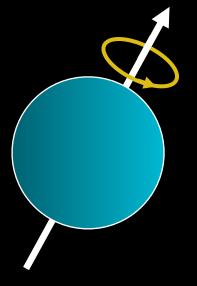
Introduction to MRI Physics

Slides originally by Karla Miller, FMRIB Centre

Modified by Mark Chiew (mark.chiew@ndcn.ox.ac.uk)



Slides available at:

http://users.fmrib.ox.ac.uk/~mchiew/teaching/

MRI Physics

Monday:

- ★ Basics of magnetic resonance
- ★ Image formation
- ★ Signal statistics (SNR)
- ★ Functional MRI

Wednesday:

- * Image contrast (T_2 and T_2^*)
- * Spin vs. gradient echo
- ★ Fast imaging
- ★ Diffusion MRI

What are we trying to achieve?

Informed decision making: You need to take responsibility for the design, implementation & execution of your study

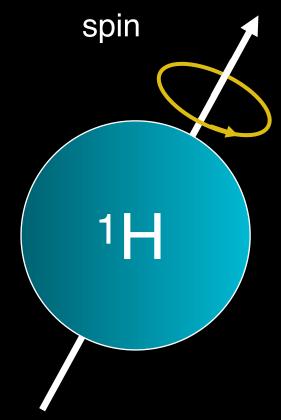
- Protocols need to be tailored to the problem
- Learning some physics will make this less daunting

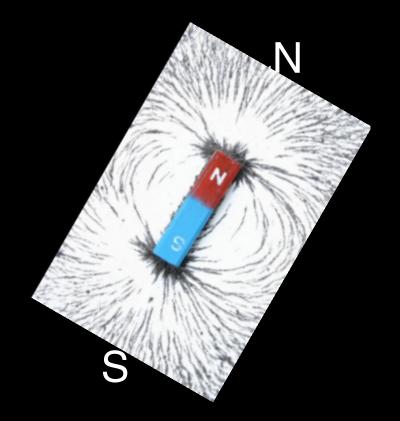
A common language: You need to be able to talk to experts

- Communicate your needs to physicists/radiographers/techs
- Build an MR vocabulary (terminology/jargon)
- Gain some intuition behind imaging concepts

MRI Physics

- ★ Basics of magnetic resonance
- ★ Image formation
- ★ Signal statistics (SNR)
- ★ Functional MRI





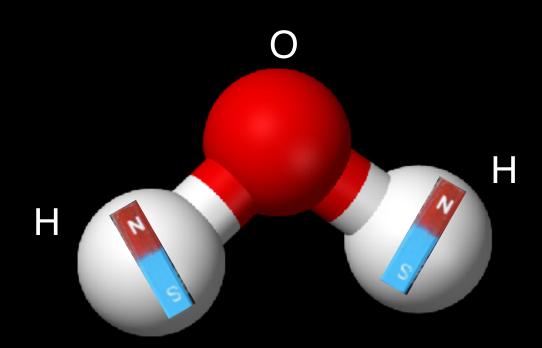
Almost all sub-atomic particles have "spin"

"Spin"

 All nuclei with odd numbers of protons/neutrons will have non-zero net spin

¹H, ¹³C, ²³Na, ³¹P

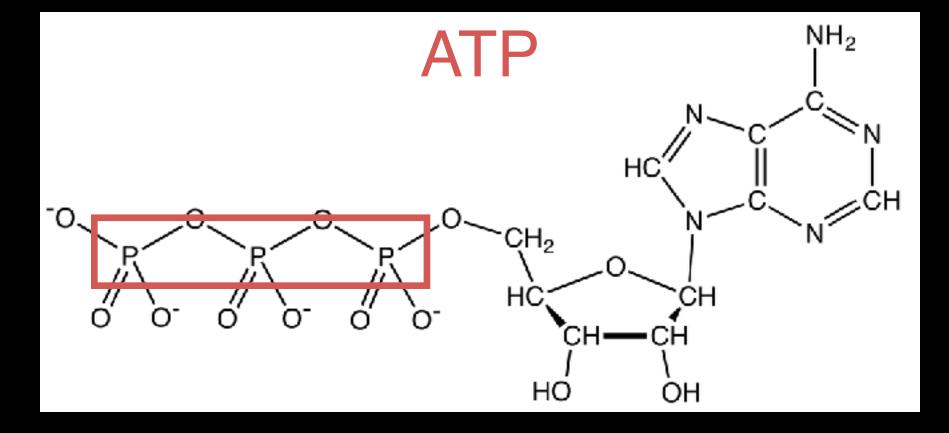
All hydrogen protons will act like little magnets



The abundance of water in the human body makes this very powerful!

¹H, ¹³C, ²³Na, ³¹P

Can also do this with phosphorous nuclei



Spins, or magnetization (when referred to in bulk) behave similarly to classic physical systems

In many ways analogous to simple oscillators, like swings or pendulums



1. Excitation

Magnetization can be moved or rotated by applying "excitation" magnetic fields (RF)

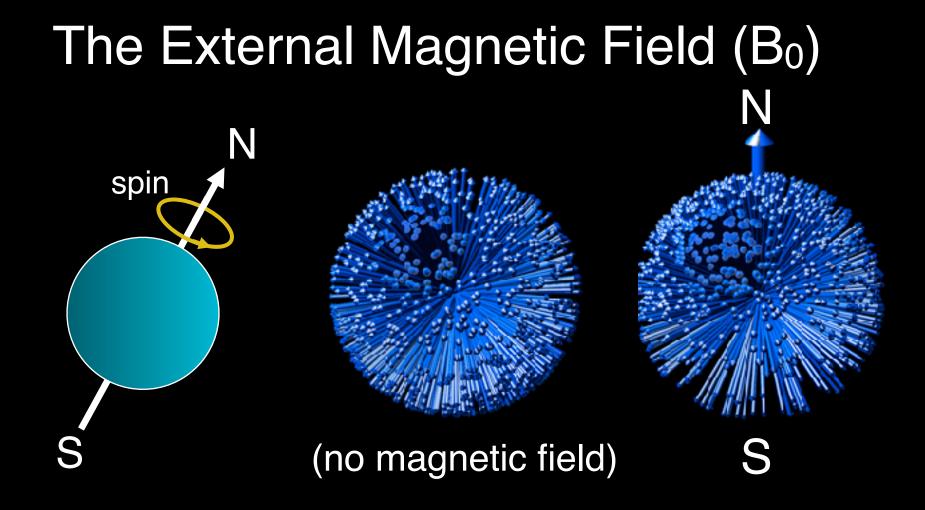
2. Resonance

Magnetization will "resonate" at a frequency proportional to magnetic field strength

3. Relaxation

The oscillations die out, i.e. magnetisation "relaxes" back to equilibrium – speed of relaxation is tissue-dependent!



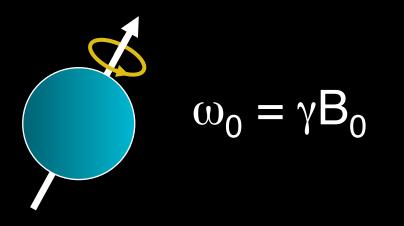


Normally: protons randomly oriented \Rightarrow no net magnetism

External field: protons align slightly \Rightarrow *net magnetization (M)* Only a few parts-per-million!

Magnetic resonance

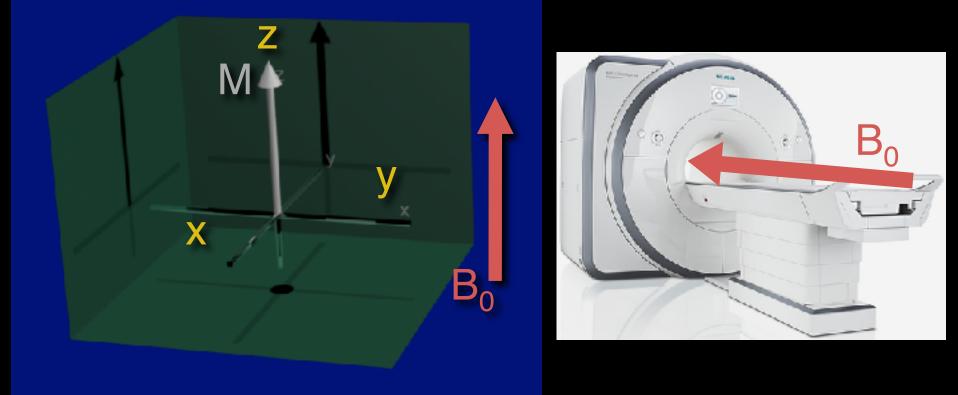
Magnetic: external field (B₀) magnetizes sample



This "Larmor Equation" defines the resonant frequency

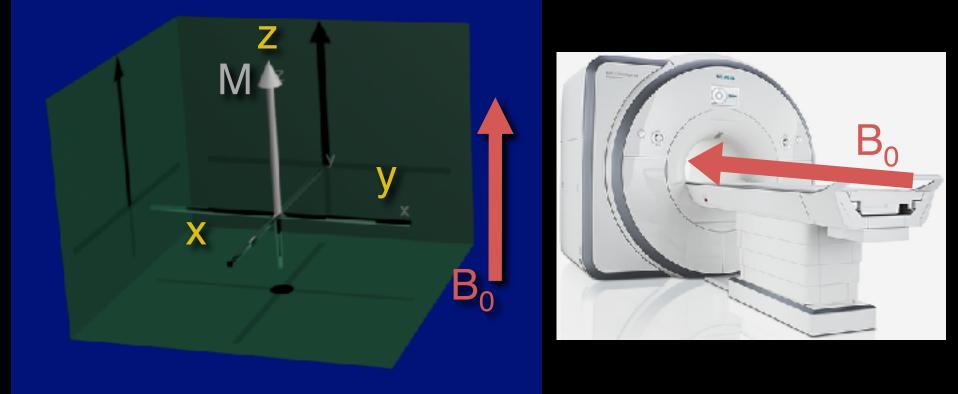
Resonance: magnetization has characteristic (resonant) frequency proportional to external field B₀

Coordinate system



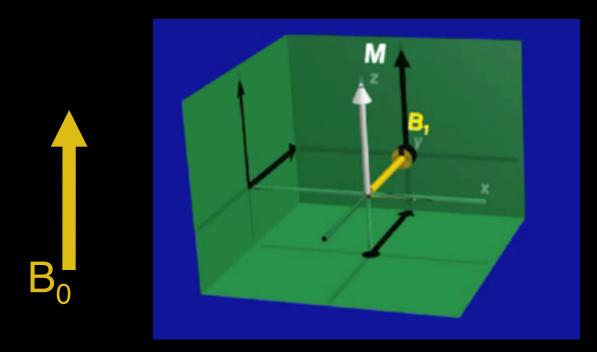
Direction of main field (B₀) defines coordinate system Longitudinal axis: parallel to B₀ (typically z) Longitudinal magnetisation: Portion of M aligned with B₀

Coordinate system



Direction of main field (B₀) defines coordinate system Transverse plane: perpendicular to B₀ (typically x,y) Transverse magnetisation: Portion of M perpendicular to B₀

