Lecture 2: Principles of Magnetic Sensing

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- I. Basic Concepts in Magnetism
- 2. Sensor Principles



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2.1 Basic concepts in Magnetism

Magnetic field sources are

- distributions of electric current (including moving charged particles)
- time-varying electric fields
- permanently magnetized material



Magnetic fields

Two sources of *H* - currents



Biot-Savart Law

$$\delta \mathbf{H} = -rac{1}{4\pi}rac{\mathbf{r} imes \mathbf{j}}{|\mathbf{r}^3|} \delta V$$

$$\delta \boldsymbol{H} = -\frac{1}{4\pi} I \frac{\boldsymbol{r} \times \delta \ell}{|\boldsymbol{r}^3|}$$



Right-hand corkscrew

Scalar product

m.r

 ∇ .B

 $m_x x + m_y y + m_z z$

 $\partial B_x / \partial x + \partial B_y / \partial y + \partial B_z / \partial z$

The 'divergence' of **B**

$$\nabla = (\partial | \partial \mathbf{x}, \partial | \partial \mathbf{y}, \partial | \partial \mathbf{z})$$



 $(yj_z - zj_y)\mathbf{e}_x - (xj_z - zj_x)\mathbf{e}_y + (xj_y - yj_x)\mathbf{e}_z \qquad (\partial B_z/\partial y - \partial B_y/\partial z)\mathbf{e}_x - (\partial B_z/\partial x - \partial B_x/\partial z)\mathbf{e}_y + (\partial B_z/\partial y - \partial B_y/\partial z)\mathbf{e}_x - (\partial B_z/\partial x - \partial B_y/\partial z)\mathbf{e}_y + (\partial B_z/\partial y - \partial B_y/\partial z)\mathbf{e}_x - (\partial B_z/\partial x - \partial B_y/\partial z)\mathbf{e}_y + (\partial B_z/\partial y - \partial B_y/\partial z)\mathbf{e}_x - (\partial B_z/\partial x - \partial B_y/\partial z)\mathbf{e}_y + (\partial B_z/\partial y - \partial B_y/\partial z)\mathbf{e}_x - (\partial B_z/\partial x - \partial B_y/\partial z)\mathbf{e}_y + (\partial B_z/\partial y - \partial B_y/\partial z)\mathbf{e}_x - (\partial B_z/\partial x - \partial B_y/\partial z)\mathbf{e}_y + (\partial B_z/\partial y - \partial B_y/\partial z)\mathbf{e$ + $(\partial B_y / \partial x - \partial B_x / \partial y) \mathbf{e}_z$

In a steady state (no time-dependent electric field)

 $\nabla \times H = j$

∮H.dI = I

 $\int (\nabla \times \mathbf{H}) d\mathcal{A} = \int \mathbf{j} d\mathcal{A} \quad \text{Stokes theorem} \rightarrow$

 $H = I/2\pi r$

The field at a distance 5cm m from a wire carrying a current of 1 A is \sim 3 A m⁻¹



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In free space \mathbf{B} = \mu_0 \mathbf{H}
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Unit of B - Tesla Unit of μ_0 T/Am⁻¹ μ_0 = 4 π 10⁻⁷ T/Am⁻¹



 $I T = I0^{7}/4\pi \approx 800,000 \text{ Am}^{-1}$

The field at a distance 5cm m from a wire carrying a current of 1 A is $4 \mu T$

 $B = \mu_0 I/2\pi r$



Two sources of *H* - magnets

The magnetic moment *m* is the elementary quantity in solid state magnetism.

Define a local moment density - magnetization - M(r,t) which fluctuates wildly on a sub-nanometer and a sub-nanosecond scale.

Define a mesoscopic average magnetization M, averaging over a few nm and ~ 1µs

$$\delta m = M \delta V$$

The continuous medium approximation

M can be the spontaneous magnetization M_s within a ferromagnetic domain

A macroscopic average magnetization is the domain average

 $\boldsymbol{M} = \sum_{i} \boldsymbol{M}_{i} \boldsymbol{V}_{i} / \sum_{i} \boldsymbol{V}_{i}$



The mesoscopic average magnetization

The magnetic moment

Ampère: A magnetic moment *m* is equivalent to a current loop.



