_{\text{bound}} = - \int_S da \hat{n} \cdot \vec{P} + \int_S da \hat{n} \cdot \vec{P} = 0$

Total bound current vanishes



$\hat{n}$ is normal to cross-section
 $\hat{n}'$ is normal to surface

total current flowing through S is

$$\int_S da \hat{n} \cdot \vec{j}_{\text{bound}} + \int_C d\vec{l} \vec{K}_{\text{bound}} \cdot \hat{n}$$

$$= c \int_S da \hat{n} \cdot (\vec{\nabla} \times \vec{M}) + c \int_C d\vec{l} \hat{n} \cdot (\vec{M} \times \hat{n}')$$

$$= c \int_C d\vec{l} \cdot \vec{M} + c \int_C d\vec{l} (\hat{n}' \times \hat{n}) \cdot \vec{M}$$

$= -\hat{x}$ unit tangent, $d\vec{l} = dl \hat{x}$

$$= c \int_C d\vec{l} \cdot \vec{M} - c \int_C d\vec{l} \cdot \vec{M} = 0$$