The Basic MRI Experiment: 1. Excitation

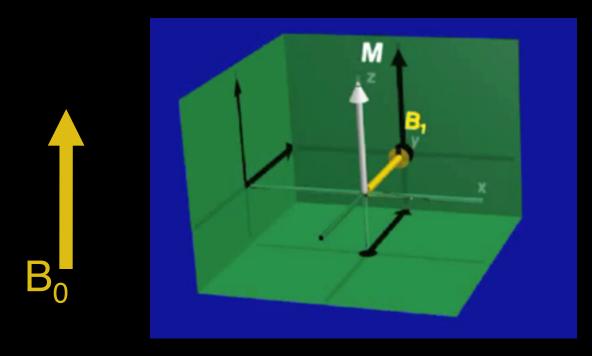


courtesy of William Overall

$$\omega_0 = \gamma B_0$$

Excitation pulse (yellow) tips magnetisation away from B₀ Excitation must occur at the resonant frequency ω_0

The Basic MRI Experiment: 1. Excitation

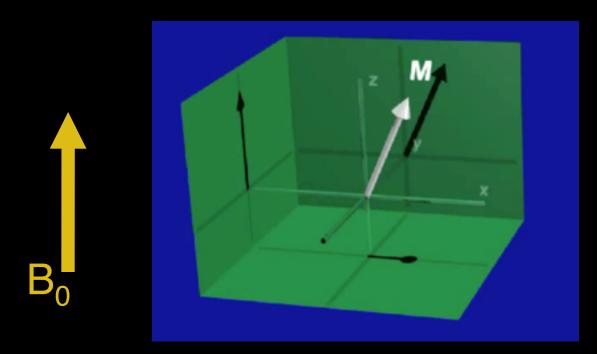


courtesy of William Overall

In a frame that rotates with B_1 , magnetisation is simply "flipped" or "tipped" out of alignment with B_0

Hence the term "flip angle" or "tip angle"

The Basic MRI Experiment: 2. Resonant Precession

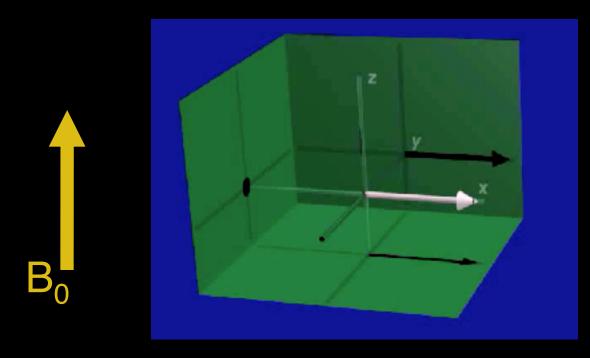


courtesy of William Overall

$$\omega_0 = \gamma B_0$$

Once excited, magnetisation precesses/oscillates/rotates at resonance frequency

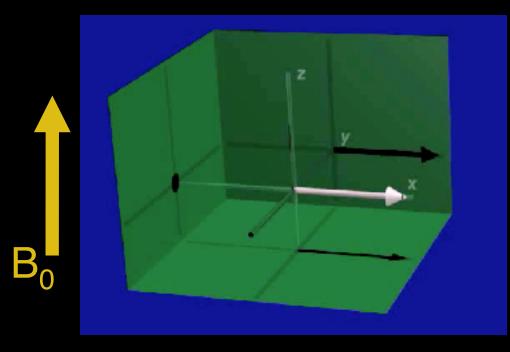
The Basic MRI Experiment: 3. Relaxation



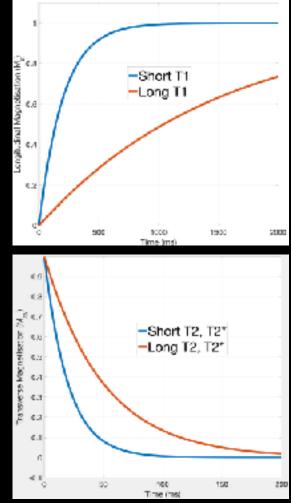
courtesy of William Overall

As it precesses, it also "relaxes" back into alignment with B_0 Speed of relaxation has time constants: T_1 , T_2 , T_2^* , which relate to the image contrast

The Basic MRI Experiment: 3. Relaxation

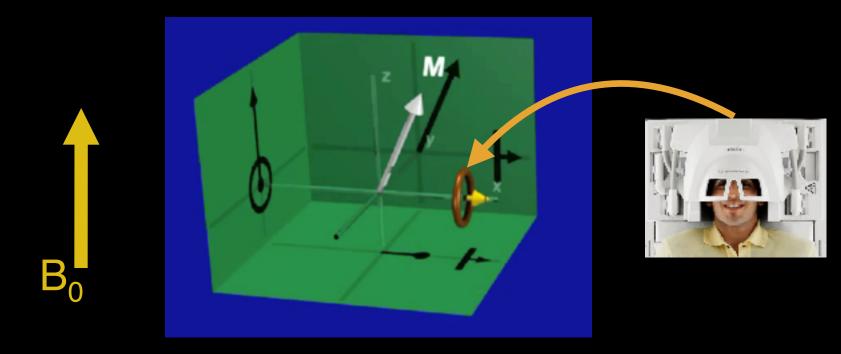


courtesy of William Overall



 T_1 : describes speed of recovery along longitudinal (z) axis T_2 , T_2^* : describe speed of signal decay in transverse (x-y) plane

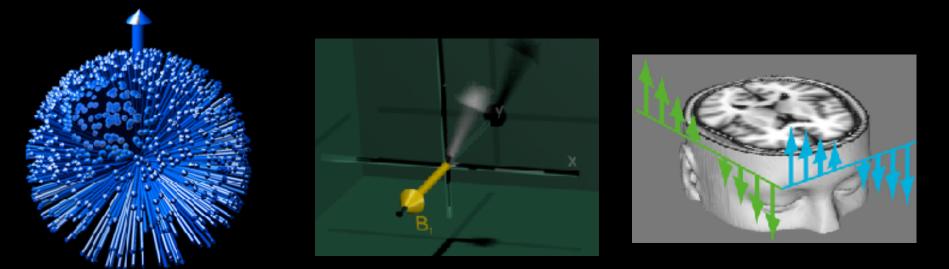
The Basic MRI Experiment: 4. Signal Detection



courtesy of William Overall

As the magnetization precesses and relaxes The precession induces a voltage in the receive coils Coils only detect rotating, transverse magnetisation

Magnetic fields everywhere...



B₀

B₁

 G_x, G_y

Main magnetic field (B₀): always on, static

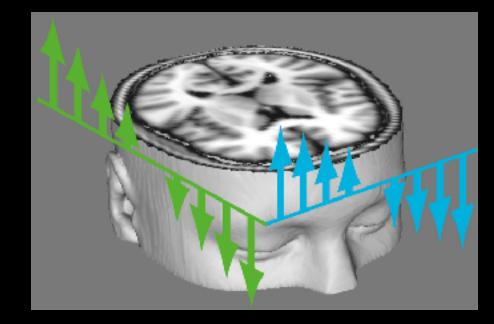
Excitation RF field (B₁): pulsed on & off, 60-300 MHz

Magnetic field gradients (G): pulsed on & off, "static"

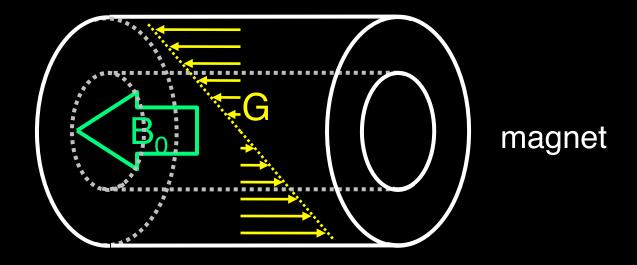
MRI scans: carefully timed RF and gradient "pulse sequences"

MRI Physics

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Magnetic Field Gradients

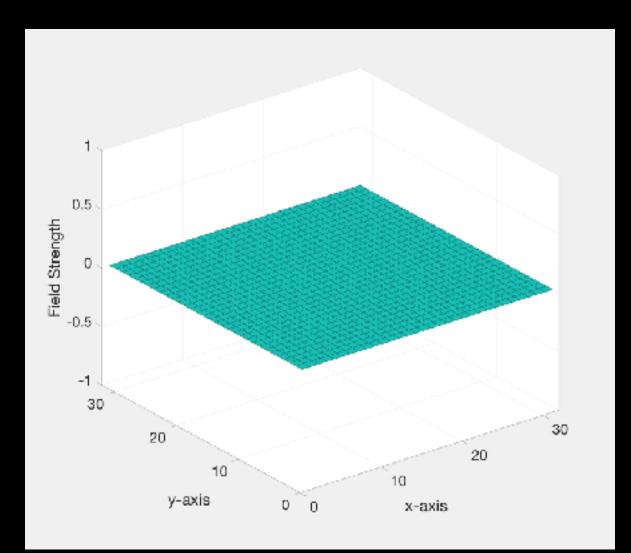


Differentiate between signal from different locations

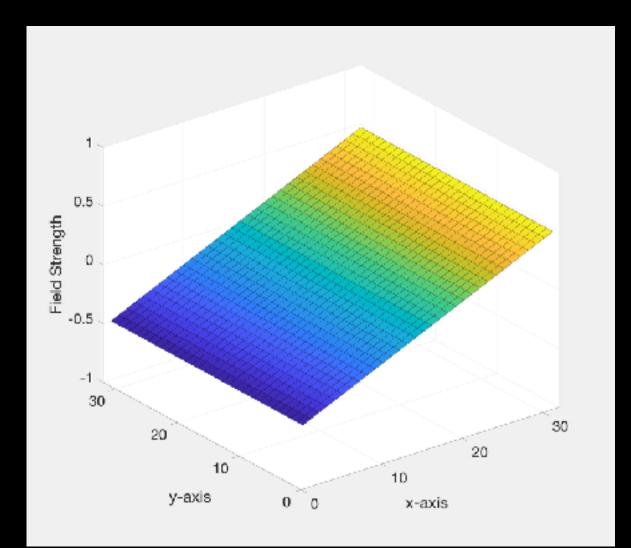
Add a spatially varying magnetic field gradient (G)

- Field varies linearly along one direction
- Gradient field adds to or subtracts from B₀