Provided the current flows in a plane

Magnetization

The local moment density **M** is the magnetization

Units: A m⁻¹

e.g. for iron M = 1710 kA m^{-1;} for BaFe₁₂O₁₉ M = 380 kA m⁻¹

e.g. for a 2.5 cc BaFe₁₂O₁₉ fridge magnet (M = 380 kA m⁻¹, $V = 2.5 \ 10^{-6} \ m^3$), $m \approx 1 \ A \ m^2$

Magnetization *M* can be induced by an applied field <u>or</u> it can arise spontaneously within a ferromagnetic domain, M_s . A macroscopic average magnetization is the domain average

The equivalent Amperian current density is

$$\boldsymbol{j}_{\boldsymbol{M}} = \nabla \times \boldsymbol{M}$$



The Magnetic Periodic Table



Susceptibilities of the elements



B, H and M

The equation used to *define* **H** is **B** = μ_0 (**H** + **M**) i.e. **H** = **B**/ μ_0 - **M**

We call the *H*-field due to a magnet; — *stray field* outside the magnet

— demagnetizing field, H_d , inside the magnet



The total *H*-field at any point is $H = H' + H_m$ where H' is the applied field

The **B** field - magnetic induction/magnetic flux density

 $\nabla . \boldsymbol{B} = 0$ Significance; It is the *fundamental* magnetic field There are no sources or sinks of **B** i.e no monopoles

Magnetic vector potential



The gradient of any scalar ϕ , $\nabla \phi$ may be added to **A** without altering **B**

Gauss's theorem: The net flux of B across any closed surface is zero

Magnetic flux $d\Phi = B.dA$ Units: Weber (Wb)

Flux quantum Φ_0 = 2.07 10¹⁵ Wb



The equation $\nabla \mathbf{x} \mathbf{B} = \mu_0 \mathbf{j}$ valid in static conditions gives:

Ampere's law $\int \mathbf{B} \cdot \mathbf{dI} = \mu_0 \mathbf{I}$ for any closed path

Good for calculating the field for very symmetric current paths.

Example: the field at a distance r from a current-carrying wire $B = \mu_0 I/2\pi r$



B interacts with any moving charge:

Lorentz force $f = q(E + v \times B)$

The **H** field

Significance; The magnetization of a solid reflects the local value of H.

In free space $\boldsymbol{B} = \mu_0 \boldsymbol{H}$; $\nabla \mathbf{x} \boldsymbol{B} = \mu_0 \boldsymbol{j}_c$

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In a medium \boldsymbol{B} = \mu_0 \mu_r \boldsymbol{H} (linear isotropic media only!)
\boldsymbol{B} = \mu_0 (\boldsymbol{H} + \boldsymbol{M}) (general case)
\nabla \times \boldsymbol{B} = \mu_0 (\boldsymbol{j}_c + \boldsymbol{j}_M),
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but $\boldsymbol{j}_M = \nabla \times \boldsymbol{M}$



The **H** field

Significance; The magnetization of a solid reflects the local value of H.

In free space $\boldsymbol{B} = \mu_0 \boldsymbol{H}$.

$$\nabla \mathbf{X} \mathbf{B} = \mu_0 (\mathbf{j}_c + \mathbf{j}_M)$$
, $\mathbf{j}_M = \nabla \mathbf{X} \mathbf{M}$ hence $\nabla \mathbf{X} \mathbf{H} = \mathbf{j}_c$ $\int \mathbf{H} \cdot d\mathbf{I} = \mathbf{I}_c$

Coulomb approach to calculate H

H has sources and sinks associated with nonuniform magnetization $\mathbf{k} \mathbf{e}_{n}$

$$\nabla . H = - \nabla . M$$

Imagine **H** due to a distribution of magnetic charges q_{m} (Am)

Field of a point 'charge'

$$H = q_{\rm m} e_r / 4\pi r^2$$

Magnetization distribution is replaced by

- surface charge distribution $\sigma_m = \mathbf{M}.\mathbf{e}_n$
- volume charge distribution ρ_m = ∇ .**M**



Ampere's law for H

Magnetic scalar potential

When H is due only to magnets i.e $\nabla \mathbf{x} \mathbf{H} = 0$ we can define a scalar potential φ_m (Units are Amps) Such that

$$H = -\nabla \varphi_{\rm m}$$
 The potential of charge $q_{\rm m}$ is $\varphi_{\rm m} = q_{\rm m}/4\pi r$

If currents are present, this cannot be done.

Poisson's equation $\nabla^2 \varphi_m = \nabla$. **M**

2.4 Boundary conditions

Gauss's law $\int_{S} \mathbf{B} \cdot d\mathbf{A} = 0$

gives that the perpendicular component of **B** is continuous.

Medium 1

 \boldsymbol{H}_{11}

B_

It follows from from Ampère's law

 $\int_{\text{loop}} \boldsymbol{H} \cdot d\boldsymbol{I} = I_{\text{c}} = 0$

(there are no conduction currents on the surface) that the parallel component of H is continuous. $(H_1-H_2) \times e_n=0$

Conditions on the potentials

Medium 2

Since $\int_{S} \mathbf{B} \cdot d\mathbf{A} = \int_{loop} \mathbf{A} \cdot d\mathbf{I}$ (Stoke's theorem)

 $({\bf A}_1 - {\bf A}_2) \times {\bf e}_n = 0$

The scalar potential is continuous

$$\varphi_{m1}$$
 = φ_{m2}

Hysteresis



The hysteresis loop shows the irreversible, nonlinear response of a ferromagnet to a magnetic field M = M(H). It reflects the arrangement of the magnetization in ferromagnetic *domains*. The magnet cannot be in thermodynamic equilibrium anywhere around the open part of the curve! It reflects the arrangement of the magnetization in ferromagnetic *domains*. The B = B(H) loop is deduced from the relation $B = \mu_0(H + M)$.

