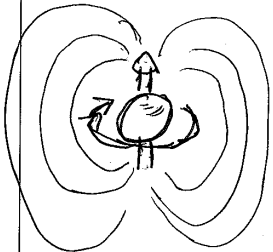


CLASSICAL DESCRIPTION OF MR

→ ODD # PROTONS / NEUTRONS have Nuclear spin
ANGULAR MOMENTUM



INTRINSIC SPIN PROPERTY!
MORE CH. 4

INTRINSIC MAGNETIC MOMENT

→ SPINNING MAGNETIC DIPOLES

→ BIOLOGICAL TISSUE MOSTLY ^1H in H_2O
SOMETIMES ^{31}P , ^{13}C , ^{23}Na exotic

→ MR IS ABOUT INTERACTIONS WITH
THREE FIELDS:

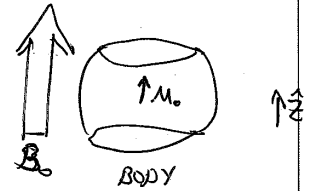
B_0 - MAIN FIELD
⇒ POLARIZATION

B_1 - RF FIELD
⇒ SIGNAL PRODUCTION / RECEPTION

\vec{G} - GRADIENT FIELDS
⇒ SPATIAL ENCODING

 B_0 - MAIN FIELD

→ PRODUCES POLARIZATION
OF SAMPLE M_0



→ SPINS EXHIBIT RESONANCE
AT LARMOR FREQUENCY

$$\omega = -\gamma B_0$$

↑ Gyromagnetic ratio

γ DEPENDS ON NUCLEUS
FOR PROTONS!

$$\frac{\gamma}{2\pi} = 4.257 \text{ MHz/G}$$

$$\gamma = 2\pi 4.257 \text{ rad/G}$$

WORTH REMEMBERING!

OTHERS: ^{23}Na $\frac{\gamma}{2\pi} = 1.127 \text{ MHz/G}$

^{13}C $\frac{\gamma}{2\pi} = 1.071 \text{ MHz/G}$

^{15}N $\frac{\gamma}{2\pi} = -0.43 \text{ MHz/G}$ Negative!

TYPICAL B₀'S

0.1 T	4.2 MHz	VERY LOW!
0.5 T	21 MHz	LOW (Permanent Magnet)
1 T	42 MHz	MILD (superconducting)
1.5 T	63 MHz	"HIGH" Diagnostic
3 T	126 MHz	"HIGH" fMRI Neuro
4 T	170 MHz	RARE (Decommissioned @ Berkeley)
7/9.4 T		VERY HIGH Research ONLY!

FOR SPECTRAL/SPATIAL LOCALIZATION

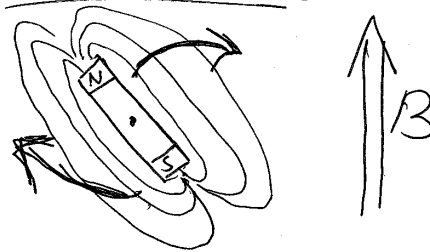
WE REQUIRE HOMOGENEITY

$\Delta B_0 \sim 1 \text{ PPM}$ 40 cm³ FOV

64 Hz @ 1.5 T

PRETTY REMARKABLE!

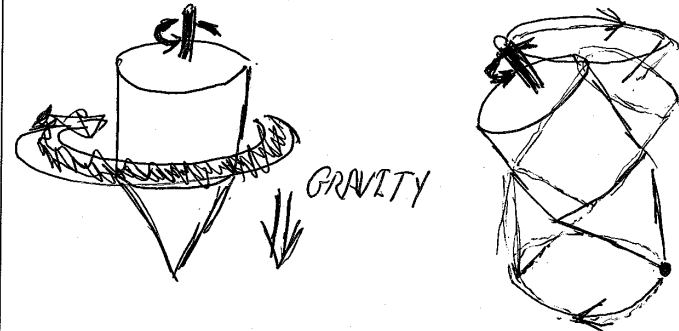
WHY RESONANCE?



→ TORQUE, BUT NO RESONANCE.

→ MISSING BAR MAGNET IS MISSING ANGULAR MOMENTUM.