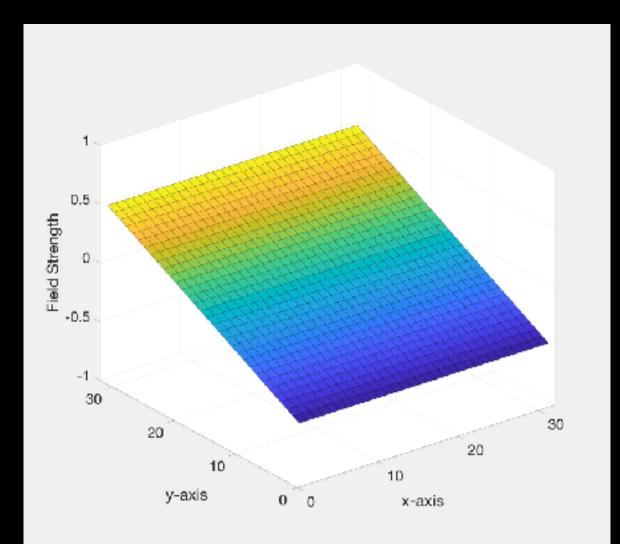
No Gradient



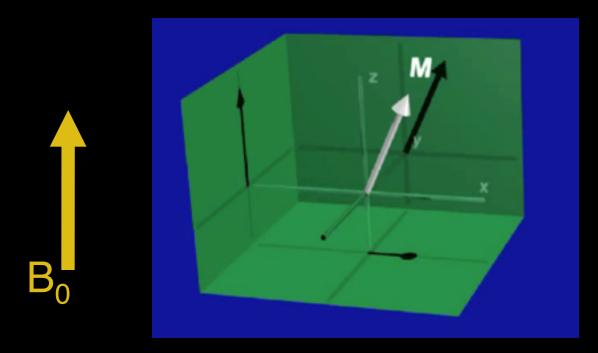
x-Gradient



y-Gradient



Precession



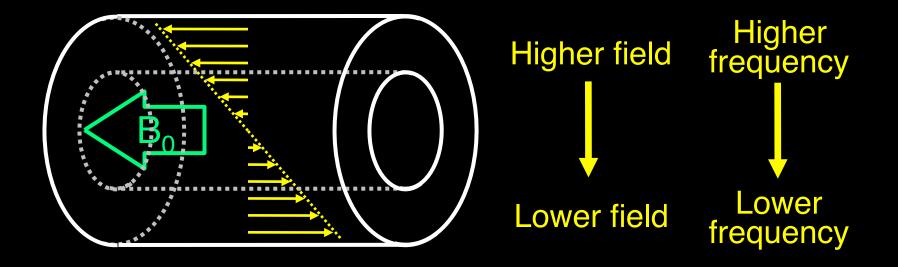
courtesy of William Overall

$$\omega_0 = \gamma(\mathsf{B}_0 + \mathsf{G}(\mathsf{x}, \mathsf{y}, \mathsf{z}))$$

Resonance frequency is proportional to total field: Static B₀ + applied gradients

Gradients and Resonance

$$\omega_0 = \gamma(\mathsf{B}_0 + \mathsf{G}(\mathsf{x}, \mathsf{y}, \mathsf{z}))$$

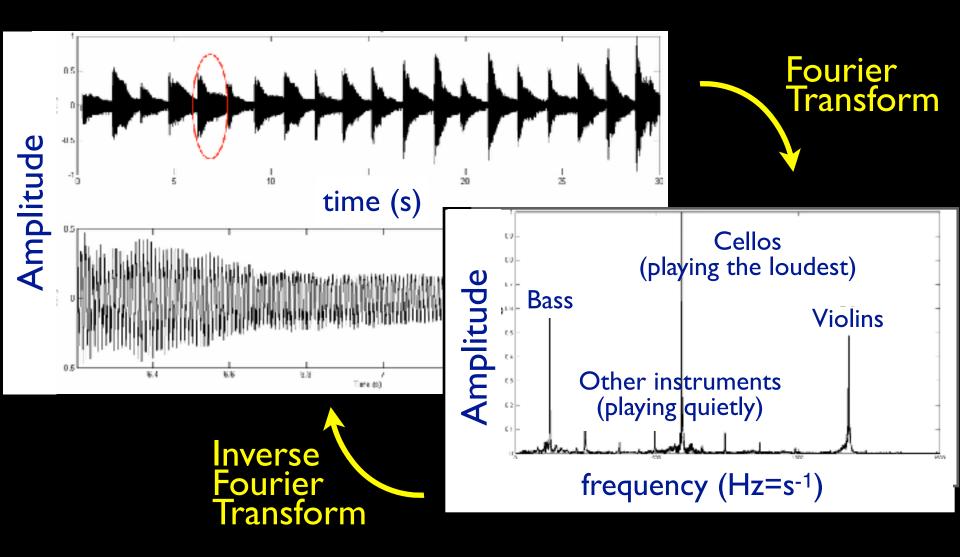


Distinguish different spatial locations by assigning different resonant frequencies to different positions

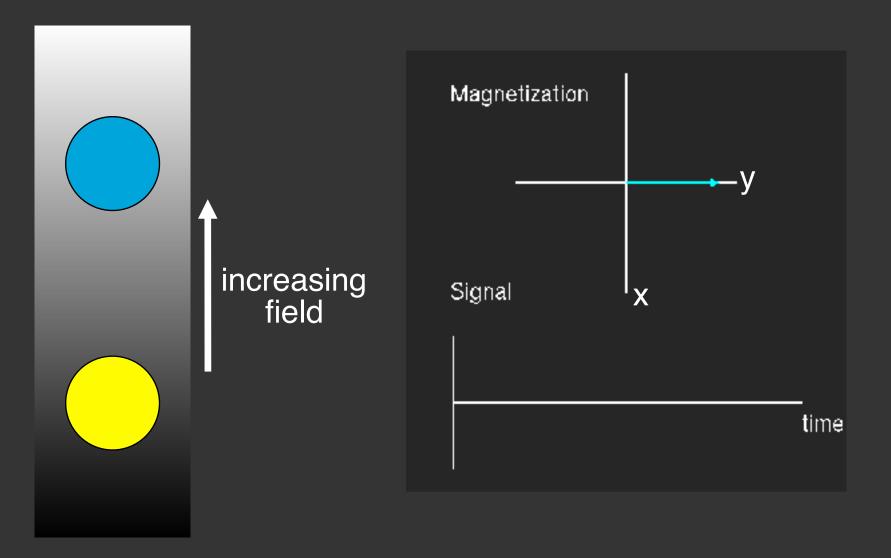
Imagine differentiating between instruments based on their frequency content!

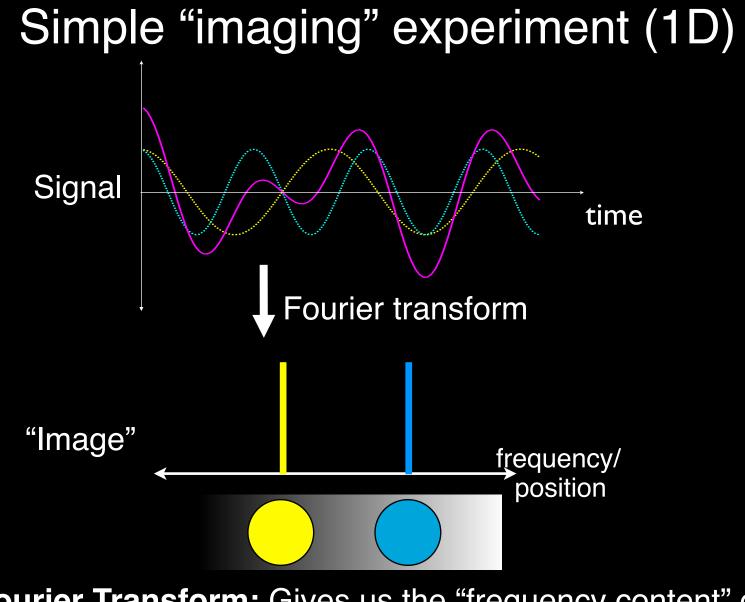


Frequency decomposition

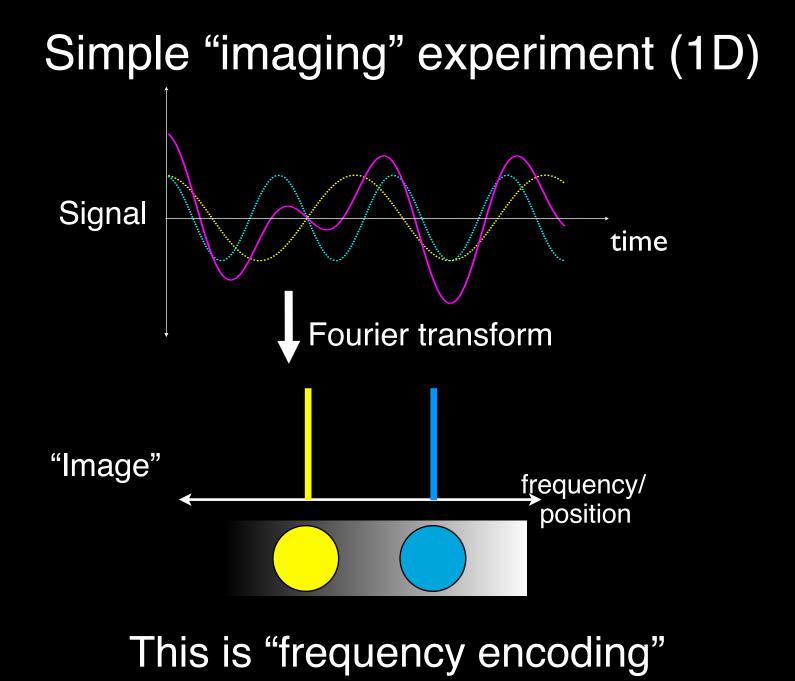


Simple "imaging" experiment (1D)

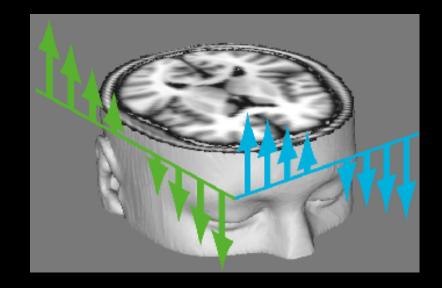




Fourier Transform: Gives us the "frequency content" of our signals



Magnetic gradients



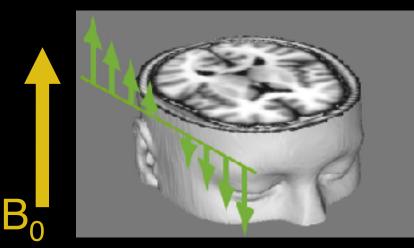
G_x, G_y

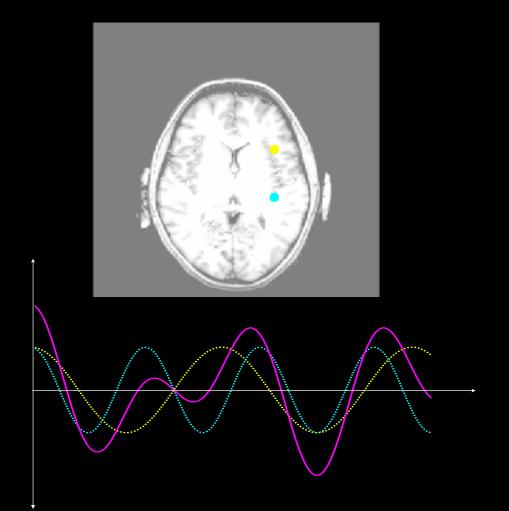
It's a bit more complex in more than 1 dimension Have 3 gradient fields (along x, y, z) Manipulate the strength & timing independently

Gradients in multiple dimensions

0.05			
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Magnetic field gradients