Maxwell's equations in a material medium



 $\nabla \cdot \mathbf{B} = 0$ $\nabla \cdot \mathbf{D} = \rho$ $\nabla \times \mathbf{H} = \mathbf{j}_c + \partial \mathbf{D} / \partial t$ $\nabla \times \mathbf{E} = -\partial \mathbf{B} / \partial t$

Written in terms of the four fields, these are valid in any medium. In vacuum $D = \varepsilon_0 E$, $H = B/\mu_0$, ρ is charge density (C m⁻³), \mathbf{j}_c is conduction current density (A m⁻²)

In magnetostatics there is no time-dependence of **B**, **D** or ρ Conservation of charge $\nabla . \mathbf{j} = -\partial \rho / \partial t$. In a steady state $\partial \rho / \partial t = 0$ Magnetostatics: $\nabla . \mathbf{j} = 0$; $\nabla . \mathbf{B} = 0$ $\nabla \times \mathbf{H} = \mathbf{j}_c$ Constituent relations: $\mathbf{j}_c = \mathbf{j}(\mathbf{E})$; $\mathbf{P} = \mathbf{P}(\mathbf{E})$; $\mathbf{M} = \mathbf{M}(\mathbf{H})$

Magnetic materials

Ferromagnets and ferrimagnets have spontaneous magnetization within a domain.

The magnetization falls with increasing temperature, first gradually, then abruptly at the Curie point $T_{\rm C}$.



Iron Fe

bcc; $a_0 = 287 \text{ pm}$

The most important ferromagnetic material.

Main constituent of the whole Earth, 5 wt % of crust.

Usually alloyed with 6 at% Si and fabricated in

300 µm rolled laminations (isotropic or grain

oriented), castings or reduced powder,

Mainly used in electrical machines (motors, transformers) and magnetic circuits.

Production 5 Mt/yr for magnetic purposes (8 B€)

 $J_{s} = 2.0 \text{ T} \text{ (Si-Fe)}$

$$M_{s} = 1.71 \text{ MA m}^{-1} \text{ (Fe)}$$
$$T_{C} = 1044 \text{ K} \text{ (Fe)}$$
$$K_{1} = 48 \text{ kJ m}^{-3} \text{ (Fe)}$$
$$\lambda_{s} = -8 10^{-6}$$



Permalloy Fe₂₀Ni₈₀

324 pm

Multipurpose soft magnetic material, with near-zero anisotropy *and* magnetostriction Sometimes alloyed with Mo, Cu ...

Sputtered or electrodeposited films, sheet, powder.

Uses: magnetic recording; write heads, read heads (AMR), magnetic shields, transformer cores

$$J_s = 1.0 T$$
 $M_s = 0.8 MA m^{-1}$
 $K_1 \approx 2 kJ m^{-3}$ $\lambda_s = 2 10^{-6}$
 $T_C = 843 K$

Compositions near Fe50Ni50 have larger J_s but greater anisotropy

fcc; $a_0 =$



Cobalt Co

hcp; a = 251 pm, c = 407 pm

Highest-T_C ferromagnet, anisotropic, expensive (¤50 /kg), strategic. Useful alloying addition Sputtered nanocrystalline thin films (with Cr, Pt, B additions) are used as media for hard discs

$$J_s = 1.8 T$$
 $M_s = 1.44 MA m^{-1}$
 $K_1 = 530 kJ m^{-3}$
 $T_C = 1360 K$



Most common magnetic mineral, source of rock magnetism, main constituent of lodestones..

A ferrimagnet. with Fe²⁺ and Fe³⁺ disordered on B -site above the Verwey transition at $T_v = 120$ K, ordered below; A-B superexchange is the main magnetic interaction

$$\begin{bmatrix} Fe^{3+} \end{bmatrix}_{tett} \{Fe^{2+} Fe^{3+} \}_{oct} O_4 \\ \downarrow \uparrow \uparrow \\ -5 \mu_B + 4 \mu_B + 5 \mu_B = 4 \mu_B \end{bmatrix}$$

A half-metal. Fe(B); \downarrow electrons hop in a t_{2g} band Used as toner, and in ferrofluids. Potential for spin electronics..

 $J_{s} = 0.60 \text{ T} \qquad m_{0} = 4.0 \ \mu_{B} \ / \ \text{fu}$ $K_{1} = -13 \ \text{kJ} \ \text{m}^{-3} \qquad \lambda_{s} = 40 \ 10^{-6}$ $T_{C} = 843 \ \text{K}$

spinel;
$$a_0 = 839 \text{ pm}$$



BaFe₁₂O₁₉; Hexaferrite

magnetoplumbite; a = 589 pm c = 2319 pm

An hcp lattice of oxygen and Ba, with iron in octahedral (12k, $4f_{2}$, 2a) tetrahedral (4f₁) and trigonal bipyramidal (2b) sites.

