



The Principles of Quantitative MRI

Geoffrey D. Clarke
Dept. of Radiology

**University of Texas Health Science
Center at San Antonio**

Overview

- Excitation & Signal Collection Process
- Gradients for Spatial Localization
- Measuring Tissue Volumes
- Measuring NMR Properties of Tissues
- Tissue Physiology Measurements
- ~~• Tissue Biochemistry Measurements~~

Overview

- **Excitation & Signal Collection Process**
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RF Nonuniformities

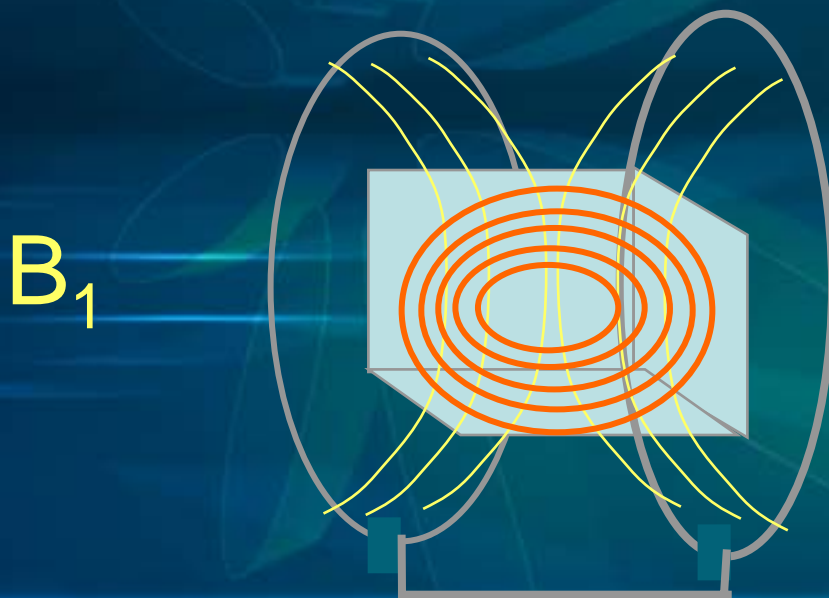
- **Radio Frequency Field Nonuniformities are the Single Biggest Cause of Errors in qMRI**
- **RF Nonuniformities Increase as the B_0 -Field Increases**
- **Dielectric Resonance Effects Become Pronounced at High B_0**

B1 TX Field

- Directly related to current in TX coil
 - Depends on Q of coil & coil loading
- Depends on TX Coil Geometry
- TX power auto-adjusted (pre-scan)
 - Values should be know to 1%
 - $1\% = 0.086 \text{ dB}$
- TX nonlinearities
- RF pulse droop

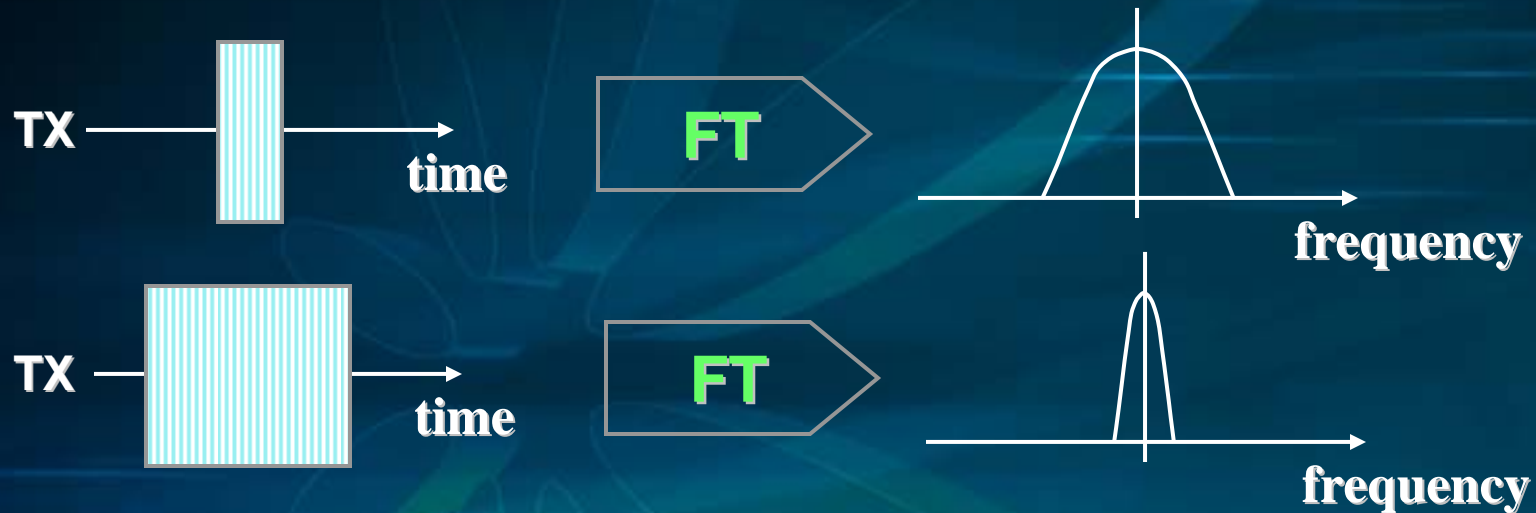
NMR Signal

$$\delta\nu(t) = \omega_0 B_{1x,y} M_{x,y} \delta V_s \cos(\omega_0 t)$$



RF Pulse Bandwidth (BW)

BW is inversely proportional to RF pulse duration:

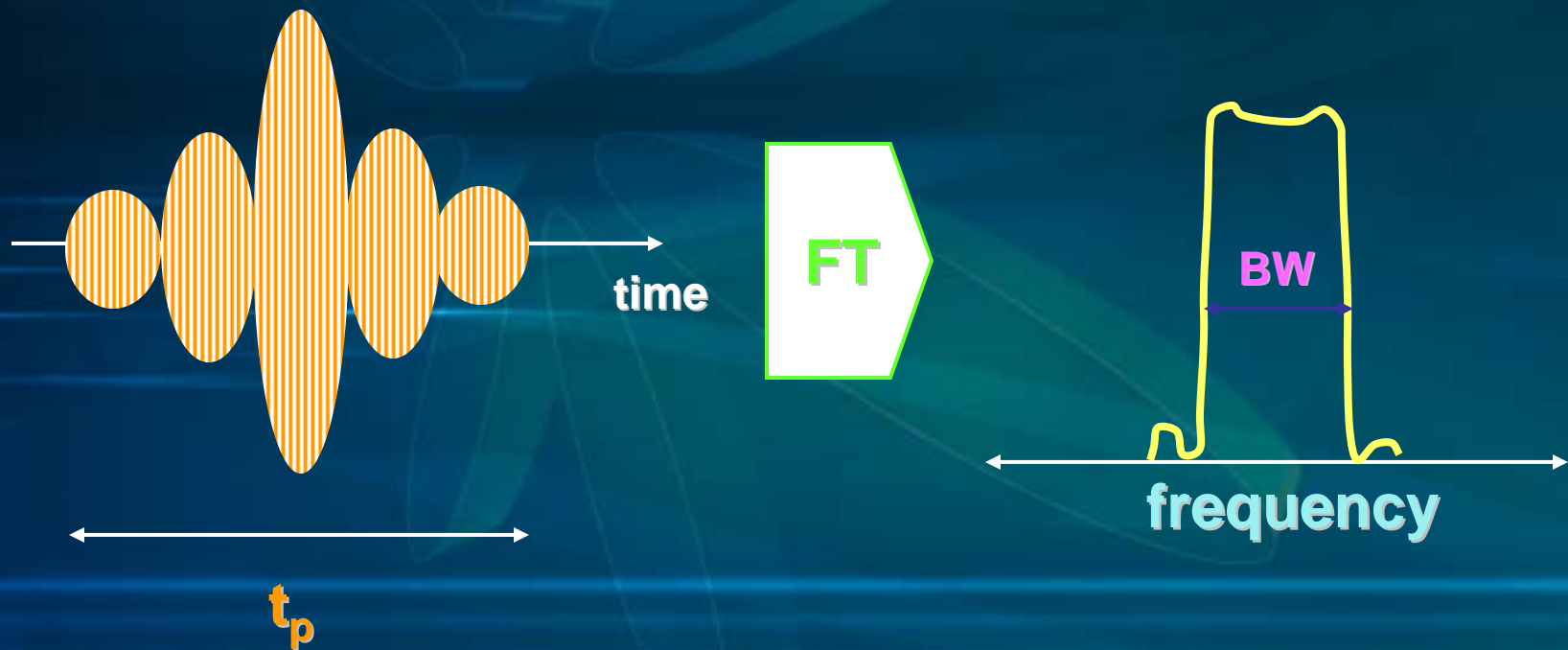


Slice select gradient is scaled based on BW & G_{ss} :

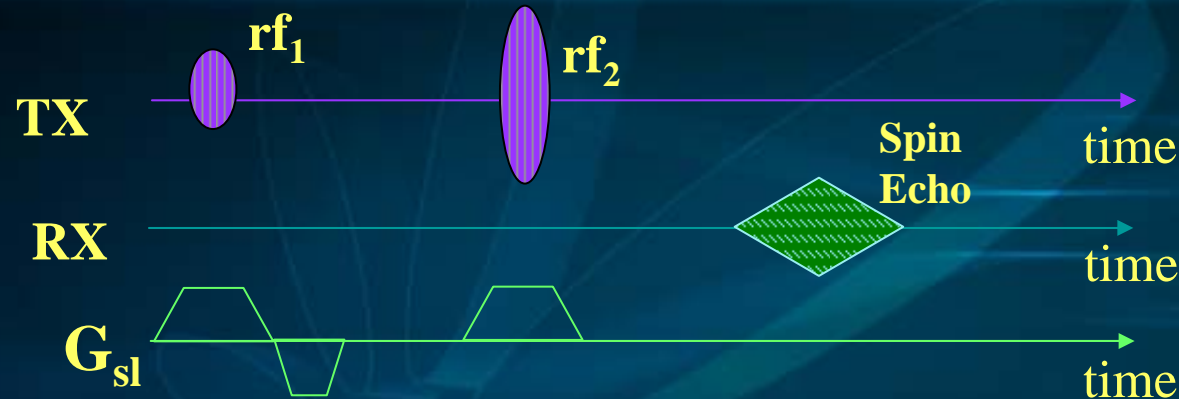
$$\text{slice thickness} = \frac{2\pi}{\gamma} \cdot \frac{BW}{G_{ss}}$$

FT Approximation

A *sinc* function ($\frac{\sin x}{x}$) envelope on the r.f. pulse produces a nearly square excitation profile of the phantom.....

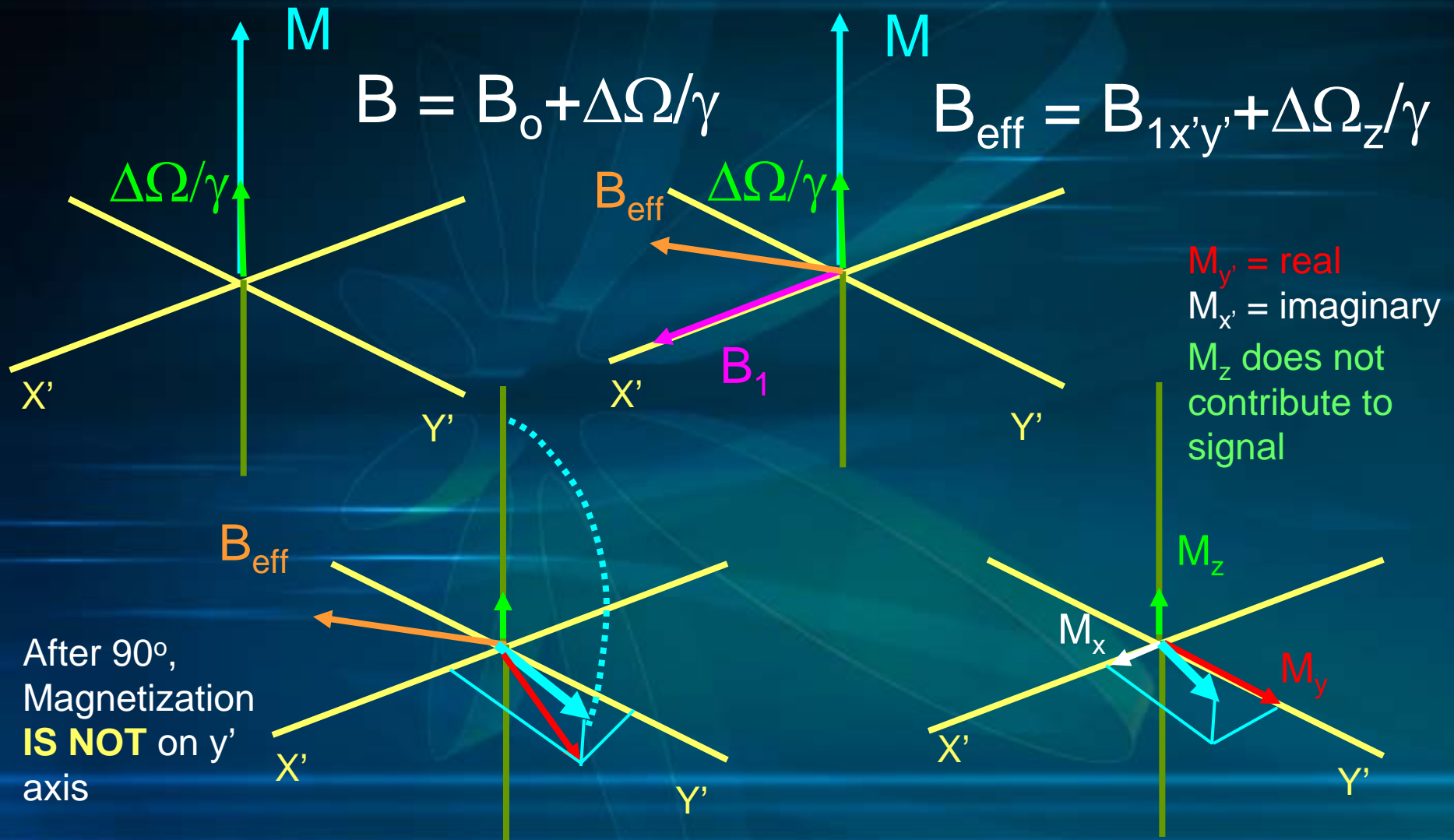


Slice Selection

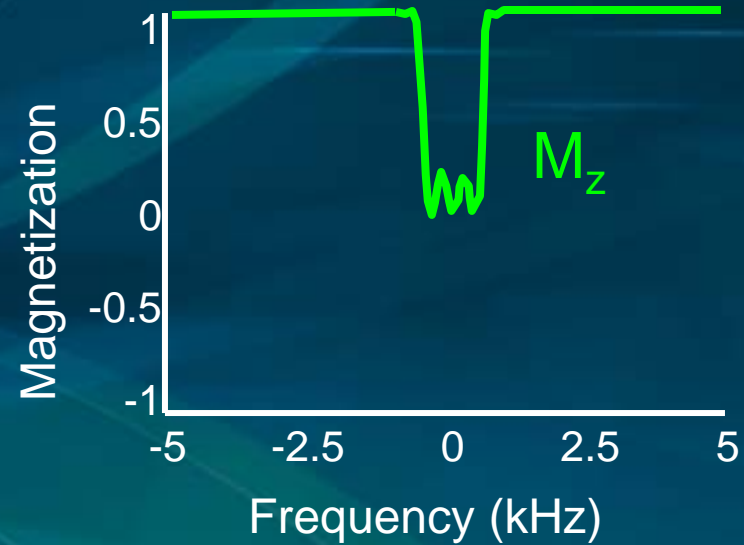
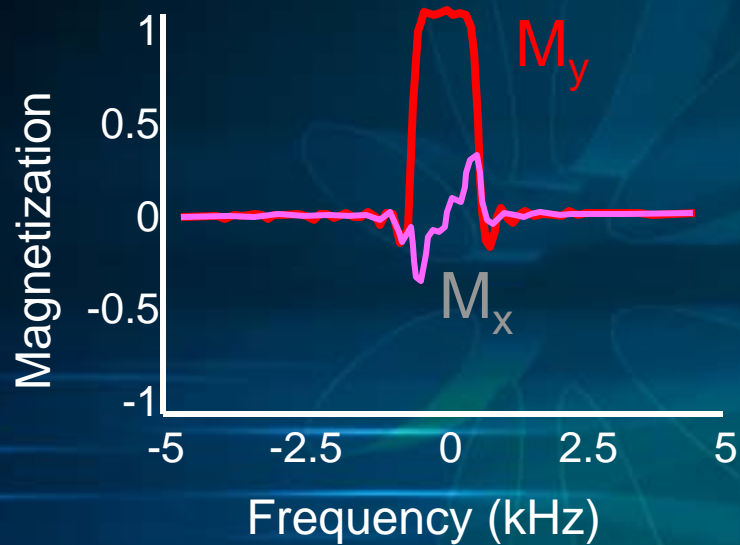