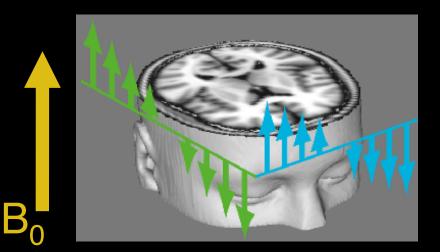


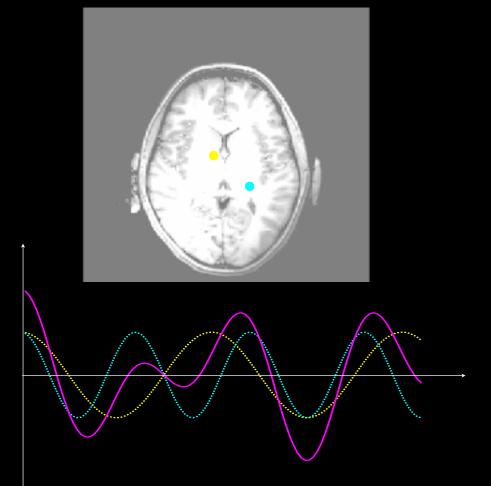


Gradients in multiple dimensions

0.05			
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Combined field gradients





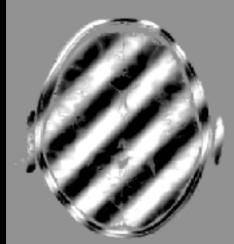
Spatial frequencies or patterns

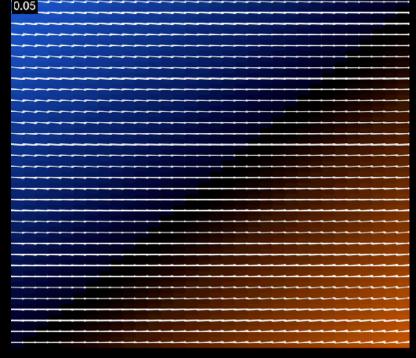
At any instant in time, signal is across space is defined by a specific "pattern" of the magnetisation phase (orientation),

i.e., its *spatial frequency* that depends on the applied gradients

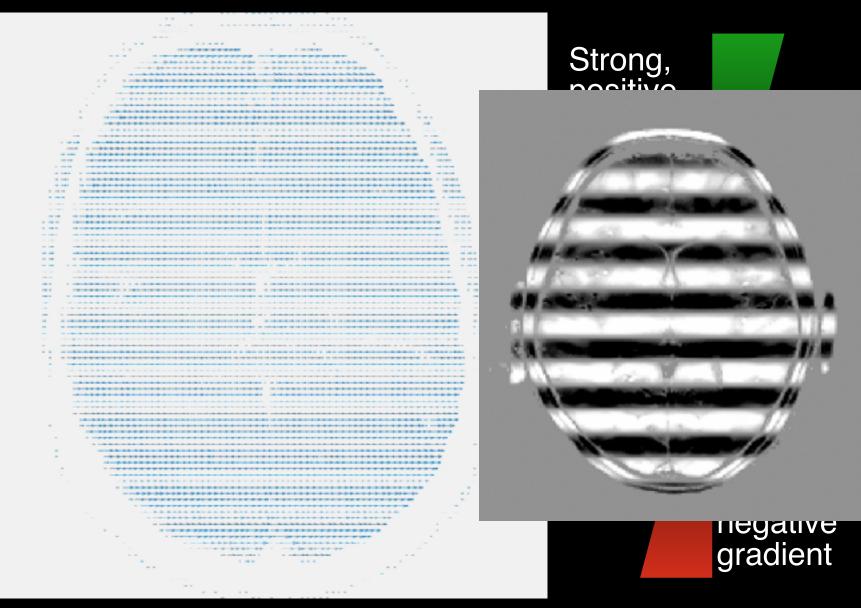
Spatial frequencies:

- sinusoidal pattern over space instead of time
- extend to multiple dimensions

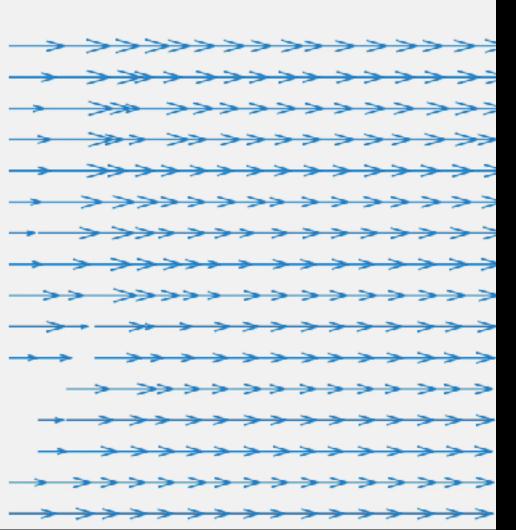




Gradients and Spatial Frequency



Gradients and Spatial Frequency



higher resonance frequency

faster precession stronger gradient magnetic field

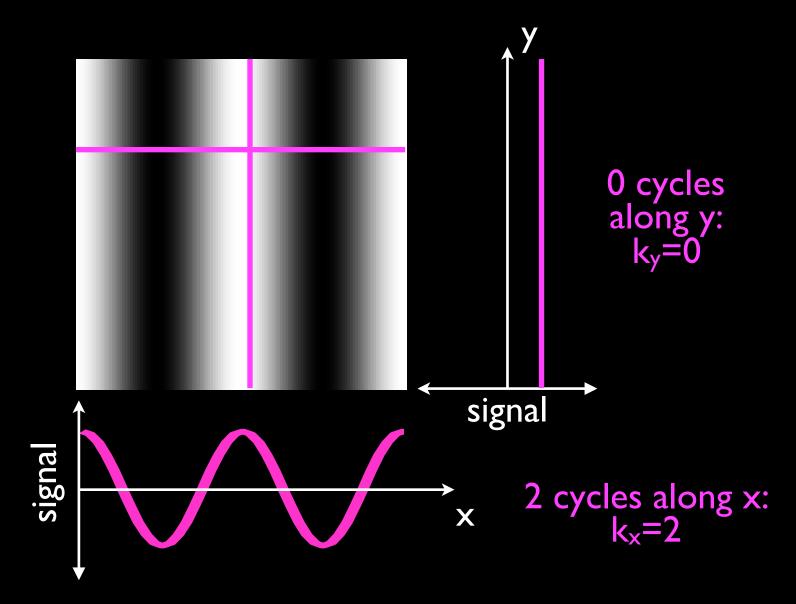
lower resonance frequency

slower precession

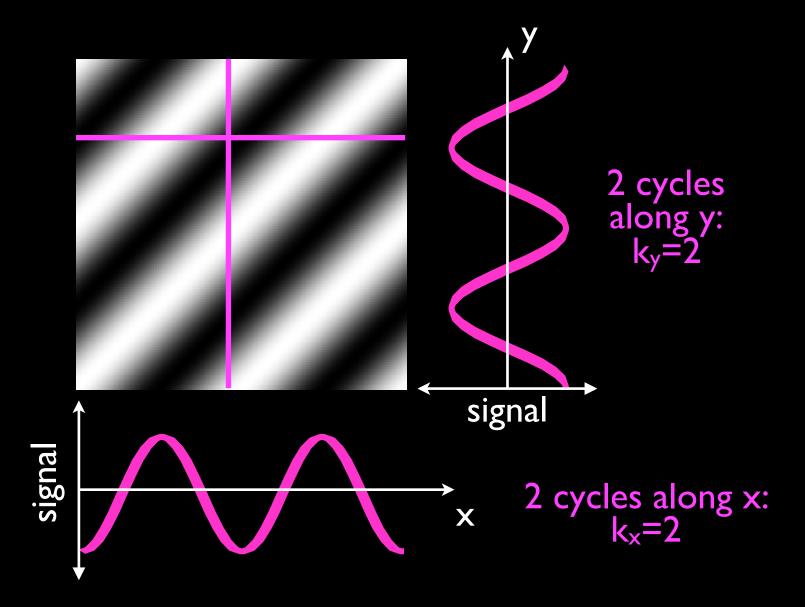
zero gradient

This is one spatial frequency... 2 cycles along y: k_y=2 signal signal 0 cycles along x: k_x=0 X

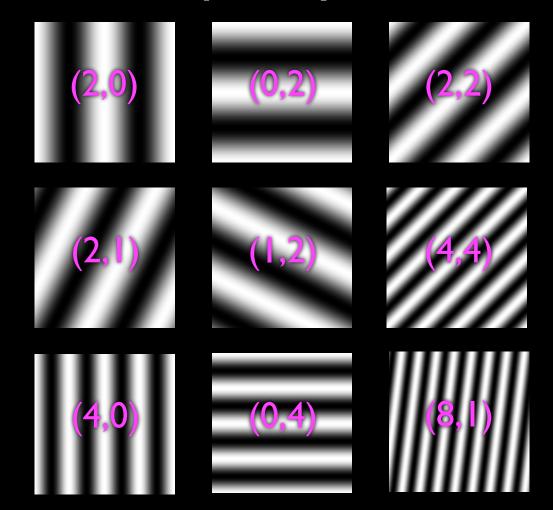
This is another one...



This is another one...

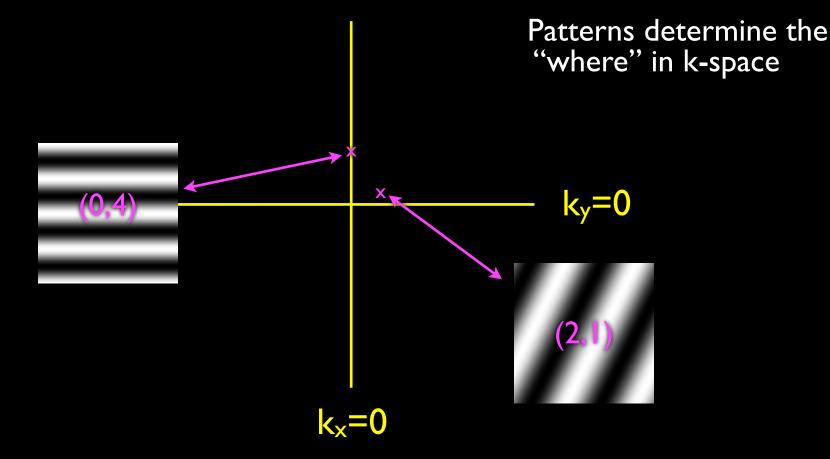


Each of these represents one 2D pattern or frequency: denote (k_x,k_y)



"k" values are the number of cycles in each direction

2D "k-space" describes contribution of each spatial frequency



2D "k-space" describes contribution of each spatial frequency

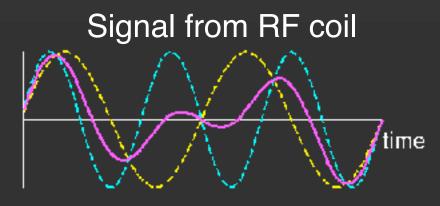
 $k_x=0$

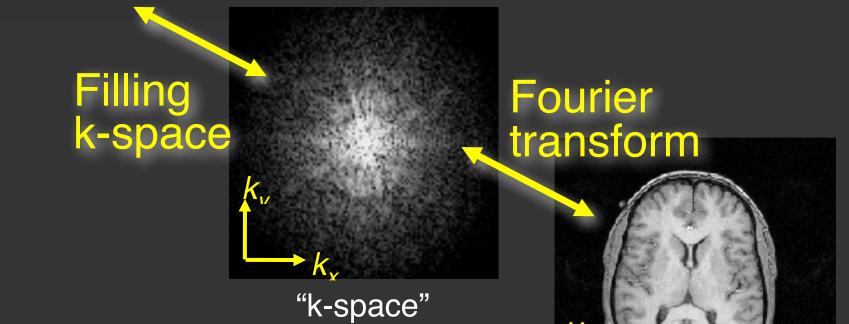
Sum total signal after application of these patterns determines the "value" of each k-space location