Brown ferrimagnetic insulator. All magnetic ions are Fe^{3+} . Also $SrFe_{12}O_{19}$ and La/Co substitution.

Structure is $12k\uparrow 2a\uparrow 2b\uparrow 4f_1\downarrow 4f_2\downarrow$

T_C = 740 K.

. . .

 $m_{
m 0}$ = 20 $\mu_{
m B}$ / fu

Low-cost permanent magnet, the first magnet to break the 'shape barrier'. 98% of all permanent magnets by mass are Ba or Sr ferrite. Found on every fridge door and in innumerable catches, dc motors, microwave magnetrons,

80g manufactured per year for everyone on earth

 $J_s = 0.48 \text{ T}$. $K_1 = 450 \text{ kJ m}^{-3}$ $B_a = 1.7 \text{ T}$



Samarium-cobalt SmCo₅

Versatile, high-temperature permanent magnet. Cellular intergrowth with Sm_2Co_{17} in $Sm(Co, Fe, Zr, Cu)_{7.6}$ alloys provides domain-wall pinning Dense sinterered oriented material. Uses: specialised electrical drives Expensive ($\approx 150 \text{ m/kg}$)

 $J_r = 1.0 T$ (BH)_{max} = 200 kJ/m³ $K_1 = 17 MJ m^{-3}$ $B_a = 30 T$ $T_C = 1020 K$

R-T exchange is direct, between the 5d and 3d shells

This is antiferromagnetic; on-site coupling of 5d and 4f spins is ferromagnetic, hence moments couple parallel for light rare earths (J = L - S) and antiparallel for heavy rare earths (J = L + S).

Useful alloys are of Pr, Nd, Sm with Fe, Co, Ni

hexagonal; a = 499 pm c = 398 pm



Neomax, Nd₂Fe₁₄B

tetragonal;
$$a = 879 \text{ pm}, c = 1218 \text{ pm}$$

The highest-performance permanent magnet.Discovered in 1983 by Sagawa (sintered) and by Croat and Herbst (melt spun)Dy, Co .. substitutions

- Dense sinterered oriented material, melt-spun isotropic flakes.
- Voice-coil actuators, spindle motors, nmr imaging, flux sources

Cost \approx 30 \in /kg, Production 50 kT/yr (1.5 B \in)

 $J_r = 1.4 \text{ T} (BH)_{max} = 200-400 \text{ kJ/m}^3$ $K_1 = 4.9 \text{ MJ m}^{-3} B_a = 7.7 \text{ T}$ $T_C = 878 \text{ K}$



Exchange interactions.

The interaction responsible for magnetic order is exchange. Basically it is a Coulomb interaction between the charges of electrons on adjacent ions 1, 2, subject to the symmetry constraints of quantum mechanics. It written in terms of their spins.

Heisenberg-Dirac Hamiltonian $\mathcal{H} = -2 J \mathbf{S}_1 \cdot \mathbf{S}_2$



Curie or Néel temperature $T_c \approx 2Z J S(S+I)/3k_B$

Exchange interactions.

The magnetic coupling in a ferromagnet can be represented by a 'magnetic stiffness' A

$$E_{ex} = \int A(\nabla \mathbf{e}_M)^2 d^3 r \qquad \mathbf{e}_M = M(\mathbf{r})/M_s \qquad (\theta, \phi)$$

$$E_{ex} = \int A[(\nabla \theta)^2 + \sin^2 \theta (\nabla \phi)^2] d^3 r \qquad (\nabla \mathbf{e}_{Mx})^2 + (\nabla \mathbf{e}_{My})^2 + (\nabla \mathbf{e}_{Mz})^2$$

$$l_{ex} = \sqrt{\frac{A}{\mu_0 M_s^2}} \qquad \text{Exchange length} \qquad A = kT_c/2a$$

$$A = 2JS^2 Z_c/a_0$$

$$A \sim 10 \text{ pJ m}^{-1}$$

$$l_{ex} \sim 2 - 3 \text{ nm}$$

Demagnetizing field

The H-field in a magnet depends M(r) and on the shape of the magnet. H_d is uniform only in the case of a uniformly-magnetized ellipsoid.

> $(\boldsymbol{H}_{d})_{i} = -\mathcal{N}_{ij}\boldsymbol{M}_{j}$ i,j = x,y,z $\mathcal{N}_{x} + \mathcal{N}_{y} + \mathcal{N}_{z} = 1$

Demagnetizing factors for some simple shapes





New icon for permanent magnets! \Rightarrow





Measuring magnetization with no need for demagnetization correction

Apply a field in a direction where $\mathcal{N} = 0$

$$\boldsymbol{H} = \boldsymbol{H}' + \boldsymbol{H}_m \qquad (\boldsymbol{H}_d)_i = -\mathcal{N}_{ij}\boldsymbol{M}_j$$


It is not possible to have a uniformly magnetized cube



When measuring the magnetization of a sample **H** is the independent variable, M = M(H).