- If constant gradient field is on during the rf pulse:
 - Larmor frequency of spins varies with position
 - The flip angle depends on the local Larmor frequency and the frequency content of the RF field pulse
 - the RF pulse can be “crafted” to contain frequencies in only a specified range

Resonant Frequency Offset

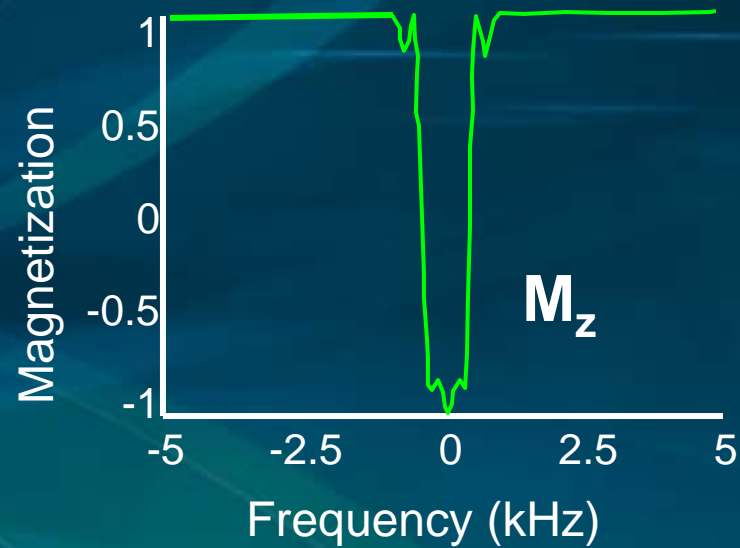
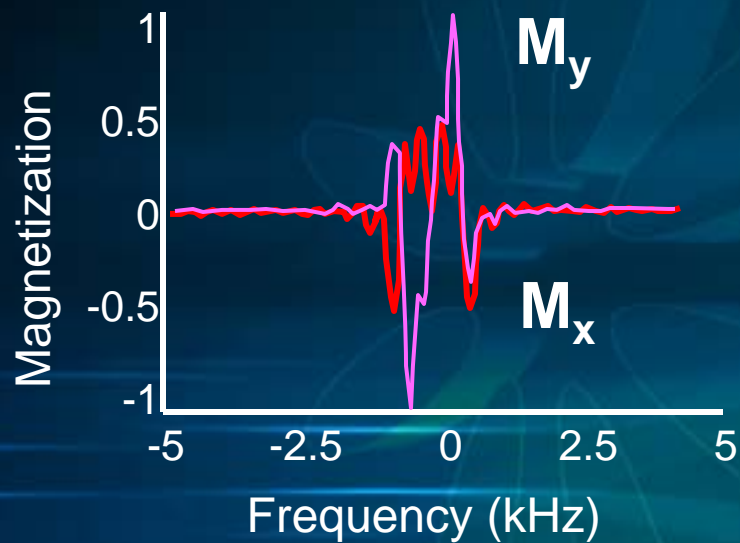


90° Sinc Pulse Profile



2 ms, 5-lobe, chemical shift refocused

180° Sinc Inversion Pulse



2 ms, 5-lobe, chemical shift refocused

Slice Profile Variations

- Flip Angle varies with location
 - Due to B_1 , B_0 field nonuniformities
- Non-linearity of Excitation (Bloch Eqns)
 - FT approximation invalid for big flip angles
 - Bloch simulator software

<http://www-mrsrl.stanford.edu/~brian/mritools.html> *

- T1-weighting of excitation profile

Bloch Equations

- A set of simultaneous differential equations that describe the behavior of the magnetization under any conditions.

$$\frac{dM_z(t)}{dt} = \gamma[M_x(t)B_y(t) - M_y(t)B_x(t)] - \frac{M_z(t) - M_0}{T_1}$$

Magnetization
along the z-axis

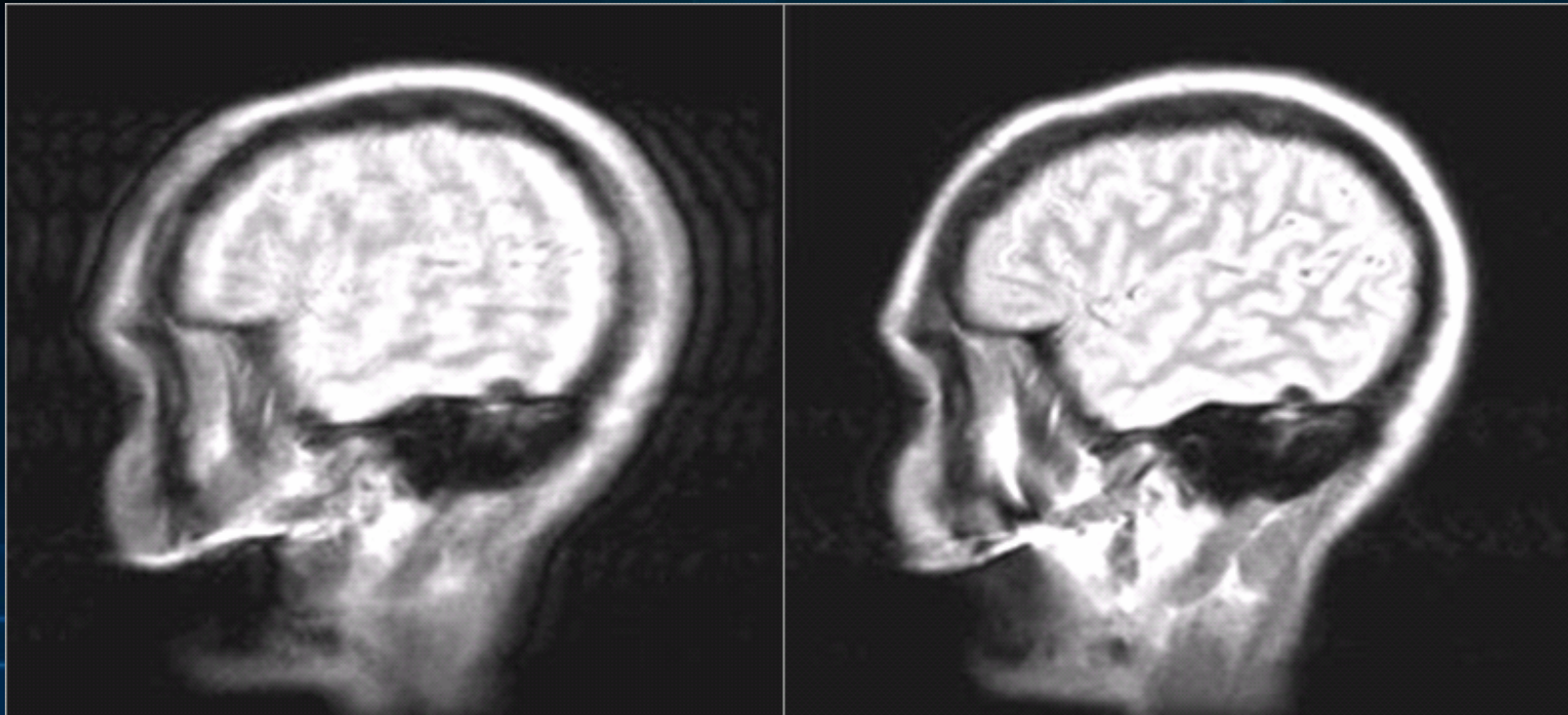
$$\frac{dM_x(t)}{dt} = \gamma[M_y(t)B_z(t) - M_z(t)B_y(t)] - \frac{M_x(t)}{T_2}$$

Magnetization
along the x-axis

$$\frac{dM_y(t)}{dt} = \gamma[M_z(t)B_x(t) - M_x(t)B_z(t)] - \frac{M_y(t)}{T_2}$$

Magnetization
along the y-axis

Poor RF Pulse Calibration



Miscalibration of FSE 180° RF pulses (left image) is corrected (right image)

B1 Field Mapping - Purpose

- a. Needed for accurate measurement of many NMR parameters, i.e. relaxation times**
- b. Enables estimation of systematic errors in parameter measurement**
- c. Enable correction of spatial sensitivity variation using reciprocity**

B1 Field Mapping - Methods

a. One-pulse read $M_{x,y}$

Venkatsen et al. Magn Reson Med 1998; 40:592

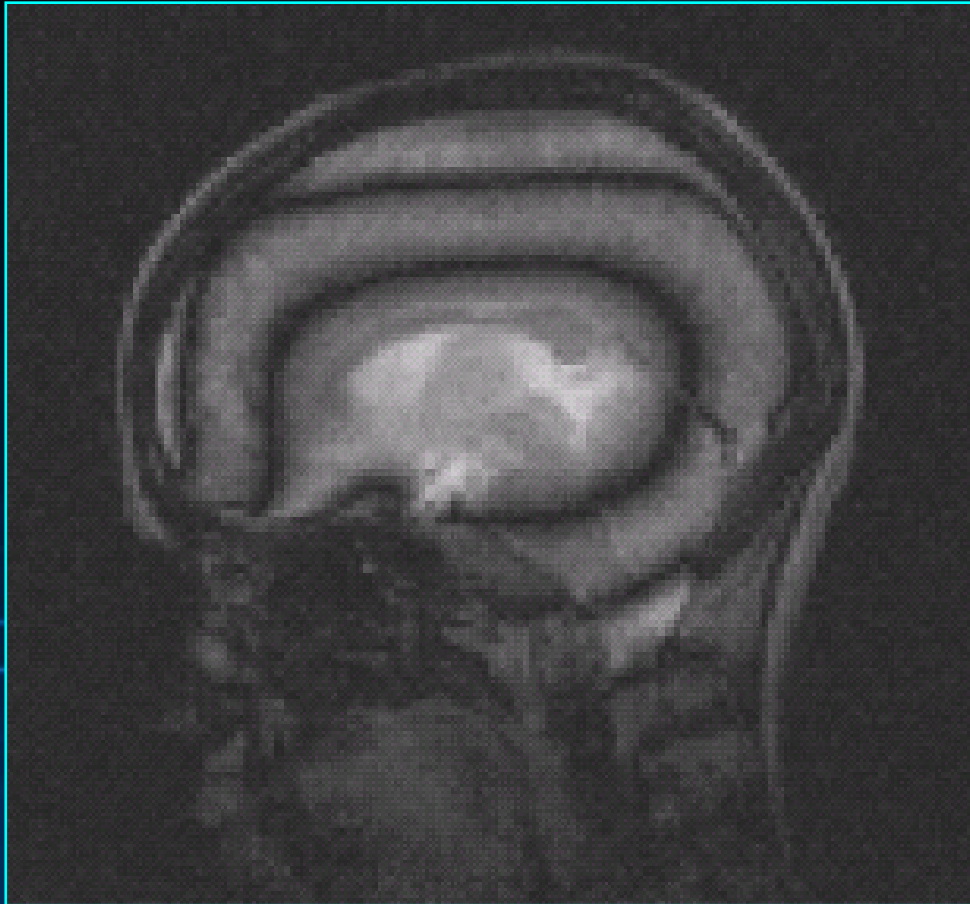
b. Spin Echo (both pulses altered)

Barker et al. BJR 1998; 71: 59-67

c. One-pulse read M_z

Vaughn et al. Magn Reson Med 2001; 46: 24

B1 Field Mapping

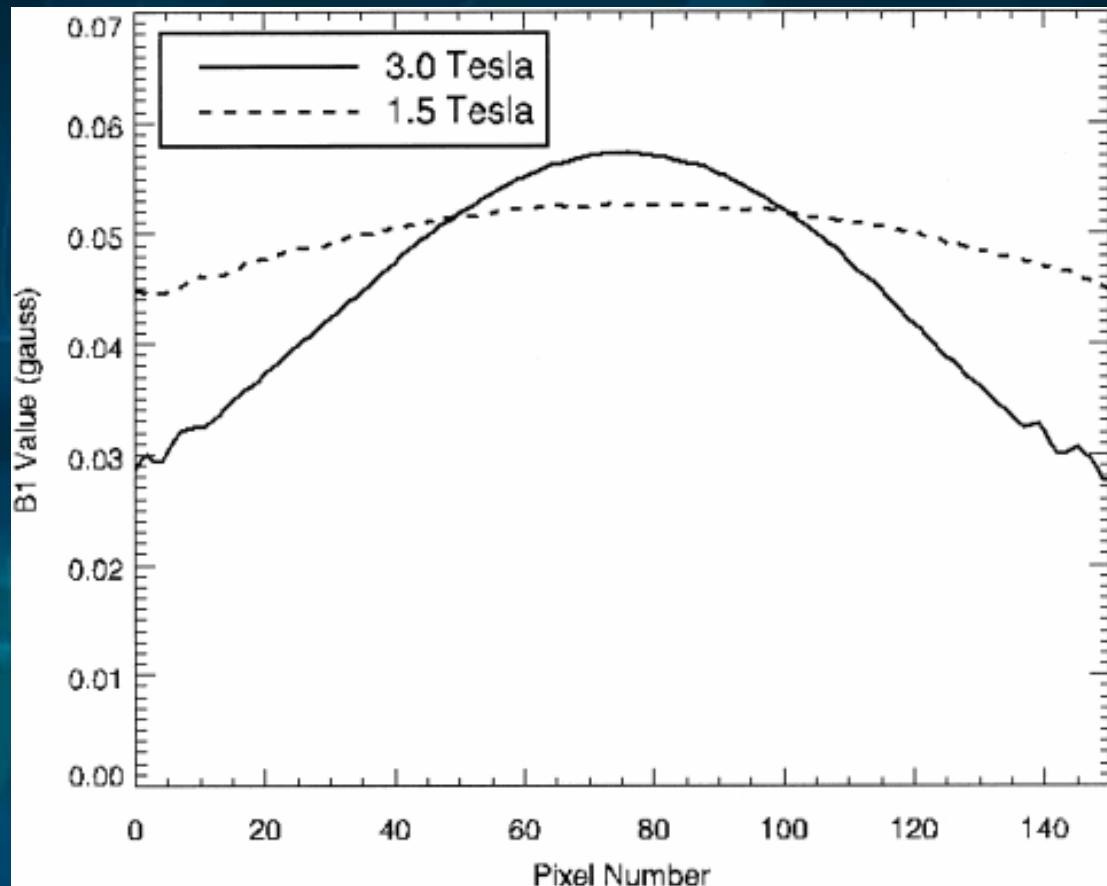
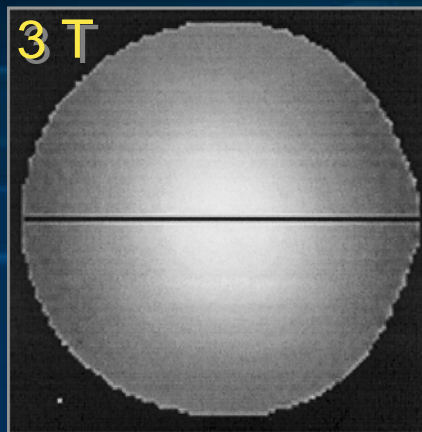
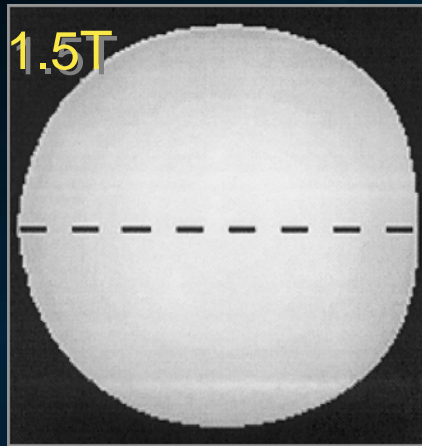


- One-pulse $M_{x,y}$ method
- Hard 180° pulse preceding 2D field echo sequence
- Bright center is maximum B_1
- Ring pattern occurs at every 5% change in B_1 -field