LIKE A SPINNING TOP!



B_1 - RF FIELD

CANT DIRECTLY DETECT M_z .

WHY? HUGE FIELD!

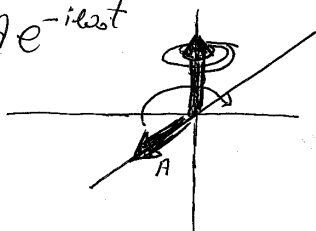
RESONANCE IS THE BIG DEAL...

B_0 IS DC SPIN RESONANCE \Rightarrow DETECTION!

\rightarrow SAMPLE RESONATES AT $\omega = \gamma B_0$.

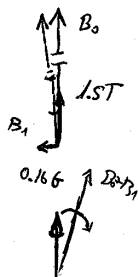
\rightarrow APPLY ROTATING RF FIELD AT $\omega = \gamma B_0$ IN THE TRANSVERSE PLANE

$B_1(t) = A e^{-i\omega t}$



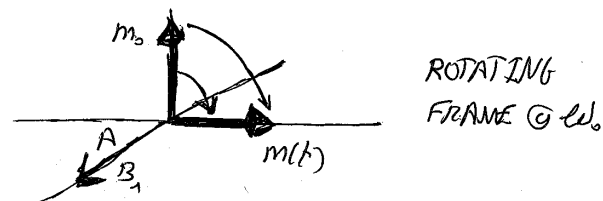
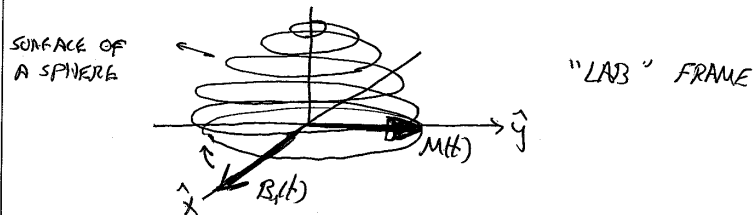
AT TIME $t=0$

AT TIME $t = \frac{1}{\omega}$



HAS TO BE ON RESONANCE TO DO SOMETHING

ILLUSTRATE BY ROTATING A STRING



\rightarrow IN ROTATING FRAME: PRECESSION ABOUT B_1

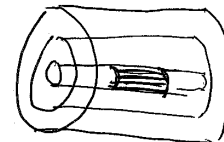
$\omega_e = \gamma B_1 = 4.257 \cdot 0.16 = 0.68 \text{ kHz}$
 $0.367 \text{ ms } 90^\circ \Rightarrow 23000 \text{ ROTATIONS}$

TYPICAL B_1 'S 0.14 - 0.35 G

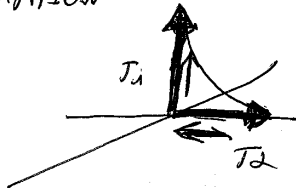
DURATION 1-3 MS LONG TIME AT 64 MHz

PEAK POWER μKW !

HEATING FROM THIS (SAR)



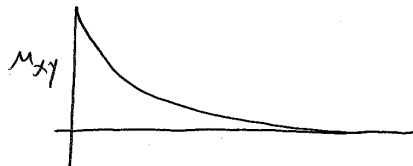
(-) MAGNETIZATION EXHIBITS RELAXATION:



T_1 ~ LONGITUDINAL $100 \rightarrow 2000$ ms In Tissue

T_2 ~ TRANSVERSE $10 \rightarrow 300$

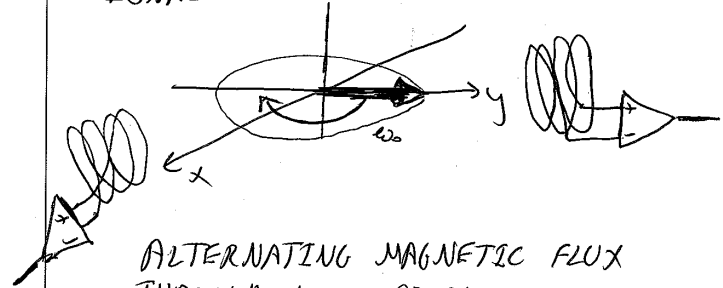
MAIN SOURCE OF TISSUE CONTRAST (MORE LATER)