Response to an applied field *H'*

Susceptibility of linear, isotropic and homogeneous (LIH) materials

 $M = \chi' H'$ χ' is external susceptibility $M = \chi H$ χ is internal susceptibility

It follows that from $H = H' + H_{d}$, dividing by M, that

$$1/\chi = 1/\chi$$
' - \mathcal{N}

Typical paramagnets and diamagnets:

$$\chi \approx \chi'$$
 (10⁻⁵ to 10⁻³)

Paramagnets close to the Curie point and ferromagnets:





In LIH meida $B = \mu H$

 $\mu = B/H$ Units: TA⁻¹m

Relative permeability

$$\mu_r = \mu/\mu_0$$

 $B = \mu_0 (H + M)$ gives $\mu_r = 1 + \chi$

 μ_0 is the permeability of free space.

• In practice it is often easier to measure the mass of a sample than its volume. Measured magnetization is usually $\sigma = M/\rho$, the magnetic moment per unit mass (ρ is the density).

• Likewise the mass susceptibility is defined as $\chi_m = \chi/\rho$

• Molar susceptibility is $\chi_{mol} = \mathcal{M} \chi_m / 1000$ \mathcal{M} is the molecular weight (g/mole)

Examples.

Susceptibility of representative materials							
Susceptibility	Units	H_2O	Al	$\rm CuSO_4{\cdot}5H_2O$	$Gd_2(SO_4)_3 \cdot 8H_2O$		
X X m X mol	$m^3 kg^{-1}$ $m^3 mol^{-1}$	$-9.0 imes 10^{-6} \ -9.0 imes 10^{-9} \ -1.62 imes 10^{-10}$	$\begin{array}{l} 2.1 \times 10^{-5} \\ 7.9 \times 10^{-9} \\ 2.1 \times 10^{-10} \end{array}$	1.41×10^{-4} 6.2×10^{-8} 1.57×10^{-8}	2.6×10^{-3} 8.7×10^{-7} 6.5×10^{-7}		

Susceptibilities of the elements



Soft and hard magnets.

The area of the hysteresis loop represents the energy loss per cycle. For efficient soft magnetic materials, this needs to be as small as possible.



Energy product.



Magnetostatic energy and forces

Energy of ferromagnetic bodies

• Magnetostatic (dipole-dipole) forces are long-ranged, but weak. They determine the magnetic microstructure.

 $M \approx 1 \text{ MA m}^{-1}$, $\mu_0 H_d \approx 1 \text{ T}$, hence $\mu_0 H_d M \approx 10^6 \text{ J m}^{-3}$

Products *B.H*, *B.M*, $\mu_0 H^2$, $\mu_0 M^2$ are all energies per unit volume.

- Magnetic forces *do no work* on moving charges $f = q(\mathbf{v} \times \mathbf{B})$
- No potential energy associated with the magnetic force.



Reciprocity theorem

The interaction of a pair of dipoles, ε_p , can be considered as the energy of m_1 in the field B_{21} created by m_2 at r_1 or vice versa.

$$\varepsilon_{p} = -m_{1}.B_{21} = -m_{2}.B_{12}$$
So $\varepsilon_{p} = -(1/2)(m_{1}.B_{21} + m_{2}.B_{12})$

$$m_{1}$$

$$B_{21}$$

Extending to magnetization distributions:

$$\varepsilon = -\mu_0 \int \mathbf{M}_1 \cdot \mathbf{H}_2 \, \mathrm{d}^3 r = -\mu_0 \int \mathbf{M}_2 \cdot \mathbf{H}_1 \, \mathrm{d}^3 r$$

Magnetic energy terms

Self energy of a magnet in its demagnetizing field $E_m = -(1/2)\int \mu_0 H_d M d^3 r$ $E_m = (1/2)\int \mu_0 H_d^2 d^3 r$

Self energy of a uniformly magnetized sample $E_m = (1/2)\mu_0 \mathcal{M}^2 V$

Energy associated with a magnetic field $E_m = (1/2) \int \mu_0 H^2 d^3 r$

Energy product of a permanent magnet

Aim to maximize energy associated with the field created around the magnet, from previous slide:

$$\frac{1}{2}\int \mu_0 H_d^2 d^3r = -\frac{1}{2}\int_V \mu_0 \boldsymbol{H}_d \boldsymbol{.} \boldsymbol{M} d^3r.$$