Dielectric Resonance Effect

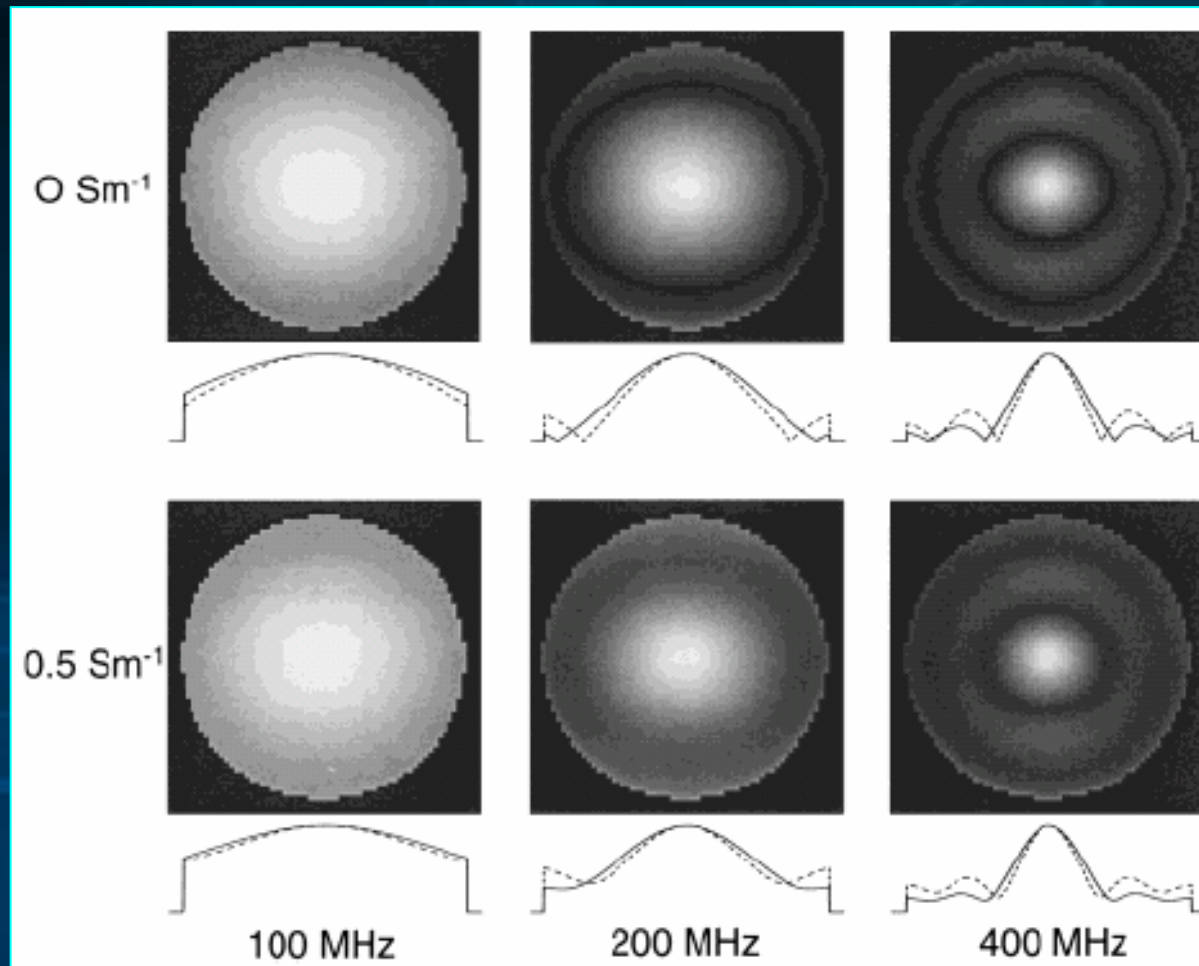
- Fields inside a sample of diameter, d , can be resonant when $d = n\lambda$
- This can make the B1 field at the center of the sample larger than at the edges
- This effect is dampened as the conductivity of the sample increases

Image Uniformity at 3 Tesla



- B1 field maps in a saline phantom (18 cm diameter)

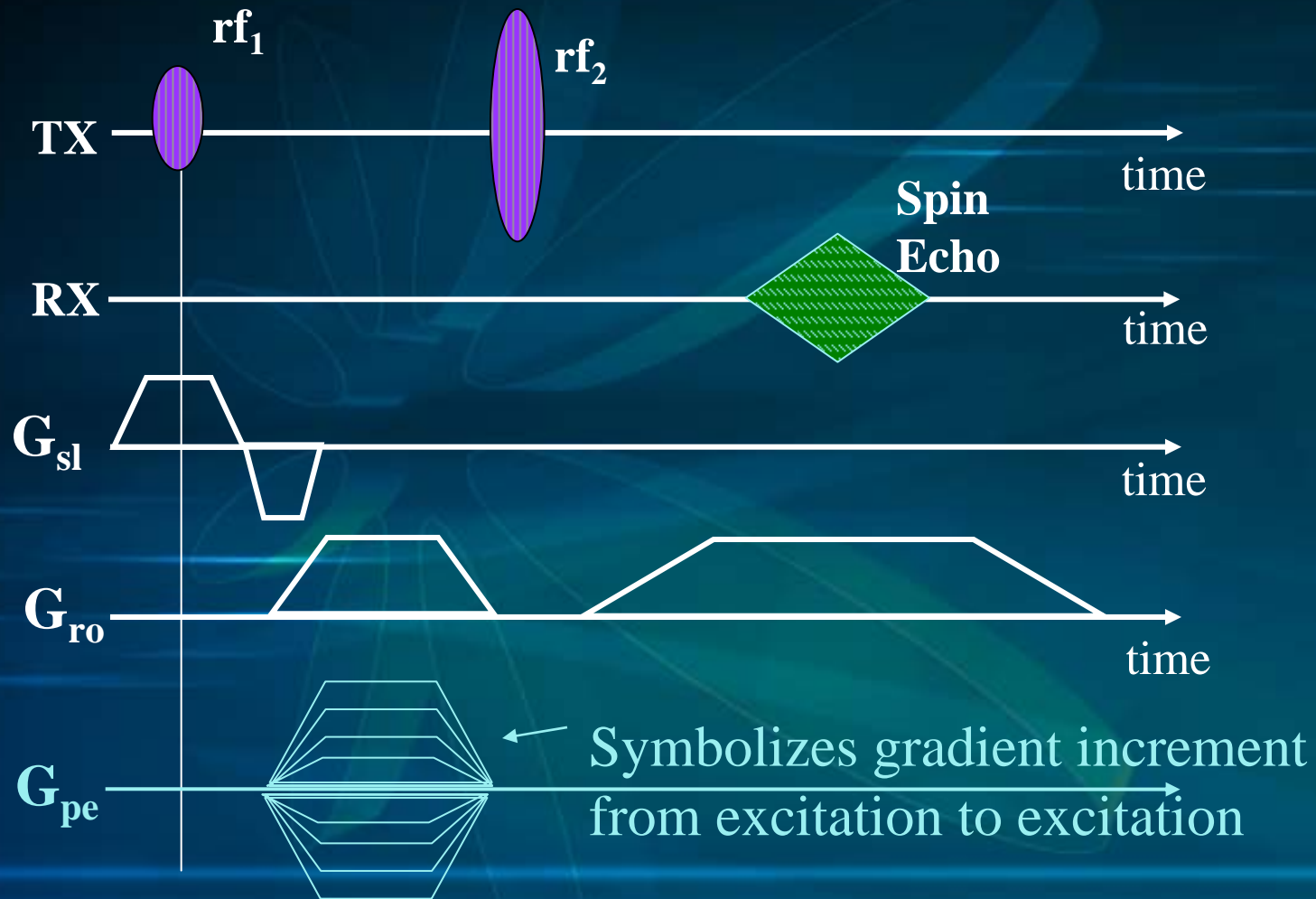
Dielectric Resonance



Overview

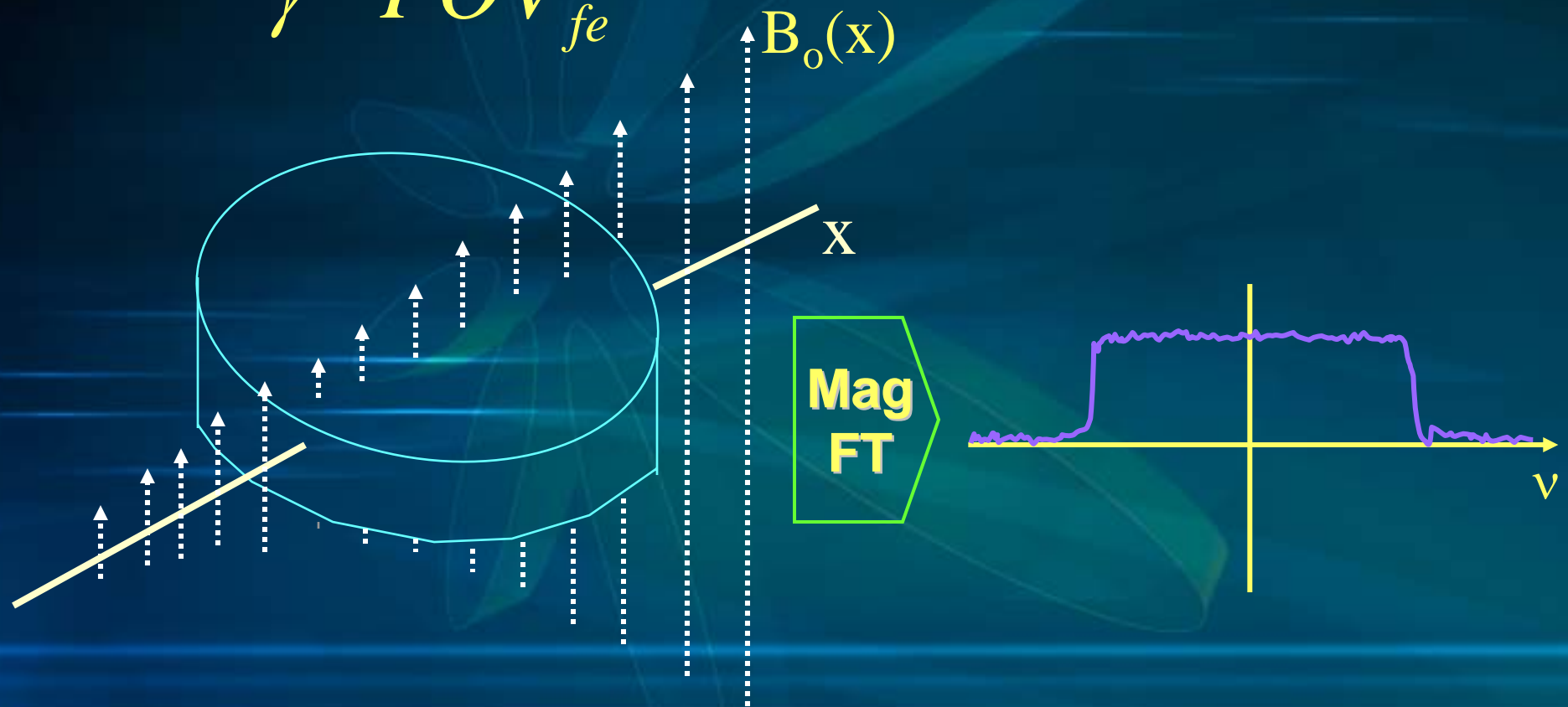
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Spin-Echo Sequence

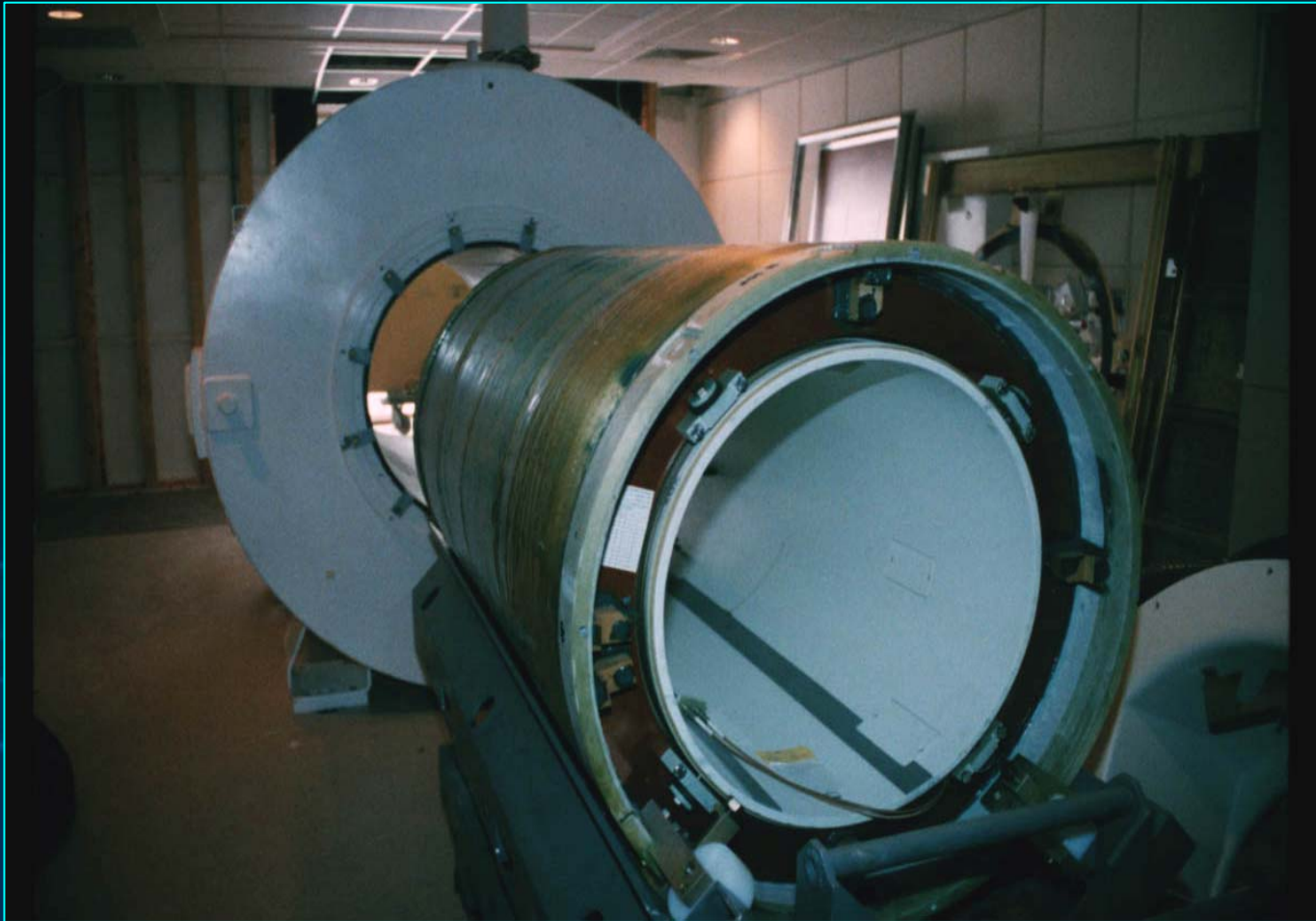


One Dimensional FT

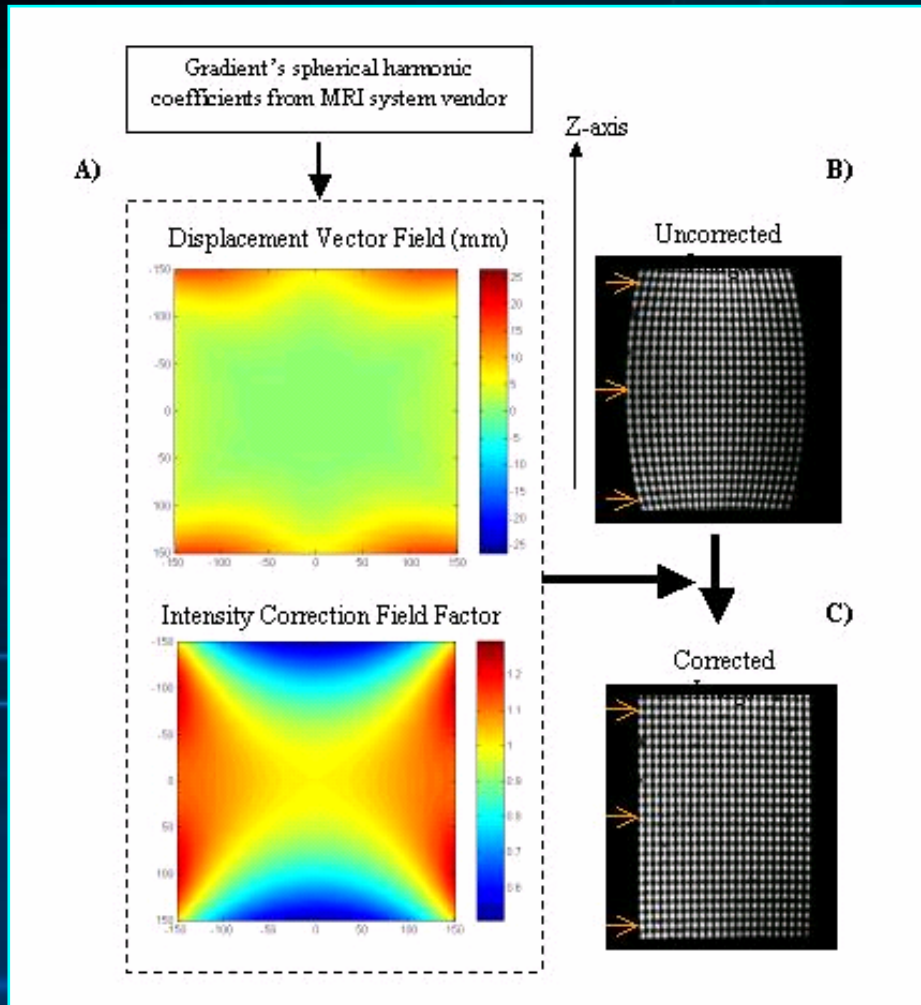
$$G_{fe} = \frac{2\pi BW}{\gamma FOV_{fe}}$$



Gradient Subsystems



MRI Gradient Fields



Gradient Nonlinearities are often tolerated as part of trade-offs with gradient field strength or coil size

Manufacturers often apply gradient distortion corrections in order to make images appear to be distortion free

Influences image quality parameters (SNR, spatial resolution, etc.)