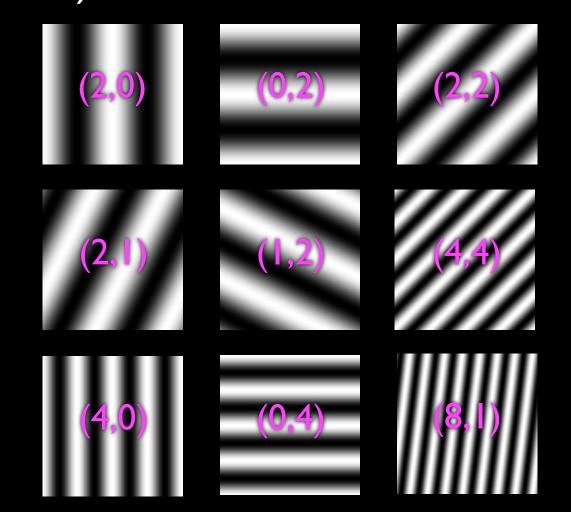




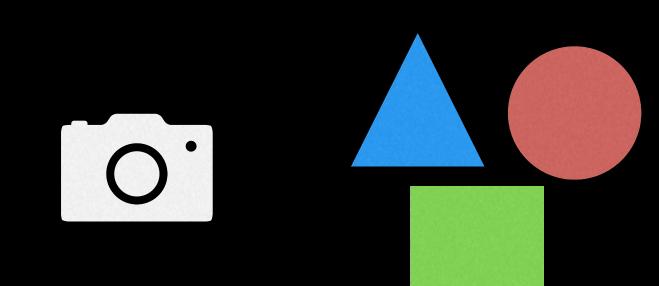


Image

Think of each pattern (k-space location) as a filter on a camera



Imagine our "camera" can only see one colour at a time



Imagine our "camera" can only see one colour at a time (blue filter)

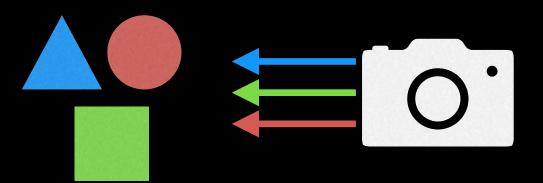
Imagine our "camera" can only see one colour at a time (red filter)

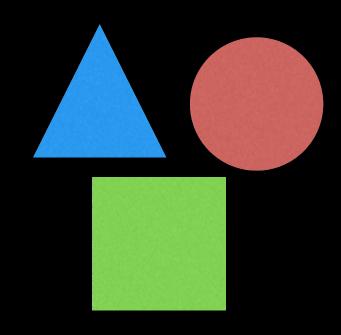
.

.

Imagine our "camera" can only see one colour at a time (green filter)

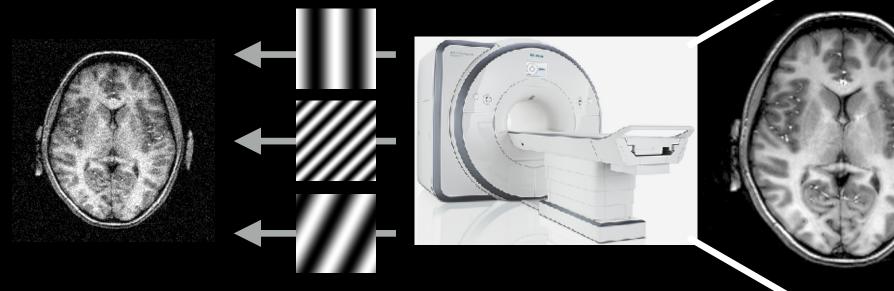
Combine the filtered images to form the final image





Scanner takes a series of measurements with each kspace "spatial filter" (as many filters as voxels)

The "spatial filters" are applied using gradients



Measurements are then combined using the Fourier Transform to form image

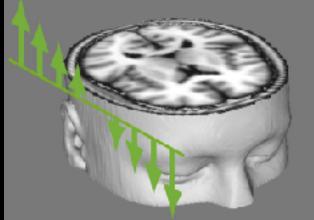
Scanner takes a series of measurements with each kspace "spatial filter" (as many filters as voxels)

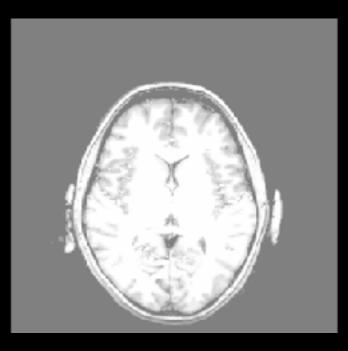
Higher resolution means "finer" features, which require "finer" filters



If gradient is on, spatial frequencies change over time

K_X



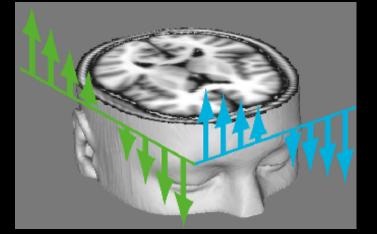


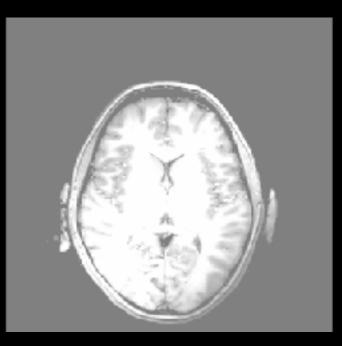
Visualize as gradients "move us" through k-space!

Ky

If gradient is on, spatial frequencies change over time

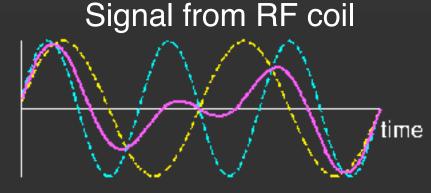
K_X



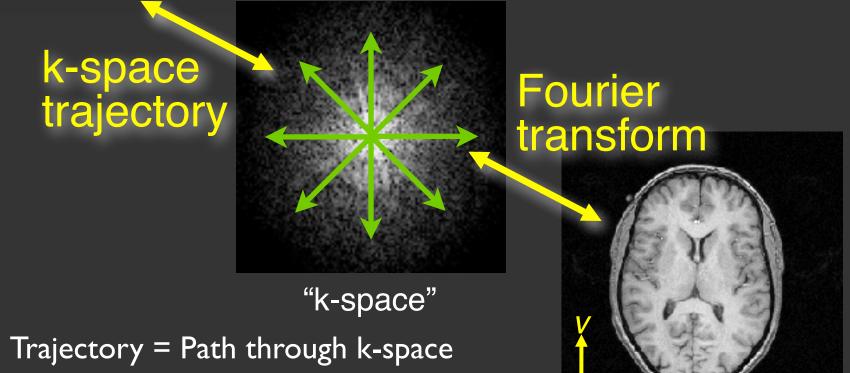


Visualize as gradients "move us" through k-space!

Ky



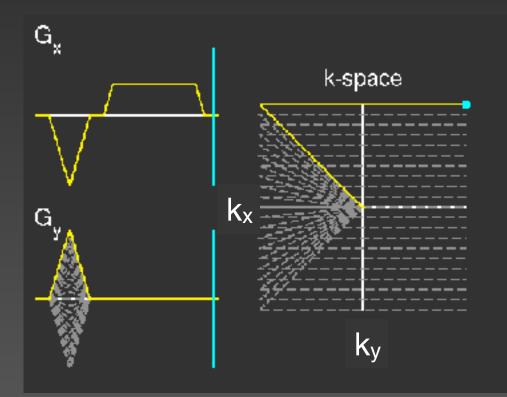
The trajectory is the ordering of k-space data acquisition



or the sequence of spatial filters

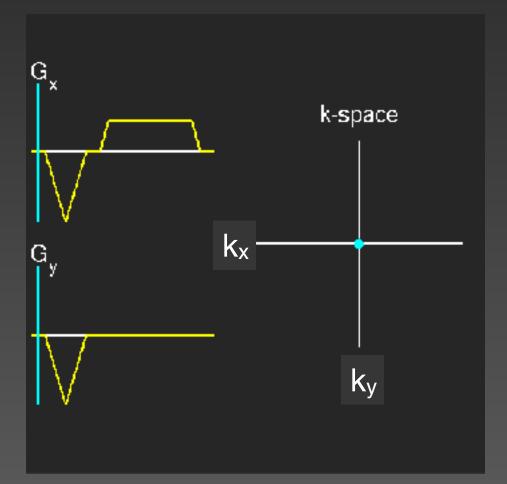
Image

Linescan (2DFT) Acquisition



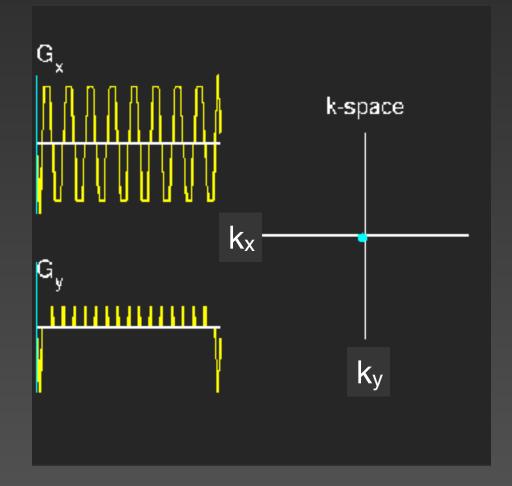
Acquire one line after each excitation

Linescan (2DFT) Acquisition



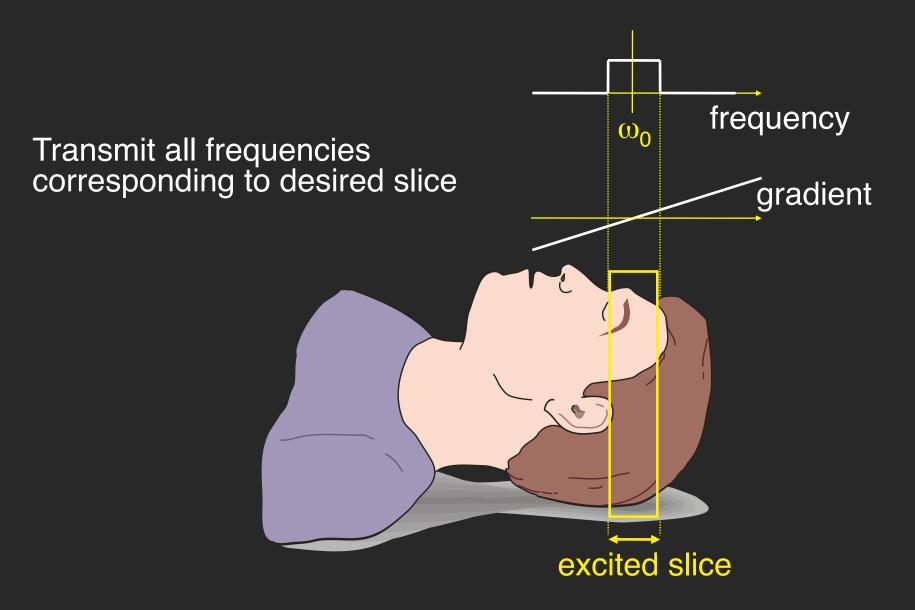
Acquire one line after each excitation Useful for structural images (minimal artifacts)