Can rewrite as:

$$\frac{1}{2} \int_{o} \mu_0 H_d^2 d^3 r = -\frac{1}{2} \int_{i} \mu_0 H_d^2 d^3 r - \frac{1}{2} \int_{i} \mu_0 \boldsymbol{M} \cdot \boldsymbol{H}_d d^3 r.$$

where we want to maximize the integral on the left. Since $\mathbf{B} = \mu_0 (\mathbf{H} + \mathbf{M})$,

Energy product: twice the energy stored in the stray field of the magnet is

$$-\mu_0 \int_i \boldsymbol{B} \cdot \boldsymbol{H}_d d^3 r$$
 (1) Optimum shape, $\mathcal{N} = 1/2$

Thermodynamics

First law: $dU = H_x dX + dQ$

dQ = TdS

 $E(H_X,S)$

Four thermodynamic potentials;

U(X,S)internal energy

enthalpy

F(X,T) = U - TSHelmholz free energ dF = HdX - SdT

 $G(H_x,T) = F - H_x X$ Gibbs free energy $dG = -XdH_{\times} - SdT$

Magnetic work is $H\delta B$ or $\mu_0 H'\delta M$

 $dF = \mu_0 H' dM - S dT$ $dG = -\mu_0 M dH' - S dT$ (U,Q,F,G are in units of Jm⁻³)



 $S = -(\partial G/\partial T)_{H'} \quad \mu_0 M = -(\partial G/\partial H')_{T'}$

Maxwell relations $(\partial S/\partial H')_{T'} = -\mu_0(\partial M/\partial T)_{H'}$ etc.

Magnetostatic Forces

Force density on a uniformly magnetized body at constant temperature

 $\boldsymbol{F}_m = \nabla(\boldsymbol{\mu}_0 \boldsymbol{H}'.\boldsymbol{M}) \qquad \quad \nabla(\boldsymbol{H}'.\boldsymbol{M}) = (\boldsymbol{H}'.\nabla)\boldsymbol{M} + (\boldsymbol{M}.\nabla)\boldsymbol{H}'$

Kelvin force

$$\boldsymbol{F}_m = \mu_0(\boldsymbol{M}.\nabla)\boldsymbol{H}'.$$

General expression, for when **M** is dependent on **H** is

$$oldsymbol{F}_m = -\mu_0
abla \left[\int_0^H \left(rac{\partial M \upsilon}{\partial \upsilon}
ight)_{H,T} dH
ight] + \mu_0 (oldsymbol{M}.
abla) oldsymbol{H}.$$

v = 1/d d is the density

Anisotropy.

shape, magnetocrystalline, induced, strain

The ferromagnetic axis lies in some particular direction determined by shape or some intrinsic anisotropy related to crystal or atomic structure.

 $E_{\rm a} = K_{\rm I} \sin^2\theta + K_{\rm 2} \sin^4\theta + \dots$

The shape contribution is derived from the energy expression $E_m = (1/2)\mu_0 \mathcal{M} P^2 V$

The magnetization lies along the direction for which $\mathcal N$ is smallest - the axis of a long bar.

The magnetocrystalline contribution for uniaxial crystals is given by a similar expression with different K_1

Shape anisotropy.

The shape contribution is derived from the energy expression $E_m = (1/2)\mu_0 \mathcal{M} P^2 V$

 \mathcal{N} is the demagnetizing factor for the easy direction - the axis of a long bar. (1/2)[1 - \mathcal{N}] is the demagnetizing factor for the perpendicular direction (assuming an ellipsoid)

Hence
$$\Delta E_{\rm m} = (1/2) \ \mu_0 M^2 V\{(1/2)(1 - \mathcal{N}) - \mathcal{N}\}$$

so $K_{\rm sh} = (1/4) \ \mu_0 M^2 (1 - 3\mathcal{N})$

The biggest it can be is $(1/4)\mu_0 M^2 J m^{-3}$

~ 2 10⁵ J m⁻³ for
$$\mu_0 M = 1 T$$

Magnetocrystalline anisotropy

The magnetocrystalline contribution is ultimately caused bu spin-orbit coupling which connects the crystal structure (electron orbits) and magnetic moment direction.

Hexagonal: $E_a = K_1 \sin^2 \theta + K_2 \sin^4 \theta + K_3 \sin^6 \theta + K'_3 \sin^6 \theta \sin 6\phi$, Tetragonal: $E_a = K_1 \sin^2 \theta + K_2 \sin^4 \theta + K'_2 \sin^4 \theta \cos 4\phi + K_3 \sin^6 \theta + K'_3 \sin^6 \theta \sin 4\phi$, Cubic: $E_a = K_{1c} (\alpha_1^2 \alpha_2^2 + \alpha_2^2 \alpha_3^2 + \alpha_3^2 \alpha_1^2) + K_{2c} (\alpha_1^2 \alpha_2^2 \alpha_3^2)$,





Anisotropy due to texture

Directional order of atomic constituents in a binary alloy can be induces by deposition on a magnetic field or by post depositional annealing.