MRI Gradient Fields

Sources of Error:

- Gradient Amplitude Calibration
 - Best about 1% error
- Gradient Non-linearities
- Eddy Currents

Eddy Currents

- Accelerating current in gradient coils (gradient pulse) causes induced currents in nearby metallic structures.
- These currents produce magnetic fields which, in turn, oppose the magnetic fields of the gradient coils

Eddy Currents

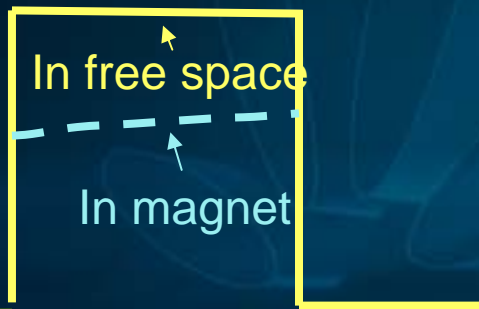
- The magnetic field produced by Eddy Currents have two time-dependent components:
 - An offset of the B_0 field
 - An additional gradient field

$$B_{ec}(\vec{r}, t) = \Delta g(\vec{r}, t) + \Delta B_0(t)$$

Gradient Waveforms

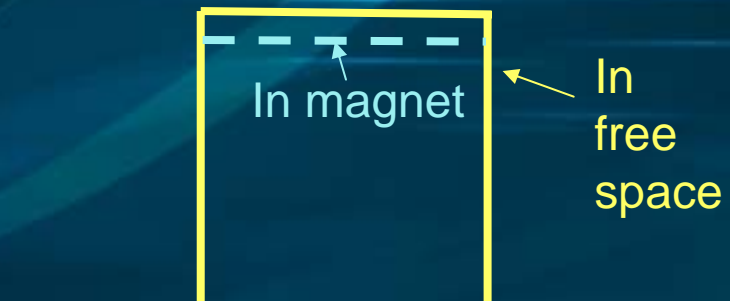
Gradient to Magnet Ratio

$G/R=0.8$

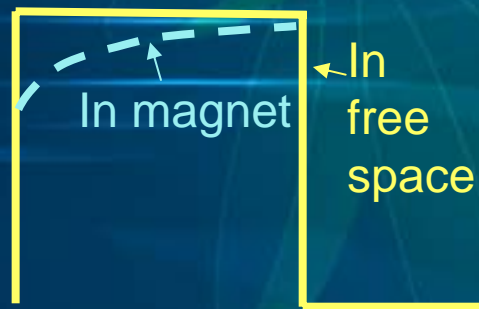


A. Whole body system.

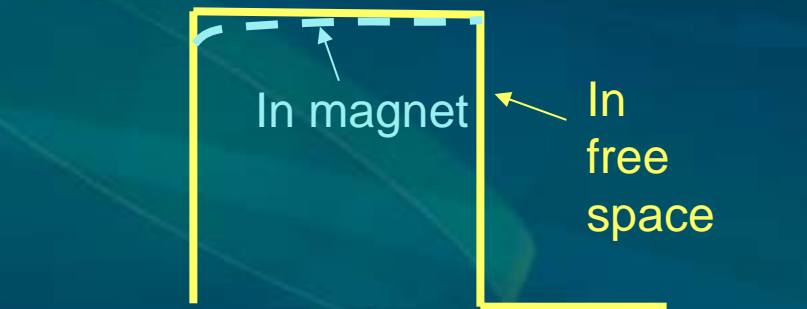
$G/R=0.6$



B. Whole body, $G/R=0.6$

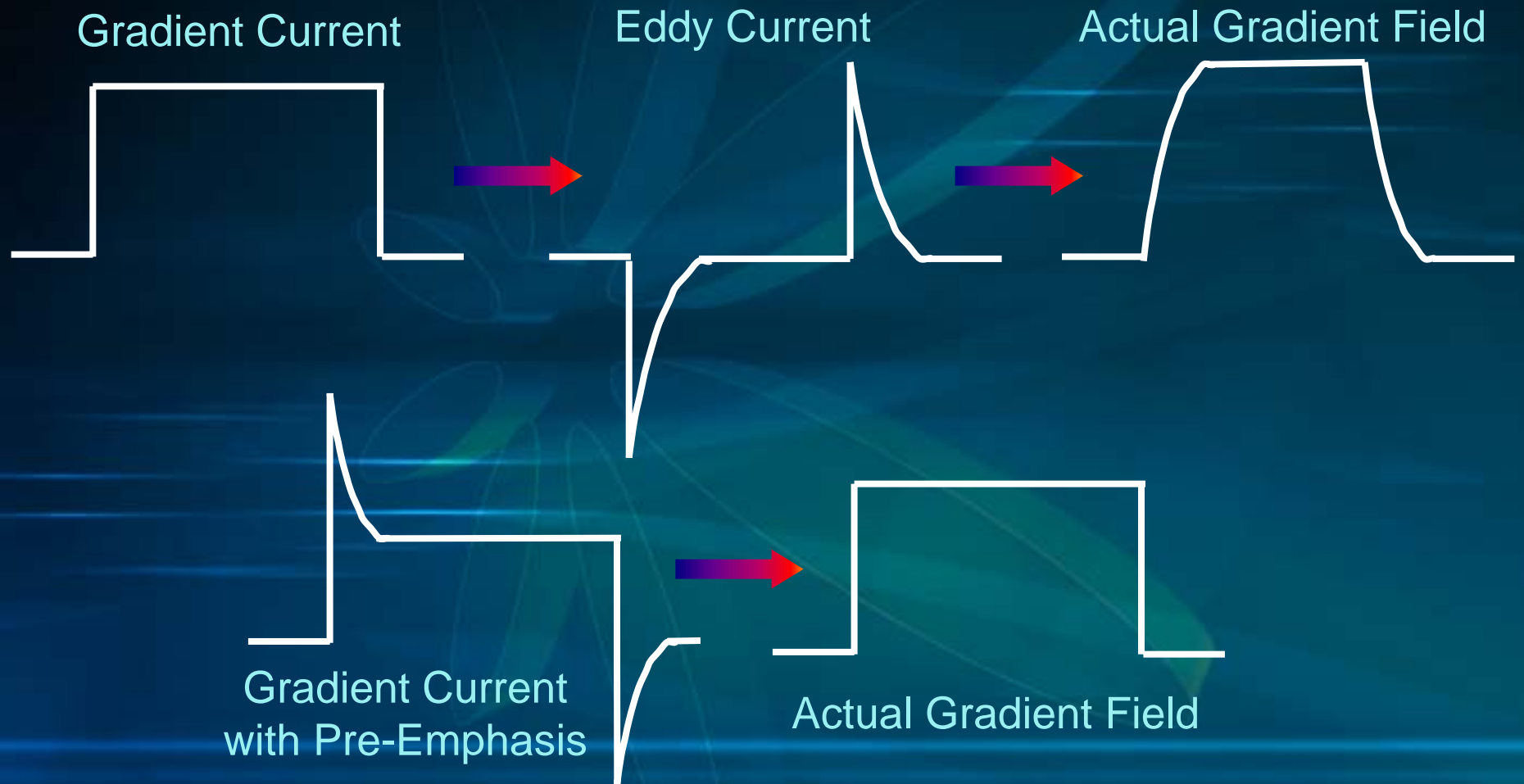


A. Microscopy system.



B. Microscopy system

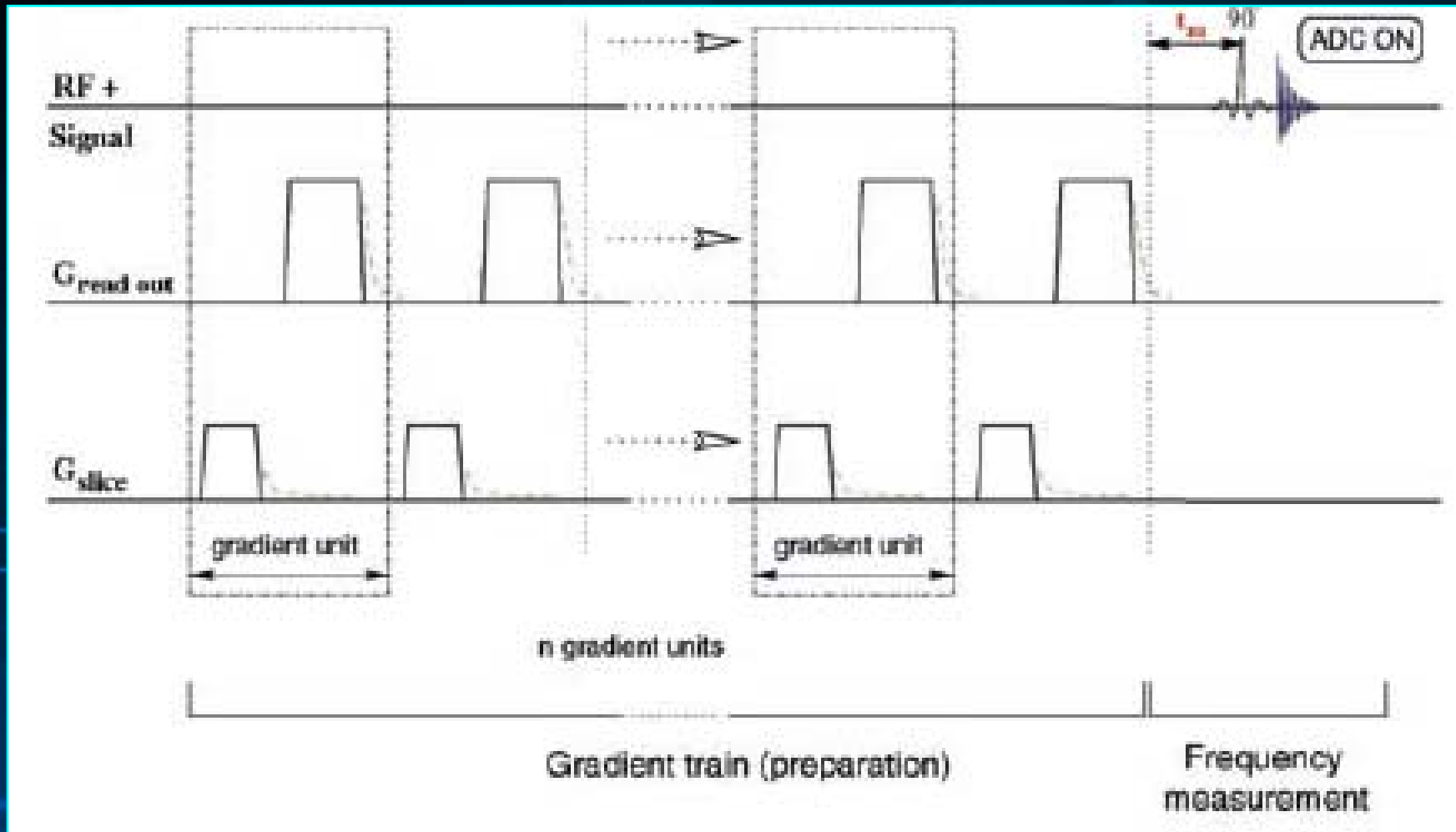
Eddy Current Preemphasis



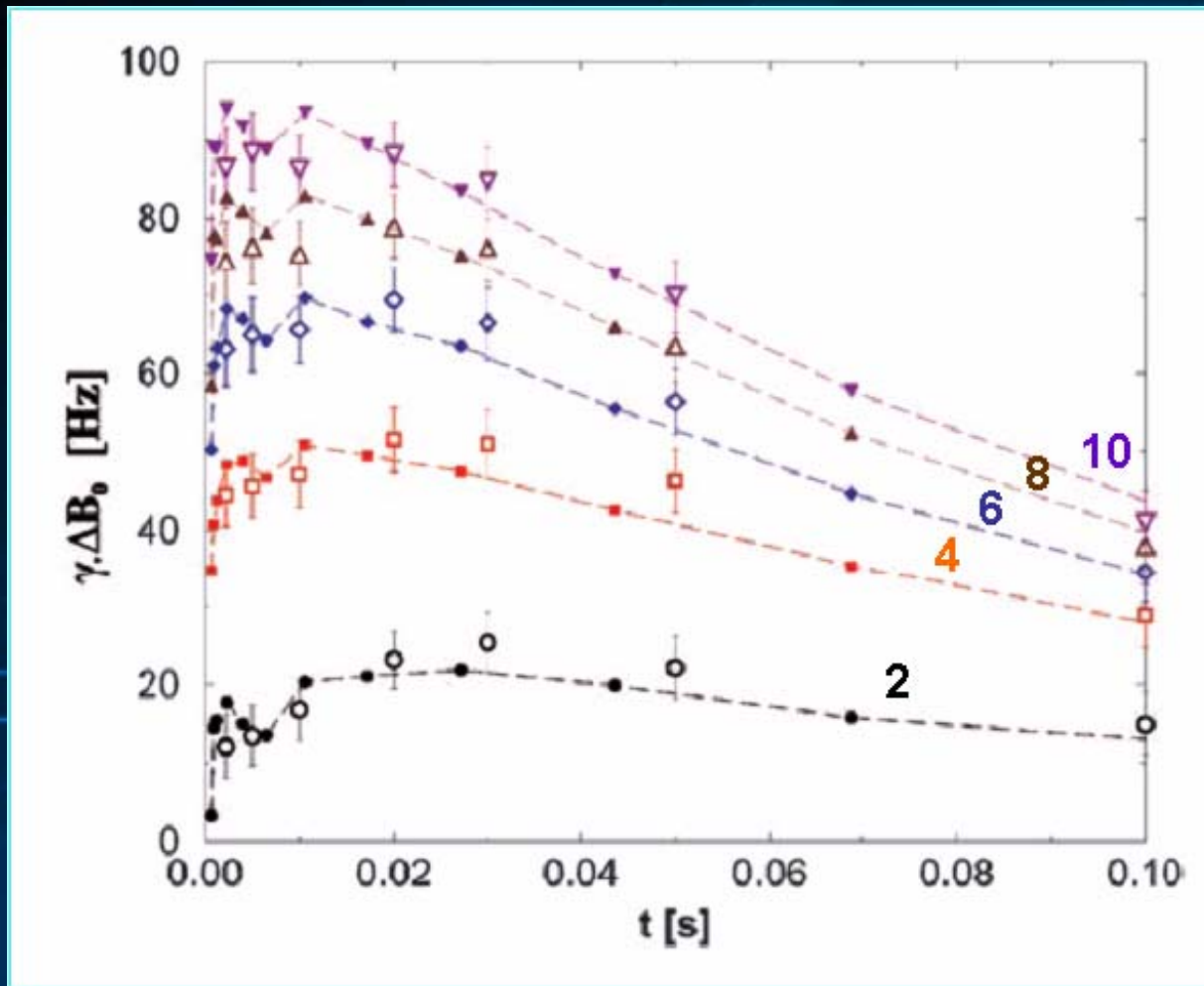
Actively Screened Gradients

- Reduce gradient field strength outside of gradient coil former
 - Current in shield is opposite polarity
- Reduces gradient field in imaging volume also
 - Improves magnet homogeneity
- Each gradient coil is associated with a screen coil
 - Twice as many amplifiers required