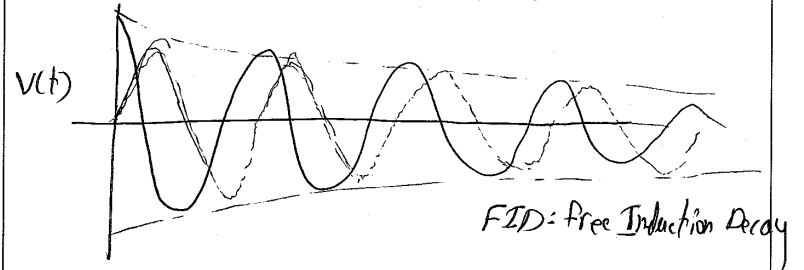
$T_2 < T_1$ ALWAYS

B₁ RECEPTION

ONCE EXCITED, WE CAN PICK UP SIGNAL



ALTERNATING MAGNETIC FLUX THROUGH A LOOP PRODUCES EMF (FARADAYS LAW $\mathcal{E} = \frac{d\phi_B}{dt}$)



THIS IS WHAT CHEMISTS USE

CHEMICAL SHIFT ~ 1 PPM
WANT B₀ FLATTER.

(-) Body \rightarrow TINY OSCILLATORS
MRI: IMAGE OSCILLATORS.

G- GRADIENT FIELDS

SPATIAL LOCALIZATION

B_1 HAS POOR LOCALIZATION $\lambda @ 64 \text{ MHz} \sim 5 \text{ cm in tissue}$

INSTEAD, CODE POSITION IN FREQUENCY

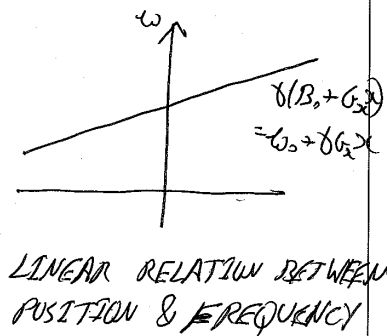
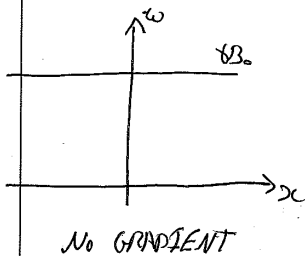
$$\omega(x) = \gamma(B_0 + G_x x)$$

\uparrow gradient in x

⊕ CHANGE IN Z COMPONENT W/ x

$$G_x = \frac{\partial B_z}{\partial x}$$

(B_x, B_y DO NOT MATTER MUCH
IN HIGH FIELD)



TYPICAL

$$\begin{aligned} G &= 1-10 \text{ G/cm} \\ &= 10-100 \text{ mT/m} \\ &= 4.2-42 \text{ kHz/cm} \end{aligned}$$

GRADIENT WAVEFORMS IN AUDIO RANGE

$$SR = 15 \text{ G/cm/ms}$$

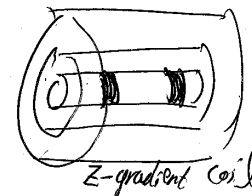
SAFETY CONCERN IS $\frac{dB}{dt}$
 $\left(\frac{dG_x(t)}{dt} \right) x$

PERIPHERAL NERVE STIMULATION

BIG AMPS: 1200 VOLTS
200 AMPS

GRADIENTS DO NOT
SATISFY MAXWELL EQN.

NOT AN ISSUE IN
HIGH FIELD



B₁ - RF FIELD ± 10% ACCURACY

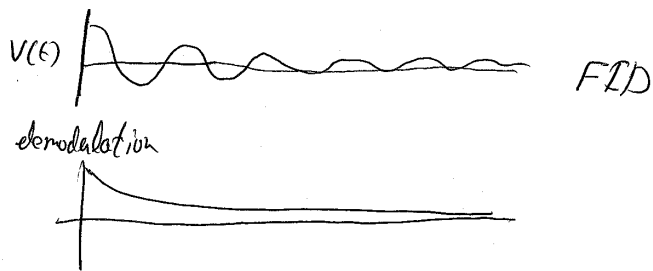
- MAGNETIZATION PRECESSES AROUND ROTATING FIELD AND IS TIPPED AWAY
- (-) VERY SMALL FIELD (MAX 0.35[G])
- (-) RESONANCE IS ESSENTIAL
- EASY TO DESCRIBE IN ROTATING FRAME.