Domains

Micromagnetic energy, wall width and structure

Domains form to minimize the dipolar energy E_d

$$E_d = -rac{1}{2}\int \mu_0 oldsymbol{H}_d \cdot oldsymbol{M} d^3r$$



Micromagnetic energy

$$E_{tot} = E_{ex} + E_K + E_d + E_m + E_{stress} + E_{ms}$$



Minimizing E_{tot} gives the mesoscale magnetic structure of the sample (monodomain, multidomaindomains or vortex)

Domain walls



Minimizing E_{tot} for two oppositely magnetized regions gives the domain wall width $\delta_w = \pi \sqrt{(A/K_I)}$

A ~ 10 pJ m⁻¹; $K \sim 10^5$ jm⁻³; $\delta_w \sim 30$ nm.



Néel relaxation $\tau = \tau_0 \exp(\Delta/kT)$

Here τ_0 is an inverse attempt frequency, $10^{-9}s^{-1}$

If $\Delta/kT = 25$, $\tau = 70$ s.

2.2 Sensor Principles

- Flux Faraday's law
- MR Lorentz force
- Hall effect Lorentz force
- AMR Spin-orbit scattering
- GMR spin accumulation
- TMR spin-dependent tunelling
- MO Faraday effect
- SQUID Flux quantization
- NMR magnetic resonance
- GMI high-frequency permeability

Sensor	Principle	Detects	Frequency	Field (T)	Noise	Comments
Coil	Faraday's law	dΦ/dt	10 ⁻³ - 10 ⁹	10 ⁻¹⁰ - 10 ²	100 nT	bulky ,absolute
Fluxgate	saturation	Н	dc - 10 ³	10 ⁻¹⁰ - 10 ⁻³	10 _P T	bulky
Hall probe	Lorentz f ce	В	dc - 10 ⁵	10 ⁻⁵ - 10	100 nT	thin film
MR	Lorentz fce	B ²	dc - 10 ⁵	10 ⁻² - 10	10 nT	thin film
AMR	spin-orbit int	Н	dc - 10 ⁷	10 ⁻⁹ - 10 ⁻³	10 nT	thin film
GMR	spin accum.n	Н	dc - 10 ⁹	10 ⁻⁹ - 10 ⁻³	10 nT	thin film
TMR	tunelling	н	dc - 10 ⁹	10 ⁻⁹ - 10 ⁻³	l nT	thin film
GMI	permability	Н	dc - 10 ⁴	10 ⁻⁹ - 10 ⁻²		wire
MO	Kerr/Faraday	Μ	dc - 10 ⁵	10 ⁻⁹ - 10 ²	Ι _Ρ Τ	bulky
SQUID It	flux quanta	Φ	dc - 10 ⁹	10 ⁻¹⁵ - 10 ⁻²	l fT	cryogenic
SQUID ht	flux quanta	Φ	dc - 10 ⁴	10 ⁻¹⁵ - 10 ⁻²	30 fT	cryogenic
NMR	resonance	В	dc - 10 ³	10 ⁻¹⁰ - 10	l nT	Very precise

2.2.1 Inductive sensors

Inductive sensors detect an emf in a coil proportional to the rate of change of flux, according to Faraday's law:

 $\mathcal{E} = -d\Phi/dt$

They provided an absolute measurement of $B = \Phi/n\mathcal{A}$, as in a search coil with an integrating voltmeter, or the rotating coil gaussmeter



Inductive read/write heads were widely used until 1990 in magnetic recording



Magnetic circuits

$\nabla \mathbf{B} = \mathbf{0}$

Assuming no flux leakage

 $B_{\rm m} \mathcal{A}_{\rm m} = B_{\rm g} \mathcal{A}_{\rm g}$ (1)

Assuming ideal soft material µ = ∞ Ampere's law ∫H.dI = 0

 $H_{\rm m}/_{\rm m} = -H_{\rm g}/_{\rm g} \qquad (2)$

Multiplying $B_m H_m V_m = - B_g^2 V_g / \mu_0$

Dividing $-B_m/H_m = \mu_0 A_g I_m / A_m I_g$ The *permeance* coefficient



Soft iron $\phi = \phi_m - \phi_g$ Airgap \mathcal{A}_{g} $|R_g|$ R_m φ_m Φ. (b) Ø R_{g} Magnet

Figure A simple magnetic circuit, and its electrical equivalent, with and without losses.