Measuring Eddy Currents

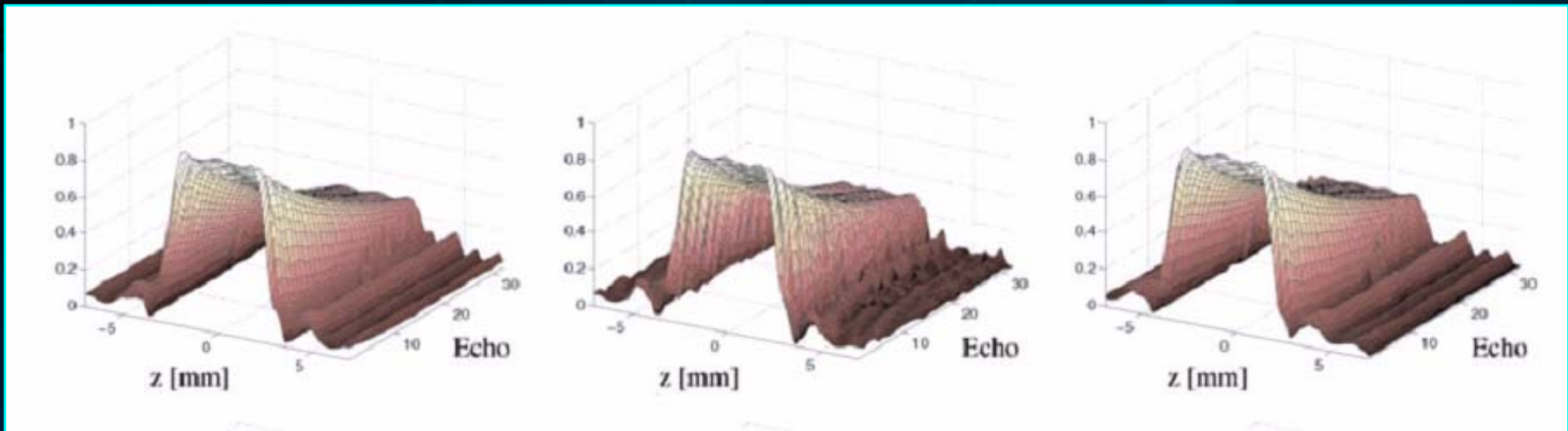


Eddy Currents



Time course
of eddy
current field
offset
following
different
numbers of
gradient pulse
units

Eddy Current Effects on Slice



Ideal – no eddy currents

worst case

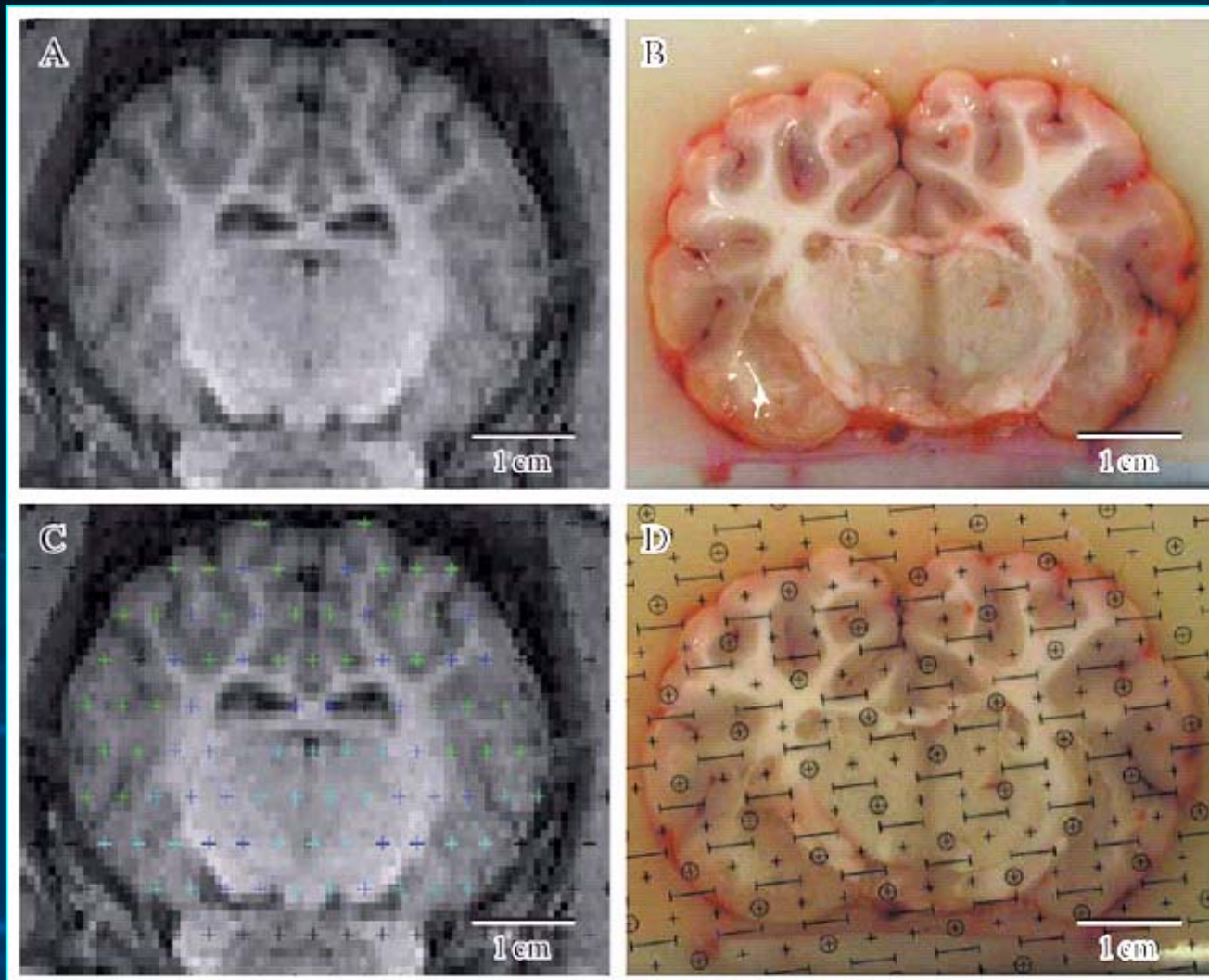
pre-gradient train

RF pulse profiles used for T2-relaxometry a) Ideal profile calculated from Bloch equations b) profile showing the influence of eddy currents; c) pre-gradient pulse train establishes steady-state which regularizes the RF pulse profile

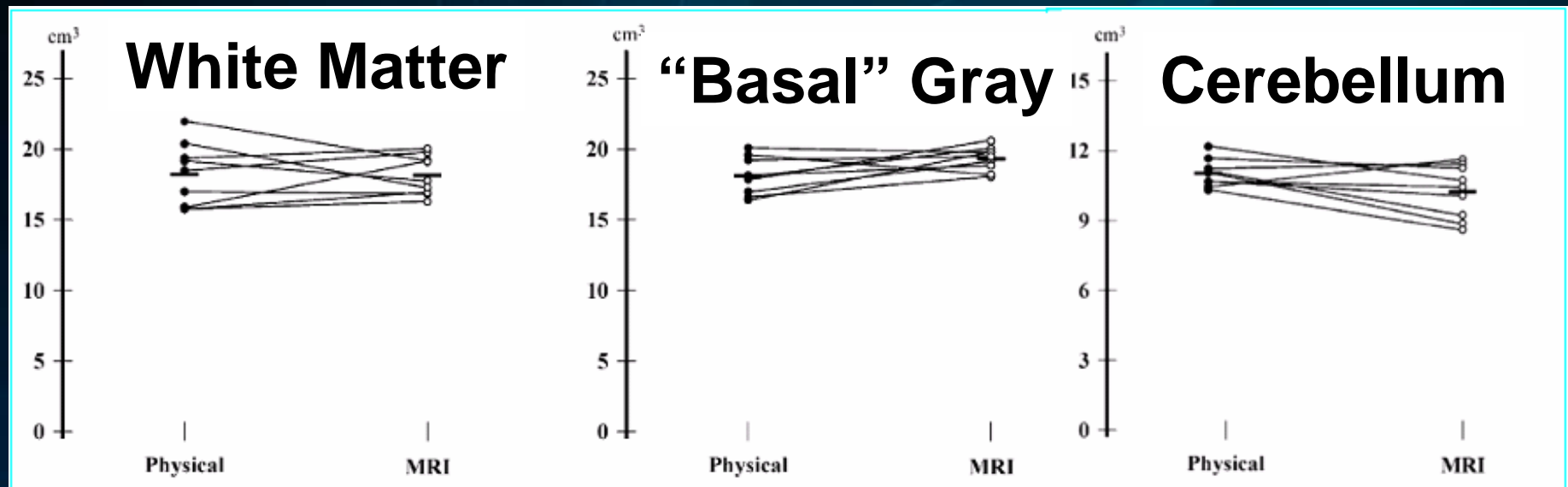
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Brain Volume Measurement

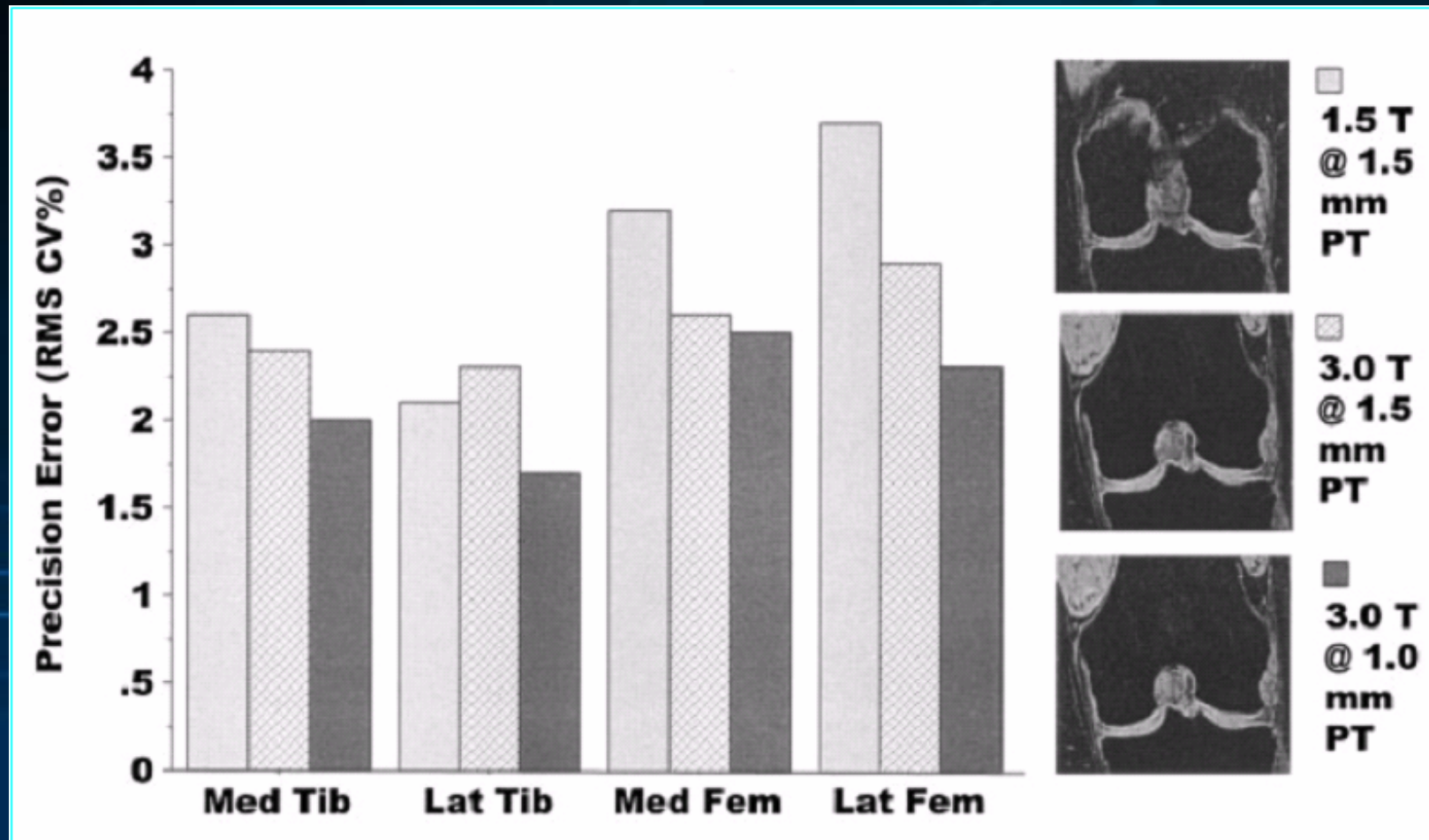


Brain Volume Relationships



- High agreement between qMRI volumetry and physical sections
- qMRI volumetry is susceptible to high inter-observer variability
- Problems greatest in those regions where tissue margins are poorly defined

Cartilage MRI

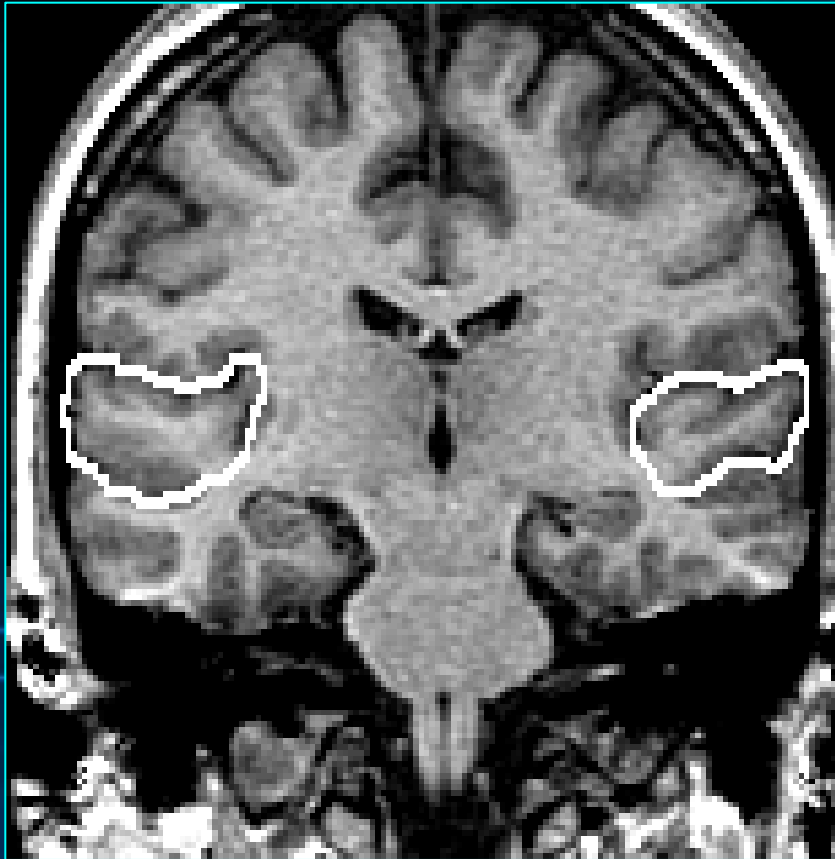


Accuracy of 3T high and tends to be more reproducible than 1.5 T

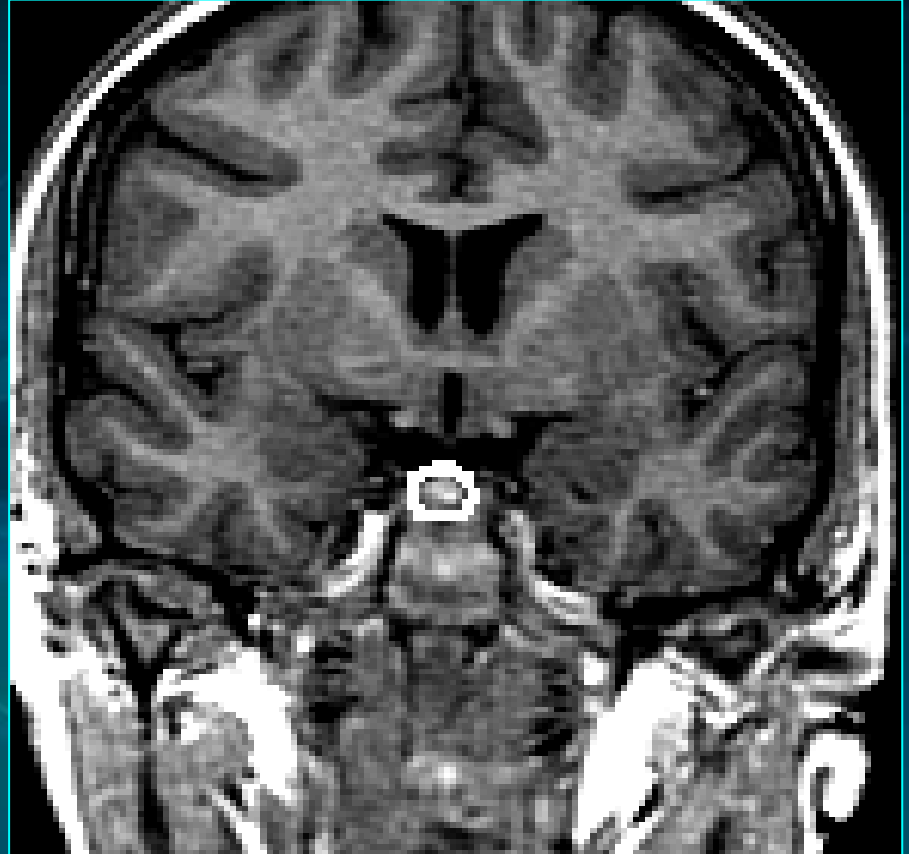
MRI in Psychiatry

- MRI is a safe and non-invasive technique to look into the to measure the human brain's normal anatomy *in vivo*
- Used in the investigation of the pathophysiology of mood disorders:
 - major depression
 - bipolar disorder
- Potential for the evaluation of psychiatric therapeutic responses