RELAXATION

- (-) LONGITUDINAL T₁
- (-) TRANSVERSE T₂

RECEPTION

(-) EMF $\mathcal{E} = \frac{d\phi}{dt}$



(*) ±10% ACCURACY LEADS TO DIFFERENT FLIPS IN SPACE

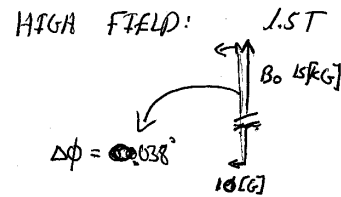


GRADIENT FIELDS

ENCODE POSITION ONTO FREQUENCY

$$G_x = \frac{\partial B}{\partial x}$$

(SMALL CONCOMITANT FIELDS B_x, B_y DO NOT CONTRIBUTE TO PRECESSION SINCE << B₀ AND NOT OSCILLATING



ASSUME $\frac{\partial \mathcal{E}}{\partial x}$

$$\|B_0 \hat{z} + B_x \hat{x}\| = 15000 \cdot 0.00033 \Rightarrow \Delta f = 14 \text{ Hz}$$

~~XXXXXXXXXX~~

FOR $G_x \sim 1 \frac{\text{G}}{\text{cm}} \Rightarrow 0.42 \frac{\text{kHz}}{\text{mm}} = 420 \frac{\text{Hz}}{\text{mm}}$

SHIFT OF 0.03 [mm]

FOR ACCURACY OF 1mm NEED

$$\frac{\Delta B_0}{B_0} \approx \frac{0.42}{63,000 \text{ kHz}} \sim 6.5 \text{ PPM} \quad \text{USUALLY 1PPM!}$$

WITH $G = \frac{46}{\text{cm}} \sim 26 \text{ PPM}$

GRADIENT FIELDS

$\pm 20x$

ENCODE POSITION ONTO FREQUENCY

$$G_x = \frac{\partial B_1}{\partial x}$$

SMALL CONCOMITANT FIELDS B_x, B_y DO NOT CONTRIBUTE TO PRESSION ABOUT B_0 - NOT OSCILLATING

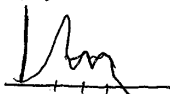
ANECDOTE

1973 Lauterbur

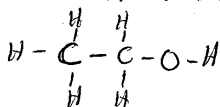
"Zeugmatography" - clever!

PAUL SOMEWHAT LAZY (AKA EFFICIENT) YOUNG PROFESSOR AT STONY BROOK CHEMISTRY. LOTS OF MEASUREMENTS!

MEASURED



H H H DIFFERENT e^- SHIELDING

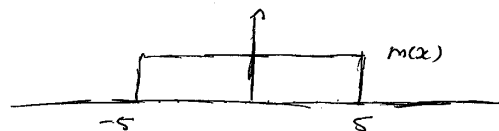


TO SPEED THINGS UP, HE PUT 2 TEST TUBES AND ADDED LINEAR GRADIENT



HE REALIZED THIS WAS IMAGING!

EXAMPLE SQUARE PULS



$$s(t) = \int_{-5}^{5} m(x) e^{-i(\omega_0 + \delta(G_x x)t)} dx =$$

$$= e^{-i\omega_0 t} \int_{-5}^{5} m(x) e^{-i\delta(G_x x)t} dx =$$

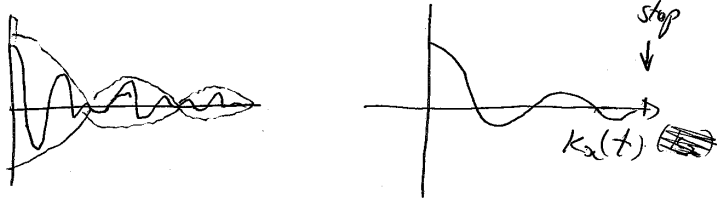
$$= e^{-i\omega_0 t} \int_{-5}^{5} m(x) e^{-i2\pi \left(\frac{\delta}{2\pi} G_x t\right) x} dx =$$

$$= e^{-i\omega_0 t} \mathcal{F}\{m(x)\} \Big|_{k_x = \frac{\delta}{2\pi} G_x t}$$