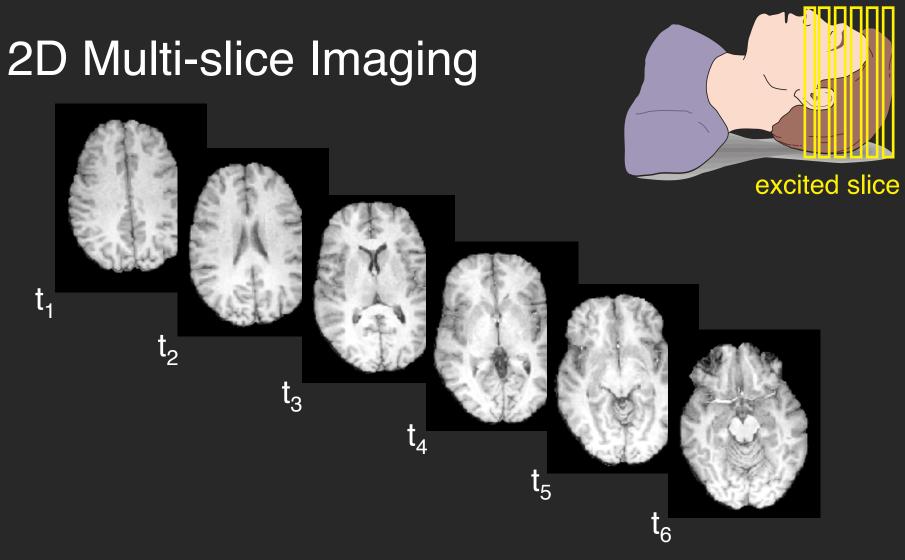
Echo-planar Imaging (EPI) Acquisition



Acquire all of k-space in a "single shot" Used for FMRI, diffusion imaging

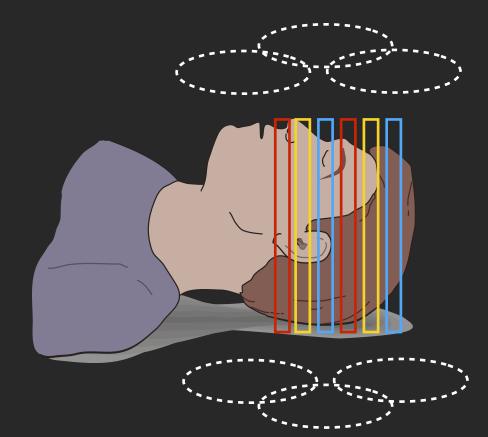
Slice Selection



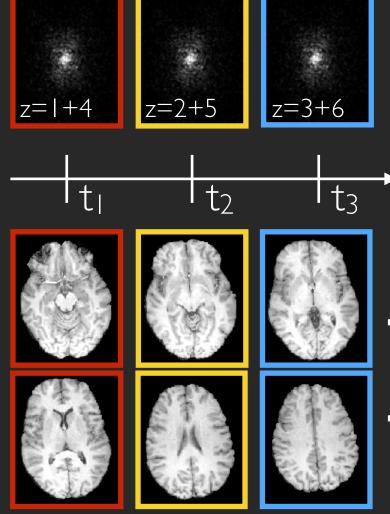


Slices excited and acquired sequentially (separately) Most scans acquired this way (including FMRI, DTI)

Simultaneous Multi-slice Imaging

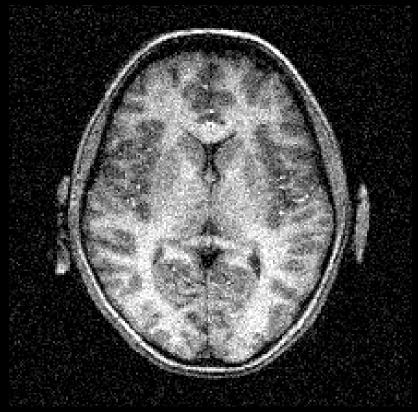




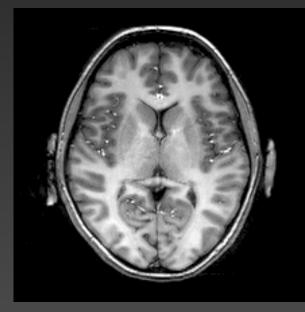


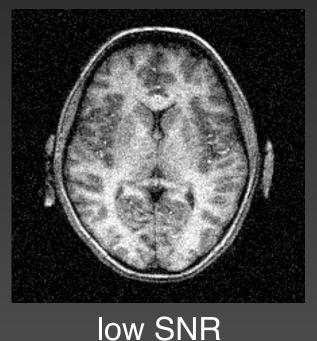
MRI Physics

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Signal-to-noise ratio (SNR)





high SNR

 $SNR = \frac{Signal}{\sigma_{noise}}$ (magnitude) (standard deviation)

Signal-to-noise ratio: describes signal "robustness" All else being equal, we want to maximize SNR!!

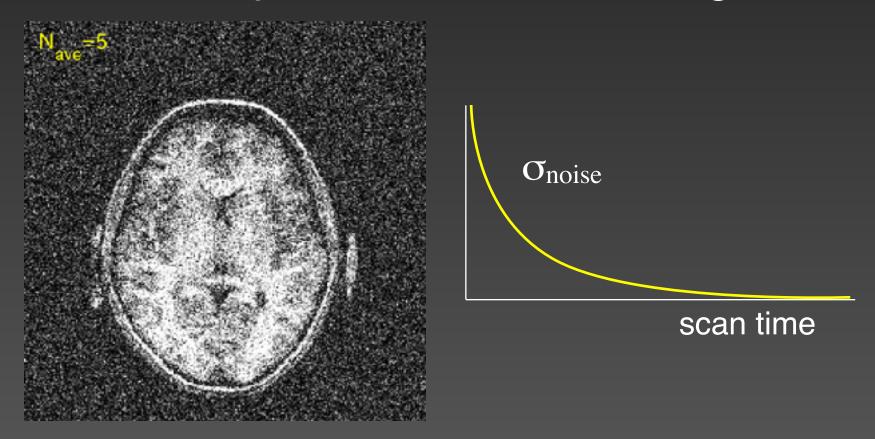
Signal-to-noise ratio (SNR)

SNR = 1SNR = 2SNR = 5SNR = 10 SNR = 50 SNR = 20

Protocol choices affecting SNR...

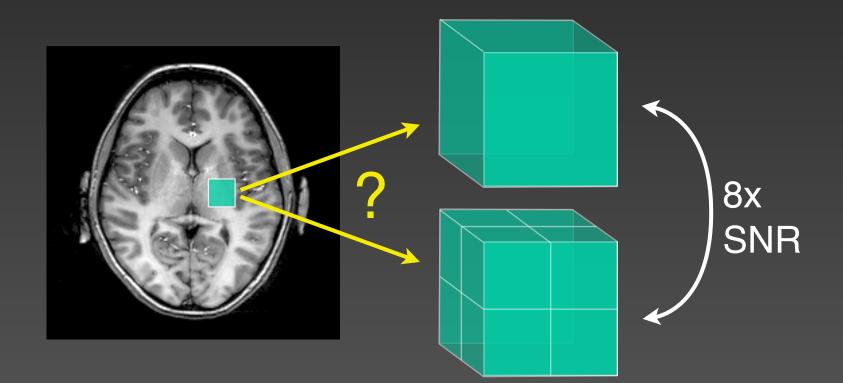
- RF receive coil & field strength
- Timing: bandwidth, TE & TR
- Voxel volume
- Scan duration (imaging time)
- Anything affecting signal!!!

SNR and acquisition time or averages



Longer acquisition \Rightarrow less noise \Rightarrow higher SNR SNR improves with the *square root* of scan time i.e., to *double* SNR you need to scan 4x longer

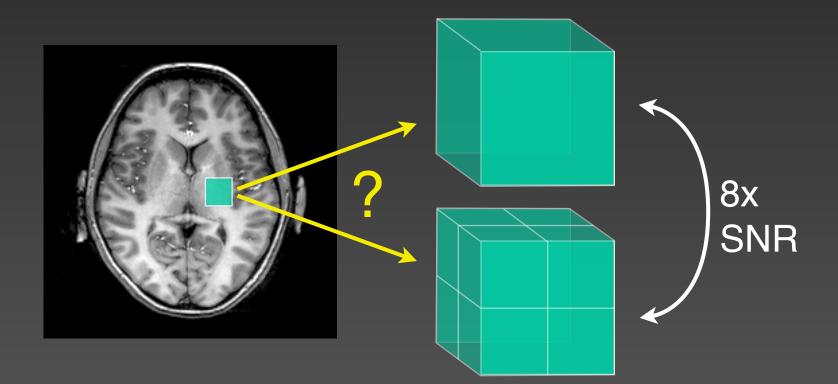
SNR and voxel volume



Larger voxels have signal from more tissue!

- Signal proportional to voxel volume
 - 2x2x2mm has 8x higher SNR than 1x1x1mm!