MRI Volume Measurements



Superior temporal gyrus tracing



Pituitary gland tracing

Volumes in Bipolar Disease

	Bipolar	Healthy Controls	F	p
Total left STG volume (cm ³)	12.5±1.5	13.6±2.5	4.45	0.043
White matter left STG volume (cm ³)	2.79±0.56	3.12±0.73	4.23	0.048
Gray matter left STG volume (cm ³)	9.7±1.3	10.5±2.1	2.36	0.134
Total right STG volume (cm ³)	14.9±2.3	15.3±1.9	0.57	0.454
White matter right STG volume (cm ³)	4.60±0.95	4.99±0.98	4.85	0.035
Gray matter right STG volume (cm ³)	10.3±1.8	10.3±1.5	0.01	0.930

Overview

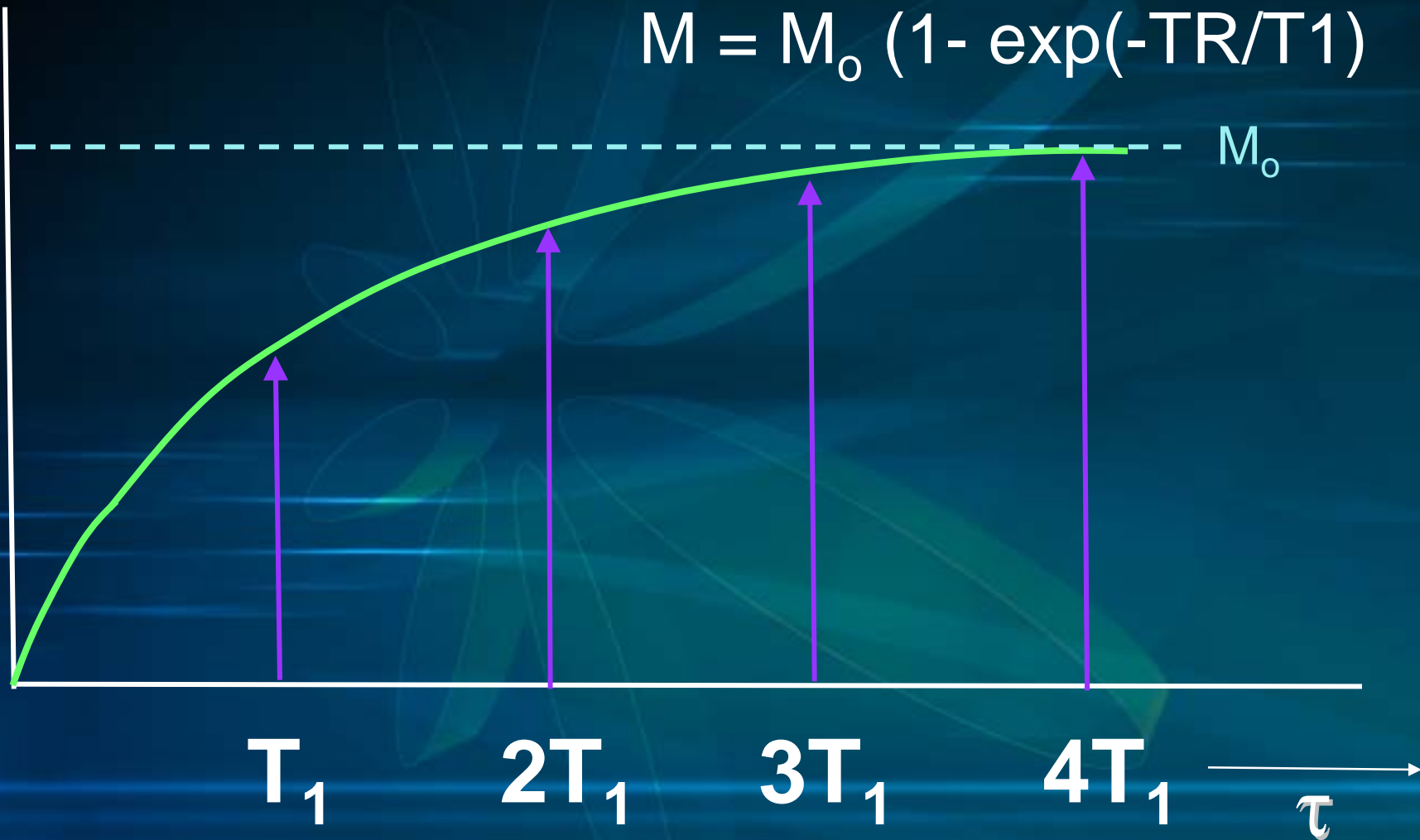
- Review of Signal Collection Process
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Relaxation Times

- T_1 : longitudinal relaxation time defines recovery of potential for next signal ($T_1=1/R_1$)
- T_2 : transverse relaxation time defines rate of dephasing of MRI signal due to microscopic processes ($T_2=1/R_2$)
- T_2^* : transverse relaxation time with B_0 inhomogeneity effects added; defines rate of dephasing of MRI signal due to macroscopic and microscopic processes ($T_2^*=1/R_2^*$)

Longitudinal Relaxation

$$M = M_0 (1 - \exp(-TR/T_1))$$



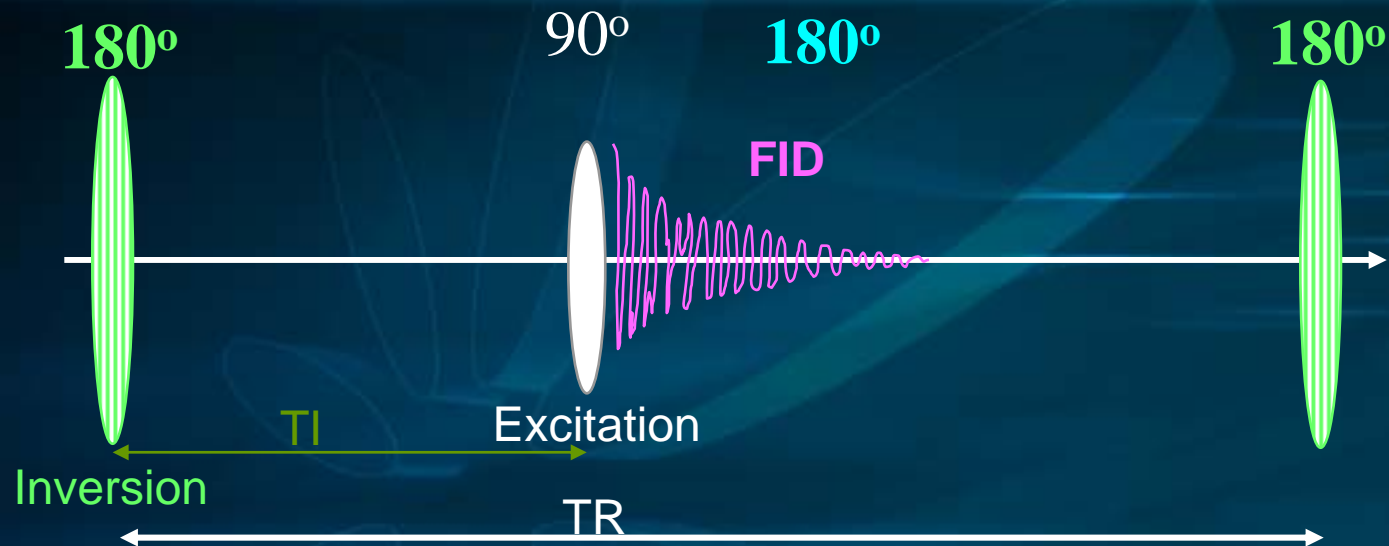
Applications for T1 Images

- Tissue characterization
- Contrast agent uptake studies
- Measurement of Tissue Perfusion
- Measurement of Blood Volume

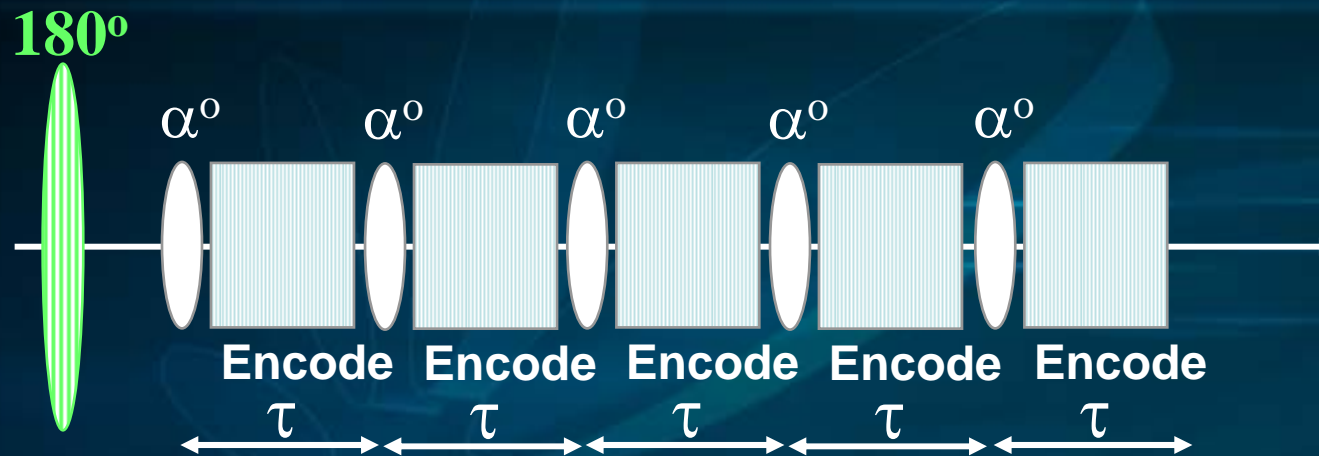
T1 Measurement Sequences

- **Inversion Recovery**
 - 180° - 90° - 180° ; the gold standard
- Saturation Recovery – 90° - 180°
- Stimulated Echo – 90° - 90° - 90°
- **Look-Locker Sequence** (see below)

Inversion Recovery



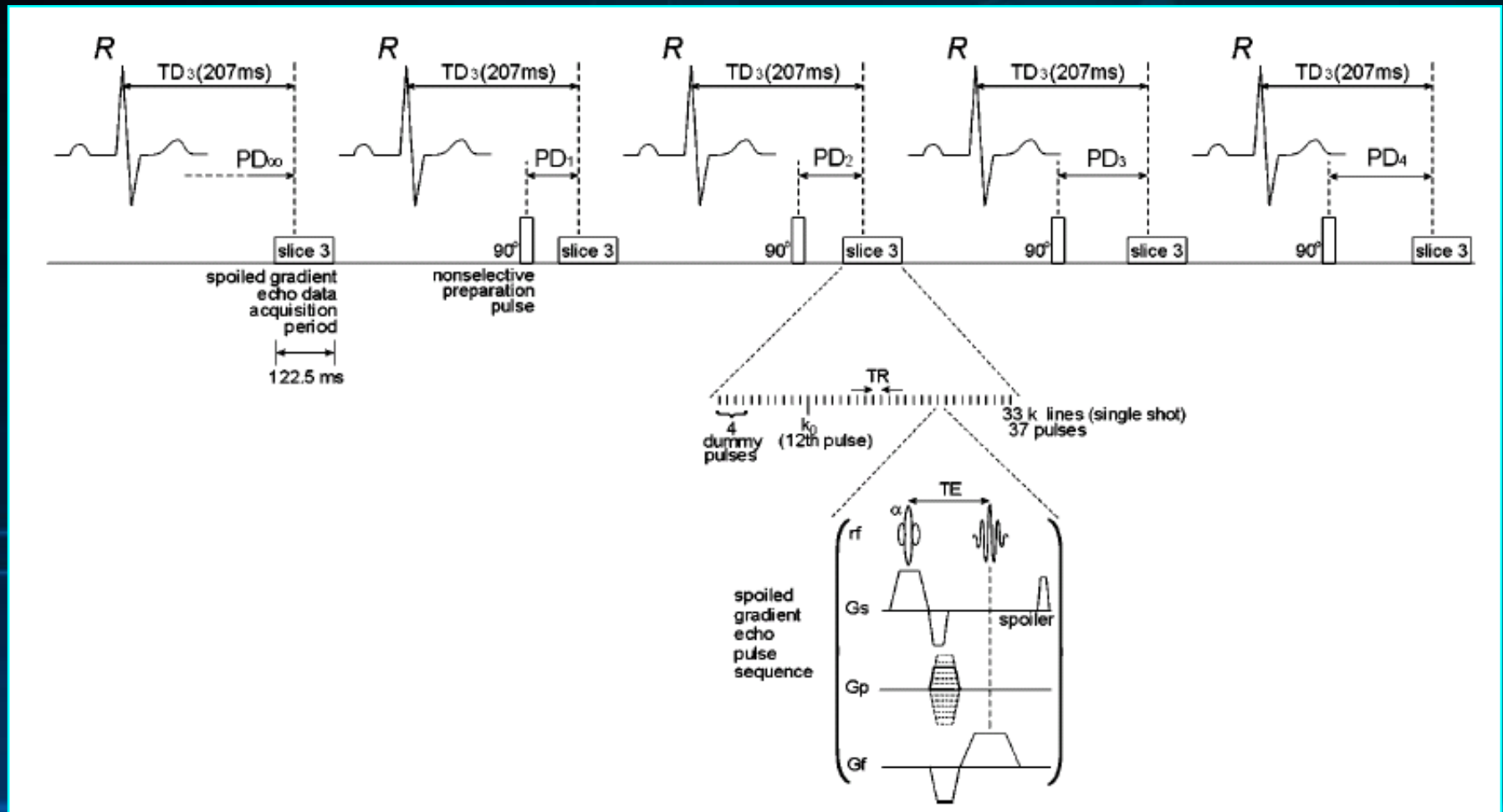
Look-Locker Sequence



- Very sensitive to RF pulse errors
- Magnetization recovery rate, T_1^* :

$$T_1^* = \frac{\tau}{\left(\frac{\tau}{T_1}\right) - \ln(\cos \alpha)}$$

Saturation Recovery for CMRI



T1 Parametric Maps



(a) T1 Map of tubes of gel doped with Gd-DTPA

(b) T1-weighted image of heart in short axis

(c) T1 parametric map image of heart in (b)

