HST.583 Functional Magnetic Resonance Imaging: Data Acquisition and Analysis Fall 2008

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HST.583: Functional Magnetic Resonance Imaging: Data Acquisition and Analysis, Fall 2008 Harvard-MIT Division of Health Sciences and Technology Course Director: Dr. Randy Gollub.

MR physics and safety for fMRI

MGH



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Outline:

Wed. Sept 24 (LLW): MR signal, Gradient and spin echo Basic image contrast Mon. Sept 29 (LLW): Encoding the image Wed Oct 1 (LLW): Fast imaging for fMRI, artifacts fMRI BOLD

What is NMR?

NUCLEAR MAGNETIC RESONANCE

A magnet, a glass of water, and a radio wave source and detector....

What is NMR? Nuclear magnetism





Wald, fMRI MR Physics

S

Compass needles





Freq = γ **B**

Gyroscopic motion



- Proton has magnetic moment
- Proton has spin (angular momentum)
- >>gyroscopic precession



Larmor precession freq. = 42.58 MHz/T

EXCITATION : Displacing the spins from Equilibrium (North)

Problem: It must be moving for us to detect it. Solution: knock out of equilibrium so it oscillates

How? 1) Tilt the magnet or compass suddenly2) Drive the magnetization (compass needle) with a periodic magnetic field

Excitation: Resonance

Why does only one frequency efficiently tip protons?

Resonant driving force. It's like pushing a child on a swing in time with the natural oscillating frequency.



The RF pulse rotates Mo the about applied field

"Exciting" the Magnetization: tip angle



Detecting the Magnetization: Faraday's Law



A moving bar magnet induces a Voltage in a coil of wire. (a generator...)

The RF coil design is the #1 determinant of the system SNR

 $V(t) = -d\Phi/dt$

$$\Phi = \mathbf{n} B_{spins} A$$

Detecting the NMR: the noise



Noise comes from electrical losses in the resistance of the coil or electrical losses in the tissue.

For a resistor: Pnoise = 4kTRB

- Noise is white. >>Noise power α bandwidth
- Noise is spatially uniform.
- R is dominated by the tissue. >> big coil is bad.



Signal to Noise Ratio in MRI

- Most important piece of hardware is the RF coil.
- SNR α voxel volume (# of spins)
- SNR α SQRT(total time of data collection)
- SNR depends on the amount of signal you <u>throw away</u> to better visualize the brain (gain image contrast)

Physical Foundations of MRI

NMR: 60 year old phenomena that generates the signal from water that we detect.

MRI: using NMR signal to generate an image

Three magnetic fields (generated by 3 coils)
1) static magnetic field Bo
2) RF field that excites the spins B1
3) gradient fields that encode spatial info G_x, G_y, G_z

Three Steps in MR:

 0) Equilibrium (magnetization points along Bo)
 1) RF Excitation (tip magn. away from equil.)
 2) Precession induces signal, dephasing (timescale = T2, T2*).
 3) Return to equilibrium (timescale = T1).

Magnetization vector during MR





Three places in process to make a measurement (image)

0) Equilibrium (magnetization points along Bo)

1) RF Excitation (tip magn. away from equil.)

2) Precession induces signal, allow to dephase

3) Return to equilibrium (timescale =T1).

Contrast in MRI: proton density

Form image immediately after excitation (creation of signal).

Tissue with more protons per cc give more signal and is thus brighter on the image.

No chance to dephase, thus no differences due to different tissue T2 values.

Magnetization starts fully relaxed (full Mz), thus no T1 weighting.



T2*-Dephasing

Wait time <u>TE</u> after excitation before measuring M.

Shorter T2* spins have dephased



T2* Dephasing

Just the tips of the vectors...



T2* decay graphs



T2* Weighting

Phantoms with four different T2* decay rates...

There is no contrast difference immediately after excitation, must wait (but not too long!).

Choose TE for max. inten. difference.



Gradient Echo (T2* contrast)

Dephasing is entirely from a spatial difference in the applied static fields.



 $B_{o} + G_{x} \times$

Red arrows processes faster due to its higher local field

Gradient Echo (T2* contrast) Dephasing is entirely from a spatial difference in the applied static fields.



Gradient Echo



7T 32ch MGH array

2D FLASH, TR/TE3 500/30 0.22 x 0.22 x 1mm (48nl) 8min at

7T 32ch MGH array



Courtesy of Dr. Christopher J. Wiggins. Used with permission.

G. Wiggins, C. Wiggins, Martinos Center MGH

2D FLASH, TR/TE₃ 500/30 0.22 x 0.22 x 1mm (48nl) 8min acq

Wald, RSNA 2007

A.A. Martinos Center, MGH Radiolog



7 Tesla 230um

2D FLASH 0.23 x 0.23 x 1.5mm 8min acq



Courtesy of Dr. Christopher J. Wiggins. Used with permission.

R

TP 0 SP F21.0 SL 1.5 FoV 208*238 896*1024s

7 T, 32ch 200um x 200um x 1mm

2D T2* weighted 200um x 200um x 1mm (1024x1024 matrix)

>Co "U-fiber" ??

Courtesy of Dr. Christopher J. Wiggins. Used with permission.

Spin Echo (T2 contrast)

Some dephasing can be refocused because its due to static fields.



Blue arrows precesses faster due to local field inhomogeneity than red arrow

Spin Echo

180° pulse only helps cancel static inhomogeneity

The "runners" can have static speed distribution.

If a runner trips, he will not make it back in phase with the others.







Other contrast for MRI

In brain: (gray/white/CSF/fat) Proton density differ ~ 20% T1 relaxation differ ~ 2000%

How to exploit for imaging?

Vary repetition rate - TR



grey matter (long T1) white matter (short T1)

T1-Weighting







white matter (short T1)



Image contrast summary: TR, TE



Source of T1 and T2 contrast in brain: Myelin content

Image removed due to copyright restrictions. Diagram showing the arrangement of nerve cells and fibers in layers and sublayers parallel to the surface in a vertical section through the human striate area or visual cortical center.

Determine functional boundaries based on MR strucure alone...

Nissel stain: cell bodies

Weigert stain: fibers

Cortical layers in Monkey at 7T





Intensity along line perpendicular To V1

MPRAGE 250um x 250um x 750um (4 hours)