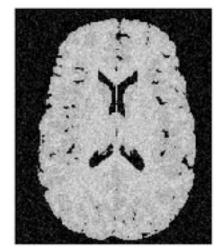
Averaging to achieve high resolution



Can we recover lost SNR by averaging? Yes! But requires a 64-fold increase in scan time (because you only get square root benefit)

Contrast-to-noise ratio (CNR)

SNR = 10, CNR = 1



SNR = 10, CNR = 6



SNR = 10, CNR = 2



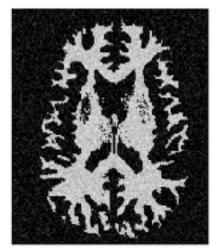
SNR = 10, CNR = 8



SNR = 10, CNR = 4

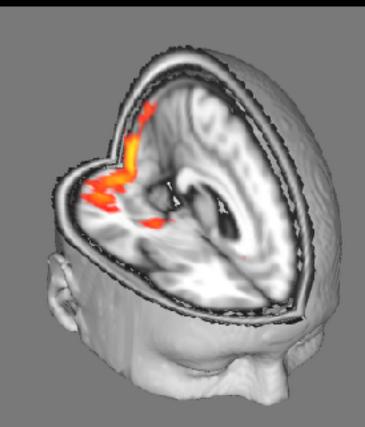


SNR = 10, CNR = 10

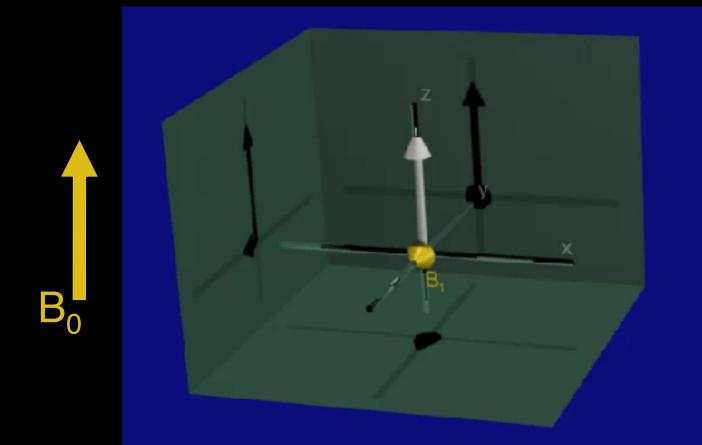


MRI Physics

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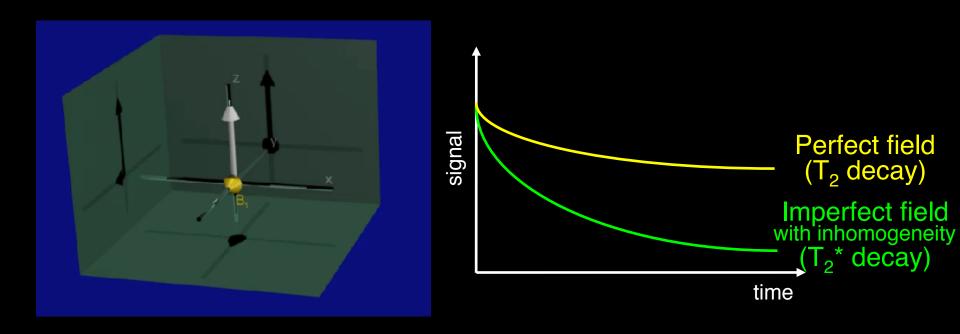


A source of signal loss: dephasing



When spins are "in-phase", they are all oriented the same way Over time, the spins within a voxel lose alignment ("dephase")

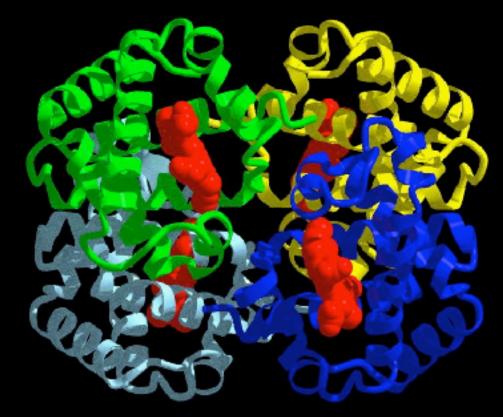
Apparent increase in $T_2 = T_2^*$



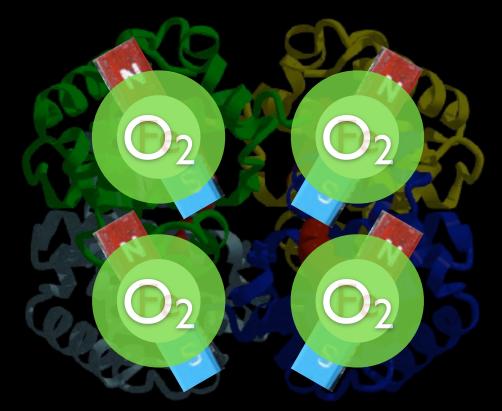
Dephasing causes magnetization vectors to partially "cancel" each other out

Dephasing results in a lower *net* signal magnitude Apparent decrease in T₂: called T₂* (more on Wednesday)

Deoxyhaemoglobin is the source of FMRI signal

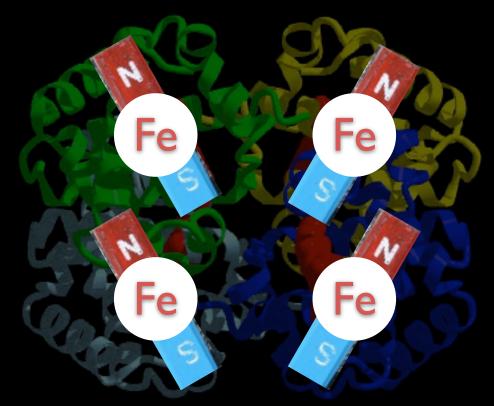


Deoxyhaemoglobin is the source of FMRI signal



When oxygen is bound to the haemoglobin, it shields the magnetic effects of iron atoms in the heme groups

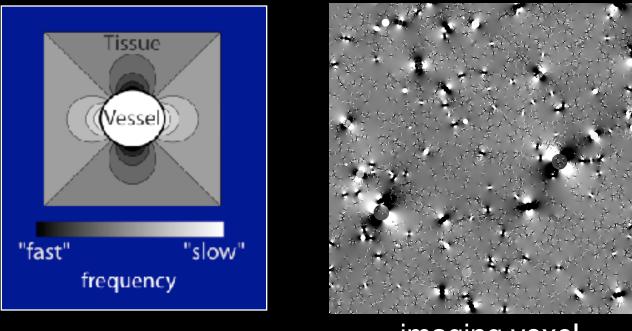
Deoxyhaemoglobin is the source of FMRI signal



Without oxygen, the iron (Fe) is exposed, causing magnetic field inhomogeneities due to its strong magnetic propertiesField inhomogeneity leads to T2* change (FMRI signals)

The BOLD Effect

[Ogawa et al, 1990]

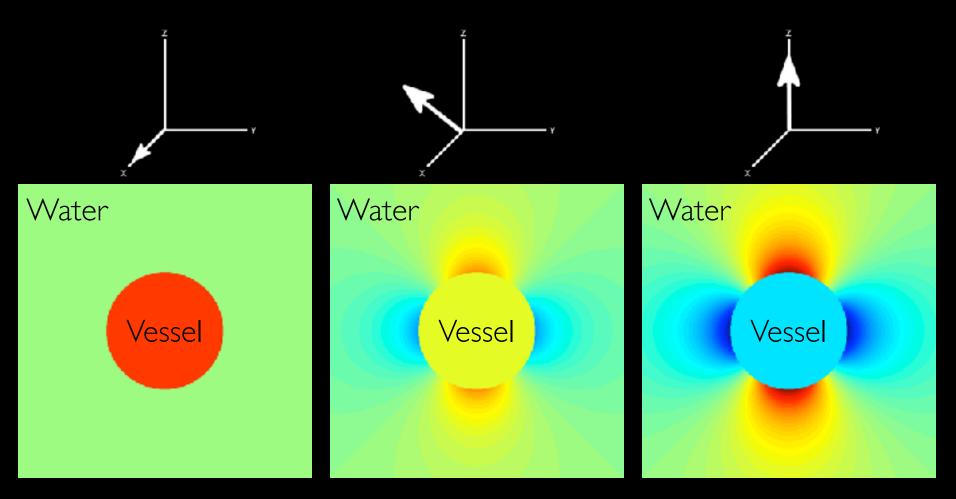


imaging voxel

Blood Oxygenation Level Dependent (BOLD) effect

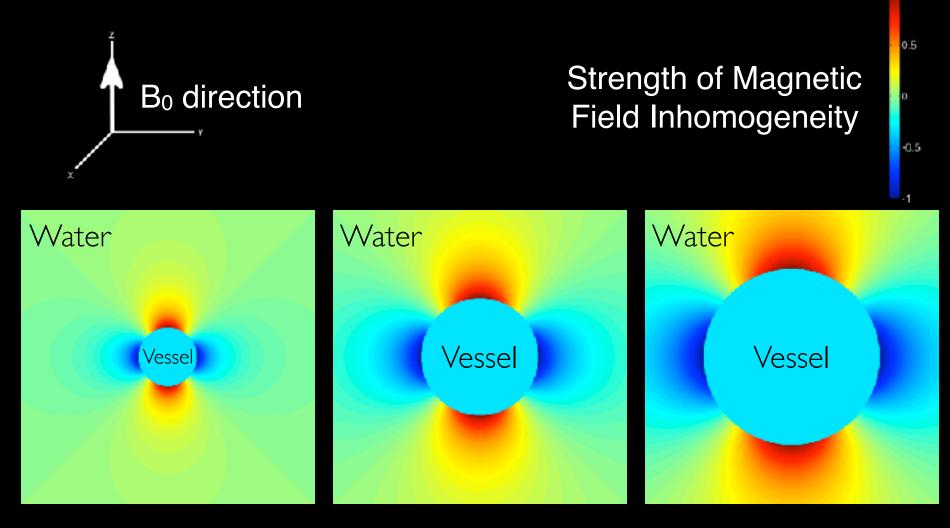
Vessels, depending on orientation and blood oxygen content will alter their local magnetic fields

BOLD Effect – vessel orientation



0°45°90°Vessel parallelVessel perpendicular

BOLD Effect – vessel size



radius = 50 μ m

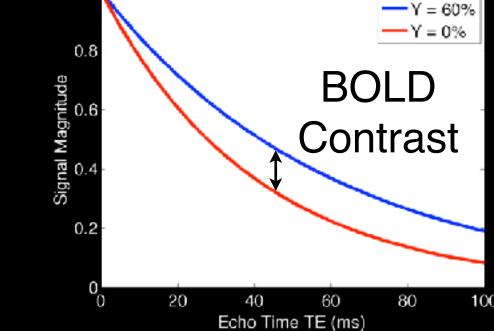
radius = 100 μ m

radius = 150 μ m

BOLD Effect – blood oxygenation level 0.5 Strength of Magnetic **B**₀ direction **Field Inhomogeneity** -0.5 Water Water Water Vessel Vessel Vessel

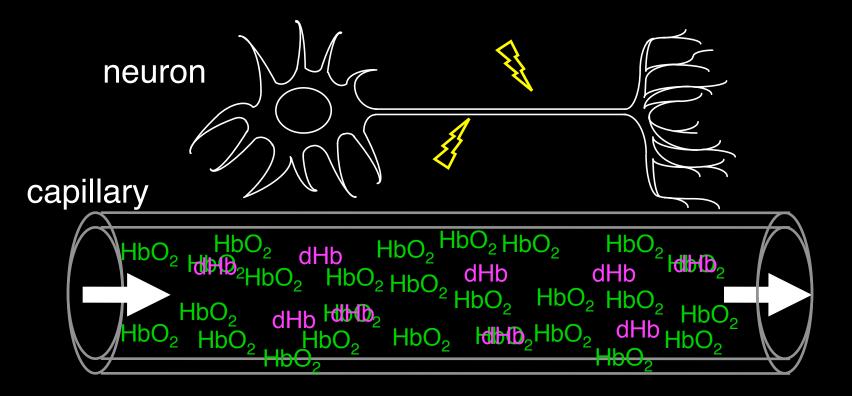
Oxygenation = 60% Oxygenation = 30% Oxygenation = 0%