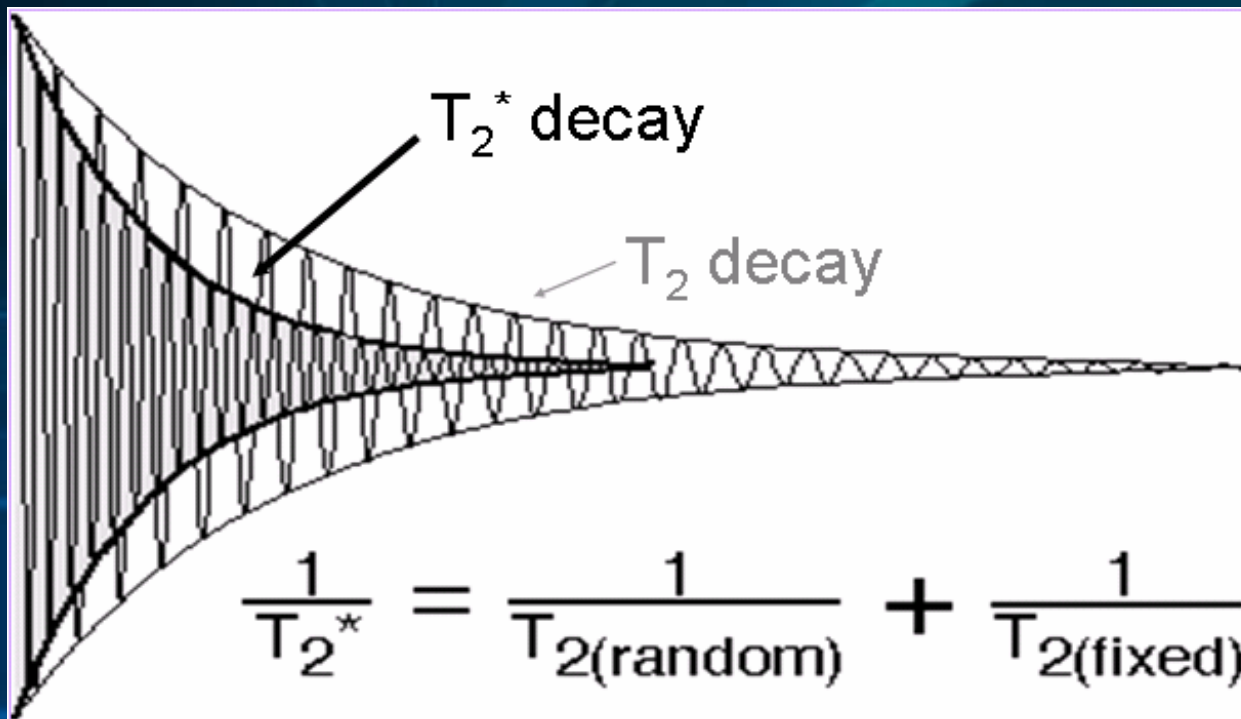
- T1's calculated from short-acquisition period T1 sequence (SAP-T1) with varying delay times

For Accurate & Precise T1

- Never Assume RF Flip Angle is Correct
 - Varies over imaged slice due to slice profile
 - Flip angle must be calibrated across slice
- Be careful in assuming magnetization has reached steady state between acquisitions
- Optimize sequence acquisition parameters to ensure maximal SNR
- Always check that fitted conforms to assumed model

Transverse Magnetization

T₂* Decay



Multi-Echo Acquisitions



Calculation of T_2

$$M_{xy} = M_o' e^{-t/T_2}$$

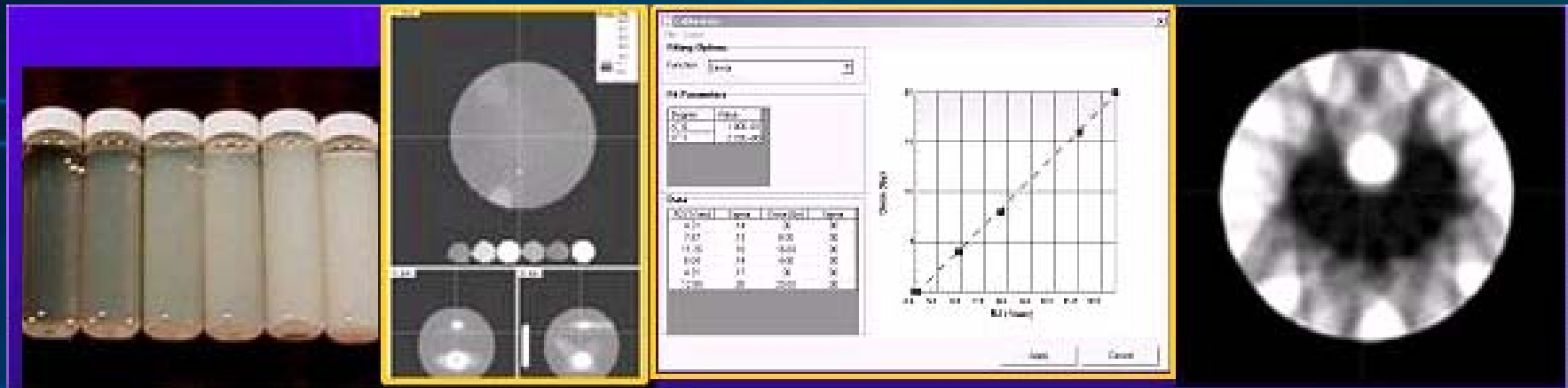
$$\ln M_{xy} = -1/T_2 t + \ln M_o'$$

$$\ln(M_{xy} / M_o') = \text{slope} = -1/T_2$$

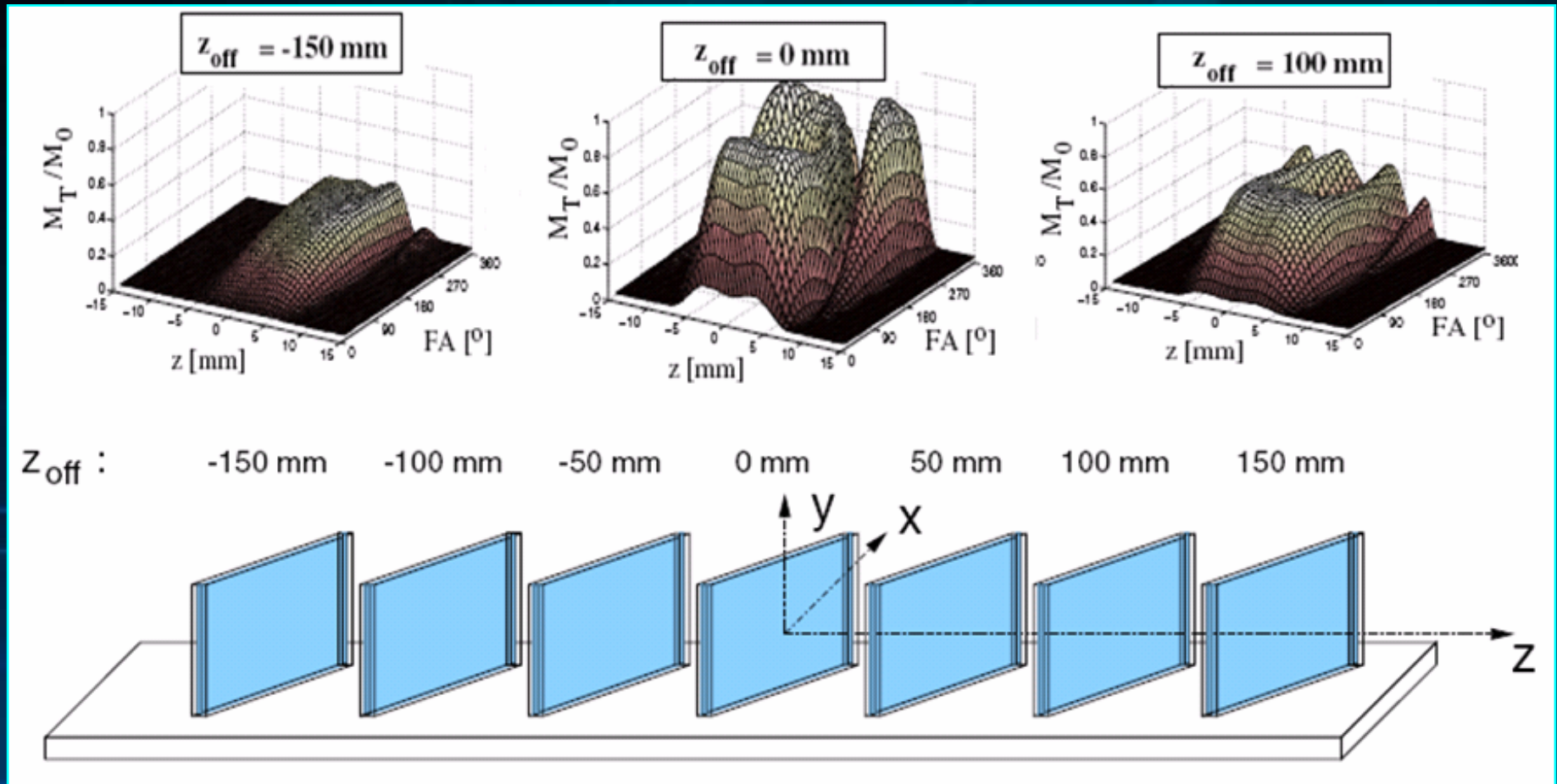
$$T_2 = -1 / \text{slope}$$

Gel Dosimeters

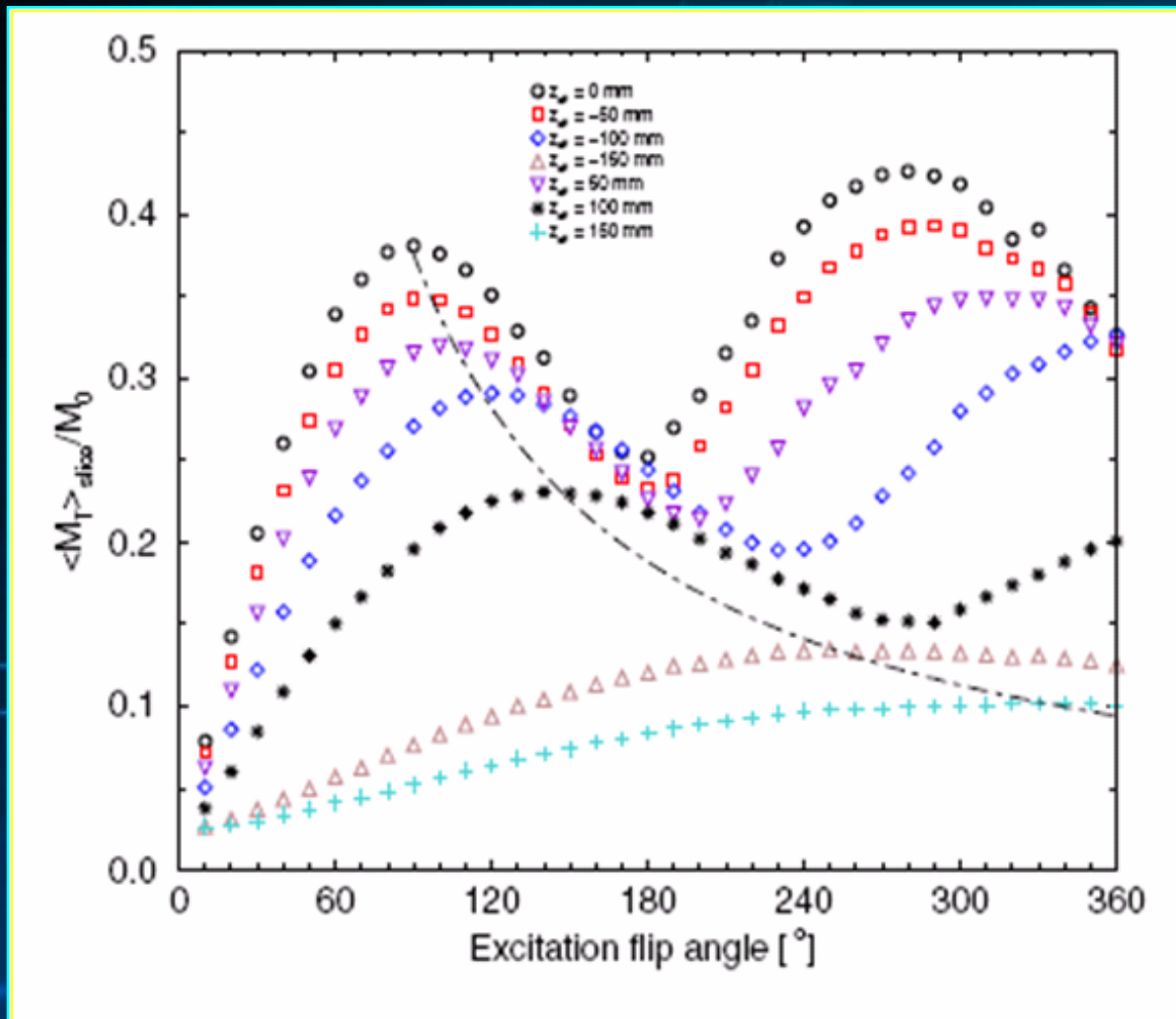
- Used for 3D Radiation Dosimetry QC
- Relies on direct relationship between relaxation rate, R_2 ($R_2=1/T_2$) of gel following exposure and dose



B1 Changes with Slice Position

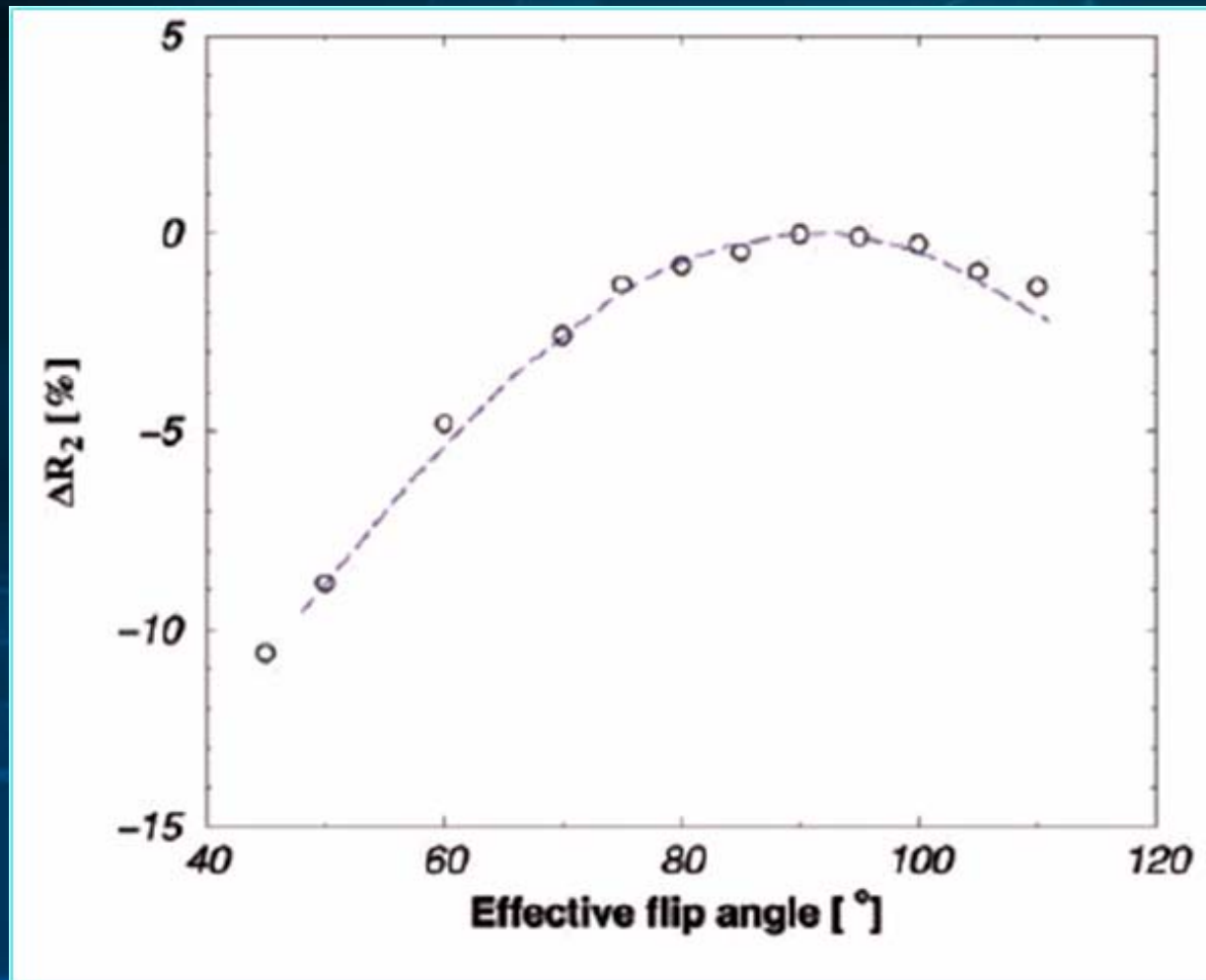


Effective Flip Angles



Average transverse magnetization within a slice as a fraction of M_0 for various slice positions for flip angles ranging from 0° to 360°