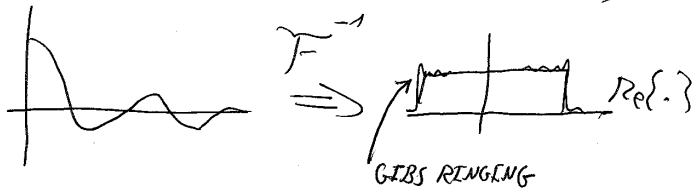
$$m(x) = \text{rect}(\cdot)$$

$$\mathcal{F}\{m(x)\} \sim \text{sinc}(\cdot)$$

$$s(t) = e^{-i\omega_c t} \underbrace{F\{m(x)\}}_{\text{BaseBand}} \Big|_{\omega_c = \frac{d}{2\pi} 60k} \leftarrow \text{CARRIER}$$



INVERSE FT OF BASEBAND SIGNAL
GIVES 1D PROJECTION (ALMOST..)



~~REASON~~ TO SPECIFIC ISSUES: COMPLEX VALUE & DEMODULATION

FOR ANALYSIS IT IS CONVENIENT TO REPRESENT
 M_x, M_y AS A COMPLEX NUMBER

RECEIVE SIGNAL (analysis):

$$S_r(t) = \underbrace{A(t)}_{\text{complex}} e^{-i(\omega_c t + \phi(t))} \quad \underbrace{\phantom{e^{-i(\omega_c t + \phi(t))}}}_{\text{real}}$$

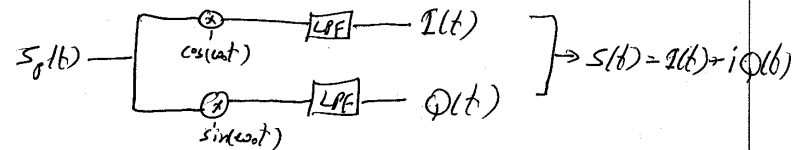
PHYSICAL SIGNAL:

$$S_p(t) = \text{Re}\{S_r(t)\} = \underbrace{A(t) \cos(\phi(t))}_{\substack{\text{I(t)} \\ \text{in phase}}} \cos(\omega_c t) + \underbrace{[-A(t) \sin(\phi(t))]}_{\substack{\text{Q(t)} \\ \text{quadrature}}} \sin(\omega_c t)$$

BaseBand SIGNAL (analysis):

$$s(t) = S_r(t) e^{+i\omega_c t} = A(t) e^{-i\phi(t)} = I(t) + iQ(t)$$

QUADRATURE PHASE-SENSITIVE DETECTION:



$$\text{Re}\{S(t)\} \rightarrow M_x \text{ in Rot Frame}$$

SO FAR

- (1) PLACE SAMPLE IN B_0
- M_z DEVELOPS $\sim ST_1$
- (2) EXCITE USING $B_1(t)$ TIP AWAY FROM \hat{z}
- (3) INSTANTANEOUS PRECESSION OF M_{xy}
PICK UP INDUCED EMF IN RF COIL
- (4) ENCODE POSITION IN FREQUENCY USING GRADIENTS
 \Rightarrow 1D PROJECTION.

~~(5) HOW DO WE GET AN IMAGE?~~

SEVERAL KEY COMPONENTS:

- ~~(1) SELECTIVE EXCITATION (dimension reduction)~~
- ~~(2) SPATIAL ENCODING~~

(3) LIMITATIONS:

GRADIENT + DURATION \rightarrow RESOLUTION
STRENGTH \uparrow
limited!

SIGNAL DECAY (T_2), FIELD INHOMOGENEITY (B_0)
DIFFUSION

TYPICAL RES: $< 1 \text{ mm}$ $\sim 20 \mu\text{m}$ TO SMALL ANIMAL SCANNERS