Oxygenation: Y=60% More Oxygenated Hb Low inhomogeneity Longer T2* Higher signal



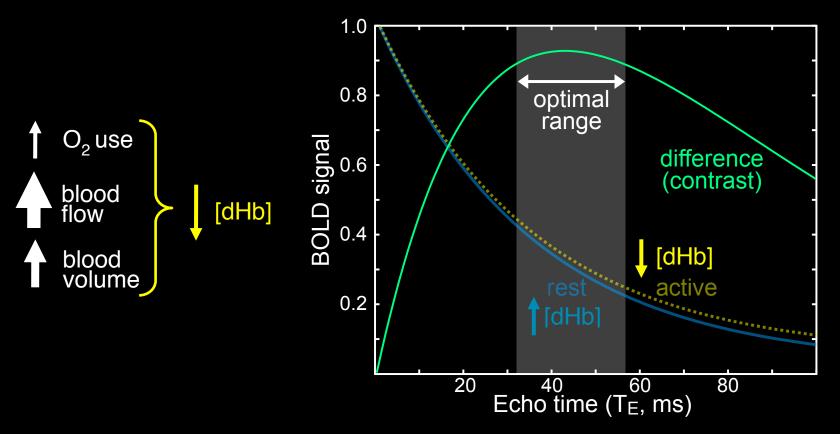
Oxygenation: Y=0% More de-oxygenated Hb **High** inhomogeneity Shorter T2* **Lower** signal

Vascular Response to Activation



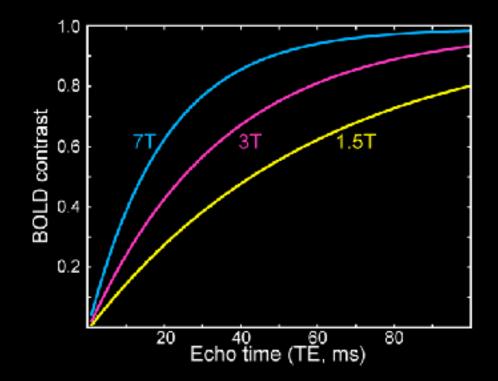
 $\begin{array}{c} & HbO_2 = oxyhemoglobin \\ & dHb \\ \hline & blood flow \\ & HbO_2 \\ \hline & blood volume \\ & HbO_2 \end{array} \end{array}$

BOLD Contrast



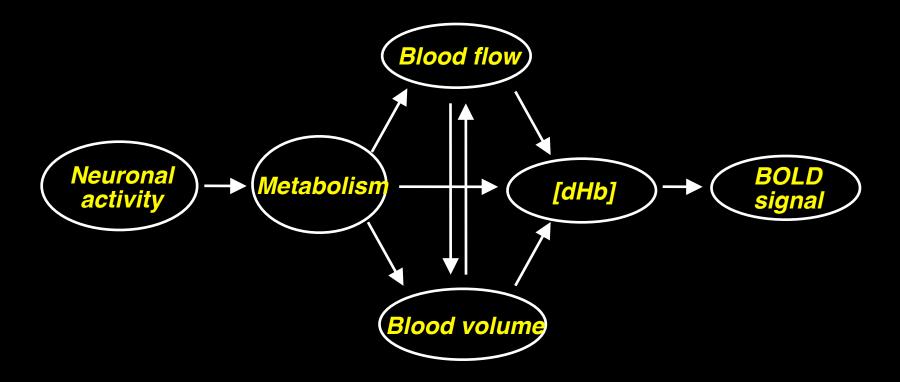
Signal increases during activation (less decay) Signal change for longer delay (T_E) Typically, 1–5% signal change

BOLD signal and field strength (B₀)



SNR and BOLD effects can increase with field strength But image artefacts get worse at higher field strength 3T is currently a good tradeoff of signal vs artefacts

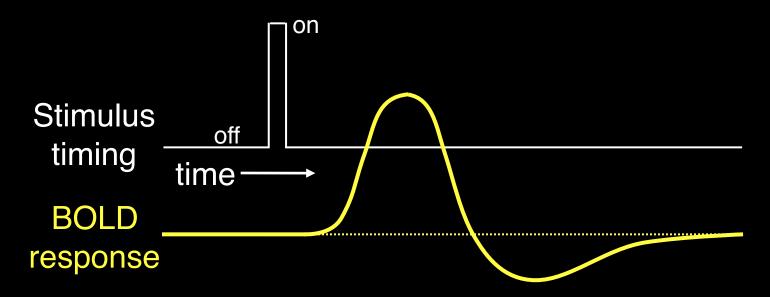
Sources of BOLD Signal



Indirect measure of activity (via metabolism!)

Subject's physiological state & pathology can change neurovascular coupling, muddying interpretation

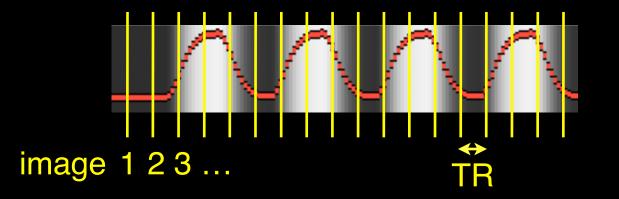
Hemodynamic response function (HRF)



Vascular response to activity is delayed & blurred Described by "hemodynamic response function"

Limits achievable temporal resolution Must be included in signal model

What is required of the scanner?



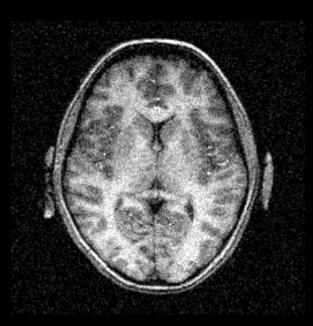
Typical stimulus lasts 1–30 s Rapid imaging: an image every few seconds Anatomical images take minutes to acquire! Acquire "single-shot" images (e.g., EPI)

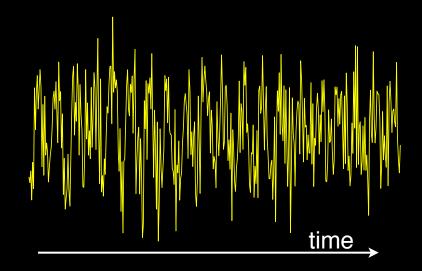
Typical* FMRI Parameters

* Typical, *not* fixed!!

Parameter	Value	Relevant points
T _E (echo time)	1.5T: 60 ms 3.0T: 30-40 ms 7.0T: 15-25 ms	Determines functional contrast, set ≈T2*
T _R (repeat time)	1–4 s	HRF blurring < 1s; Poor resolution > 4s
Matrix size / Resolution	64x64 – 96x96 2–3 mm	Limited by distortion, SNR, FOV
Scan duration	2-15 mins	Lower limit: sensitivity Upper limit: compliance

Confounds: Noise



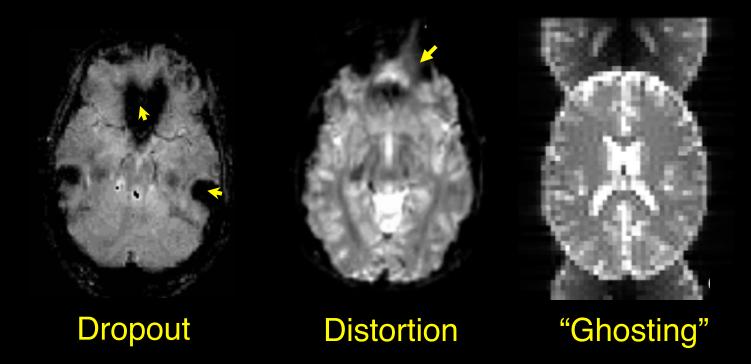


Purely random noise (example: "thermal")

Structured noise (example: "physiological")

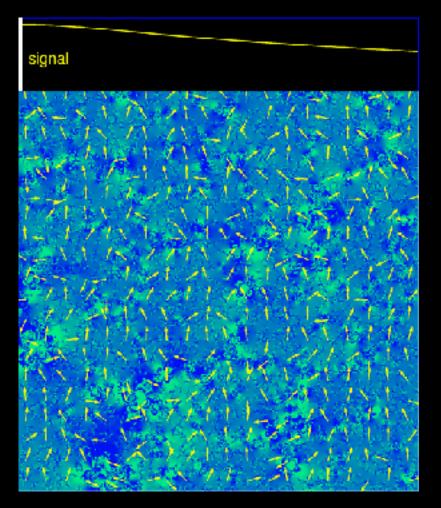
Noise: signal fluctuations leading to less robust detection with respect to statistical measures

Confounds: Artefacts



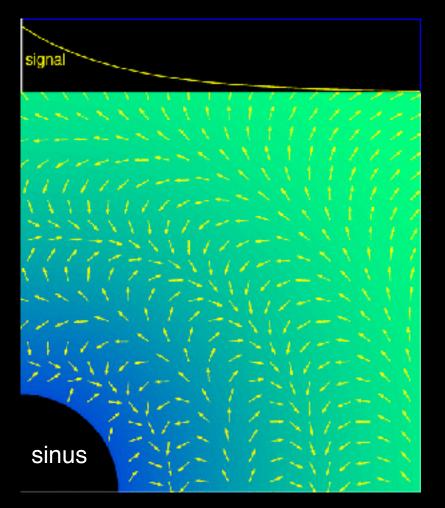
Artefacts: systematic errors that interfere with interpretability of data/images

Source of signal dropout



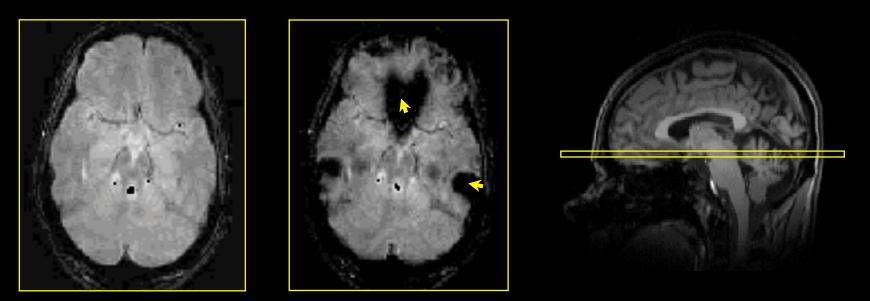
BOLD contrast is based on signal dephasing BOLD imaging requires longish delay (T_E) for contrast

Dropout is just extreme dephasing



Dephasing also occurs near air-tissue boundaries Sensitivity to BOLD means signal loss near air-tissue boundaries

BOLD Signal Dropout



Short TE

Long TE

Dephasing near air-tissue boundaries (e.g., sinuses) BOLD contrast coupled to signal loss ("black holes") Air-tissue effect is often larger than BOLD effect Dropout is not correctable post-acquisition!

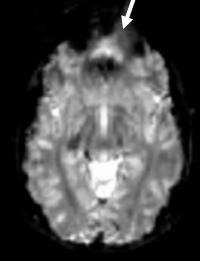
Image distortion

field offset



Field map

local warping



EPI

We think frequency maps to spatial location... So errors in frequency cause spatial mis-localization! *More on Wednesday...*

Final thoughts

Understand how different experimental parameters affect SNR and image artefacts

Tradeoffs: you can't get something for nothing, but you do have options

Get to know an engineer/physicist/radiographer: get help setting up study protocols, show them your artefacts

Quality assurance: always look at your data, even if you are running a well-tested protocol

Questions:

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