Table	Analogy betwee	n electric and m	nagnetic circuits
		Electric	Magnetic
Field		$E (V m^{-1})$	H (A m ⁻¹)
Potential		φ_e (V)	φ_m (A)
Current/flux density		j (A m ⁻²)	B (T or Wb m ^{-2})
Potential difference		$\varphi_e = \int E \cdot \mathrm{d}l$	$\varphi_m = \int \boldsymbol{H} \cdot \mathrm{d}\boldsymbol{l}$
Continuity condition		$\nabla \cdot \mathbf{j} = 0$	$\nabla \cdot \boldsymbol{B} = 0$
Linear response law		$j = \sigma E$	$B = \mu H$
Current/flux		<i>I</i> (A)	Φ (T m ² or Wb)
Resistance/reluctance	e	$R = \varphi_e / I \ (\Omega)$	$R_m = \varphi_m / \Phi (A W b^{-1})$
for a cylinder of sect	ion ${\cal A}$ and length l	$R = l/A\sigma$	$R_m = l/\mathcal{A}\mu$
Conductance/permeance		G = 1/R	$P_m = 1/R_m$

2.2.2 Fluxgates

Fluxgates depend on the nonlinear saturation of the magnetization of a soft magnetic core. Two identical cores (or a single toroidal core) have oppositely wound ac field windings. A parallel applied field leads to saturation of one of the cores, producing an ac signal linear in H.





Fluxgates are bulky but sensitive, reliable and impervious to radiation. Used, for example, in space.

2.2.3 Hall sensors

Effect discovered by Edwin Hall in 1879



Hall voltages linear in field are produced in semiconductor plates, especially Si and in 2-deg GaAs/GaAlAs structures. These are fourterminal devices, and the current source and high-gain amplifier are often integrated on a chip.

Used for secondary field measurements – each probe must be calibrated — and as proximity sensors. About a billion are produced each year.

2.2.4 Classical magnetoresistance

The simplest Lorentz force device is a semicoductor or semimetal which exhibits classical positive B^2 magnetoresiatance. High-mobility semiconductors such as InAs and InSb show large effects (~ 100 % T⁻¹). Field is applied perpendicular to the semiconductor slab, and it is possible to achieve a desired resistance by patterning a series of metallic contacts.



The sensors are nonlinear, two-terminal devices providing a good response in large fields. They are used as position sensors in brushless dc permanent magnet motors.

2.2.5 Anisotropic magnetoresistance (AMR)



Discovered by W. Thompson in 1857

 $ρ = ρ_0 + Δρcos^2 θ$

Magnitude of the effect $\Delta \rho / \rho < 3\%$ The effect is usually positive; $\rho_{||} > \rho_{\perp}$

AMR is due to spin-orbit s-d scattering

High field sensitivity is achieved in thin films of soft ferromagnetic films such as permalloy ($Fe_{20}Ni_{80}$).

Planar Hall effect

Planar Hall effect is a variant of AMR; $\rho_{\parallel} \neq \rho_{\perp} = \rho_{\parallel}$ is when $j \parallel M \dots$



$$E_{\parallel} = \rho_{\parallel} j_x \cos\theta \qquad E_{\perp} = \rho_{\perp} j_x \sin\theta$$

Components of electric field parallel and perpendicular to the current are $E_x = E_{||}\cos\theta + E_{\perp}\sin\theta$, $E_y = E_{||}\sin\theta - E_{\perp}\cos\theta$ $E_x = j(E_{||}\rho_{\perp} + \Delta\rho\cos^2\theta)$ $E_y = j\Delta\rho\sin\theta\cos\theta$

Hence

 $V_{pH} = j w \Delta \rho \sin \theta \cos \theta$

The biggest effect is when θ changes from 45 to 135 degrees.

2.2.6 Giant magnetoresistance.



Peter Grunberg and Albert Fert;



Discovery of GMR 1988 Implementation in hard disk drives 1998 Nobel Prize 2007

10⁹ GMR sensors per year






2.2.7 Tunnel magnetoresistance.



2.2.8 Giant magnetoimpedance.

GMI sensors are soft ferromagnetic wires (sometimes permalloy-plated copper wires) or films. An ac current is passed along the wire, and L is measured as a function of applied field. At high frequency, the skin depth < wire diameter. Permeability depends on f and H.



Very high field sensitivity, 10⁴ % mT⁻¹ is achievable Used in Wii games, three-axis compasses 2.2.9 Magneto-optic sensors.

Optical fibre field sensors, based on magneto-optic Faraday effect. They are bulky, and used for large fields.

Rotation sensors based on Sagnac effect.

2.2.10 Superconducting quantum interference devices.

SQUIDs detect the change of flux threading a flux-locked loop. The flux is generally coupled to the SQUID via a superconducting flux transformer. The device is sensitive to a small fraction of a flux quantum. SQUIDSs offer ultimate field sensitivity. They generally operate with a flux-locked loop.



2.2.11 Nuclear magnetic resonance.



Torque creates precession at the Larmor frequency $f_L = \gamma B/2\pi$

Protons (in water for example) can be polarized by a field pulse, and then allowed to precess freely (fid) at the Larmor frequency in the field to be measured. f_L in the Earth's field is ~ 2 kHz.

Rb or Cs vapor can be magnetized by optical pumping with circularly-polarized light, and the nuclear precession measured. The Rb-vapour magnetometer provides an extremely precise, absolute value of the magnitude of the field.

These magnetometers have been packaged on a chip.