R₂ Calibration



For Accurate & Precise T2

- Never Assume RF Flip Angle is Correct
 - Varies over imaged slice due to slice profile
 - 180° flip angle must be calibrated across slice
- Use multi-echo (vs. dual echo) approach and big TX coils whenever possible
- Analyze and understand eddy current effects on T2 measurement
- In tissues, beware of multi-exponential decay

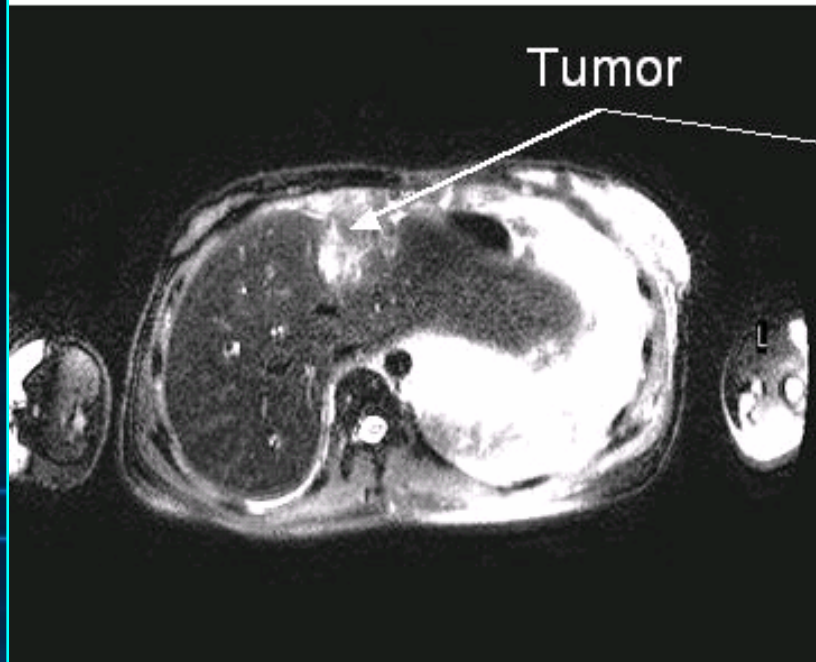
T2* Parametric Imaging

$$M_{xy} = M_o' e^{-t/T_2^*}$$

- Similar to T2 measurements but use gradient echo imaging with varying TE

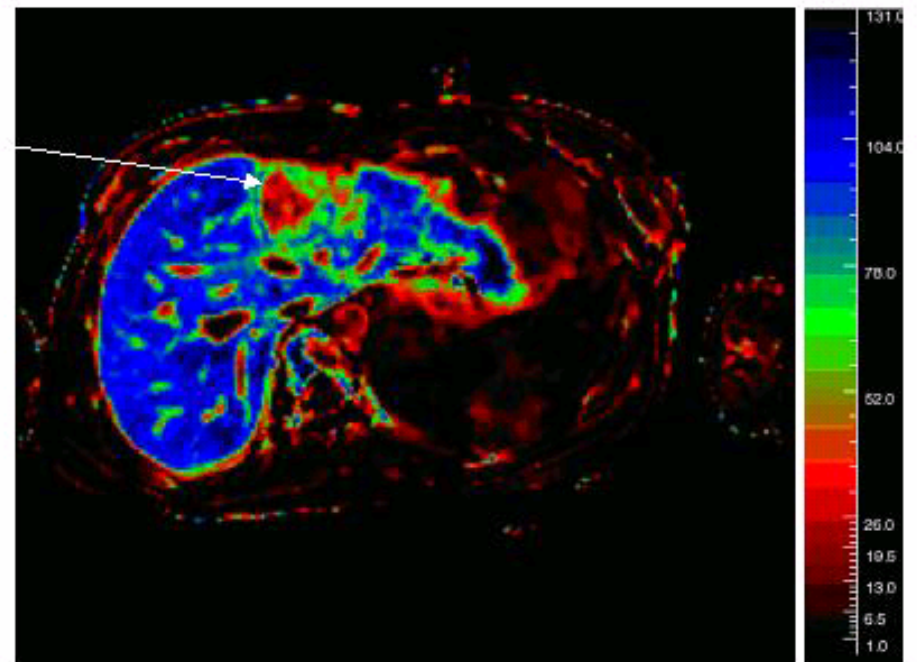
Contrast Agent Maps

contrast weighted image



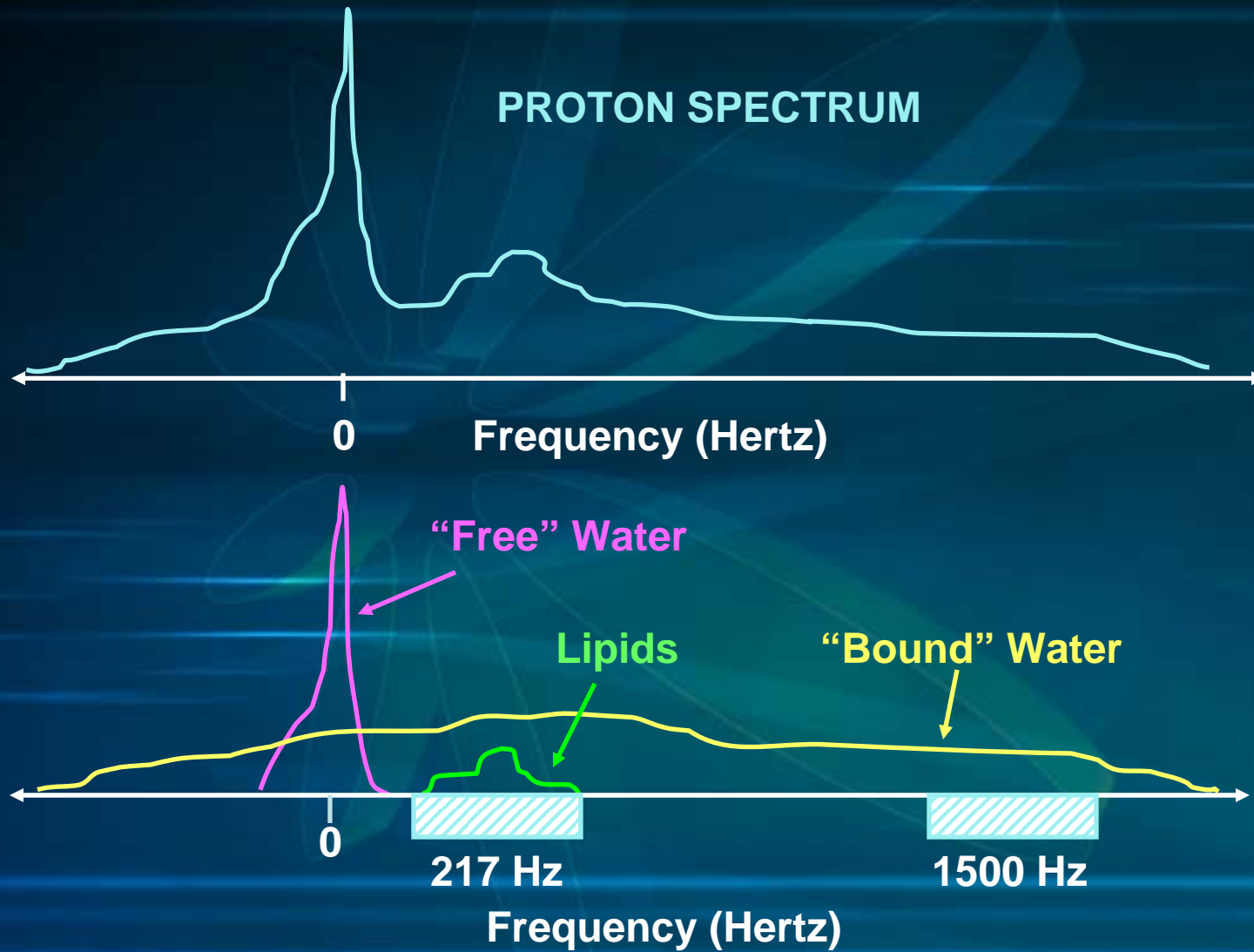
T1-weighted image

contrast agent distribution map



Parametric map of R_2^*

Magnetization Transfer



Magnetization Transfer Ratio

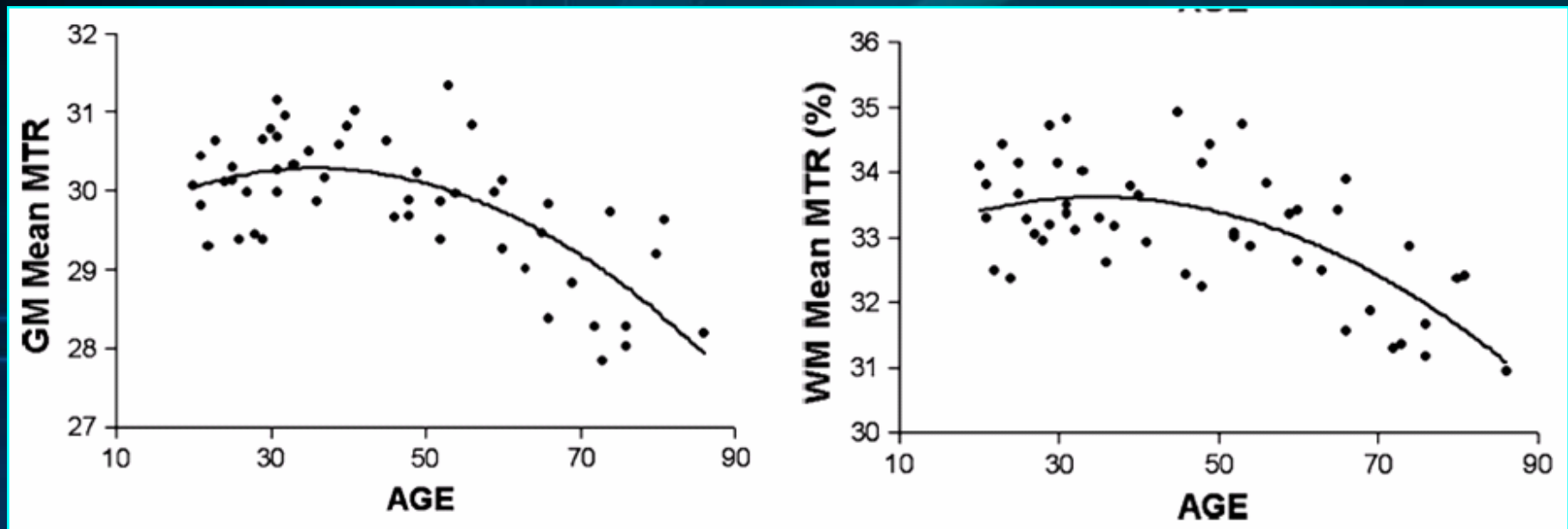
$$MTR = \frac{M_0 - M_s}{M_0}$$

Magnetization Transfer Ratio (MTR)

- the difference of the saturated versus non-saturated images relative to the signal in the normal (non-saturated images)

MTR and Aging

Gray matter and white matter MTR images reveal a quadratic change with age that is primarily attributed to normal demyelination



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Physiological Measurements

- **Flow** – bulk motion of blood and other fluids within body
- **Perfusion** – amount of blood traveling through capillaries in ml/s/gm of tissue
- **Diffusion** – random motion of spins in a homogeneous solution
 - **Apparent Diffusion Coefficient** – measured diffusion rate of water through tissue

Attenuation Due to Diffusion

$$A(TE) = A(0) \exp\left[-\gamma^2 G^2 D_{app} \delta^2 \alpha^2 \left(\Delta - \frac{\delta}{4}\right)\right]$$

Where: $\alpha = \pi/2$;

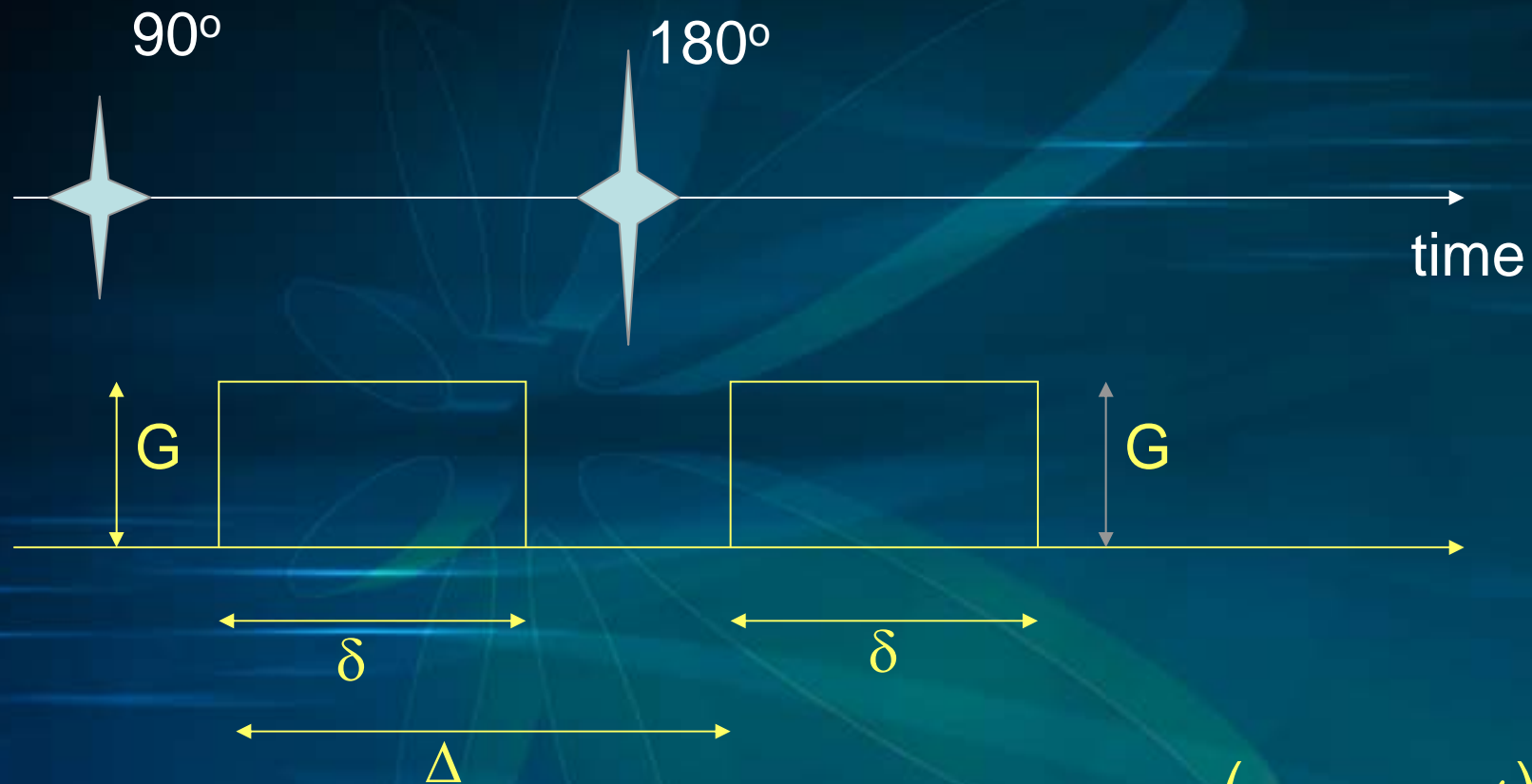
G is amplitude of diffusion sensitive gradient pulse;

δ is duration of diffusion sensitive gradient;

Δ is time between diffusion sensitive gradient pulses;

D_{app} is the apparent diffusion coefficient

DWI Basic Pulse Sequence

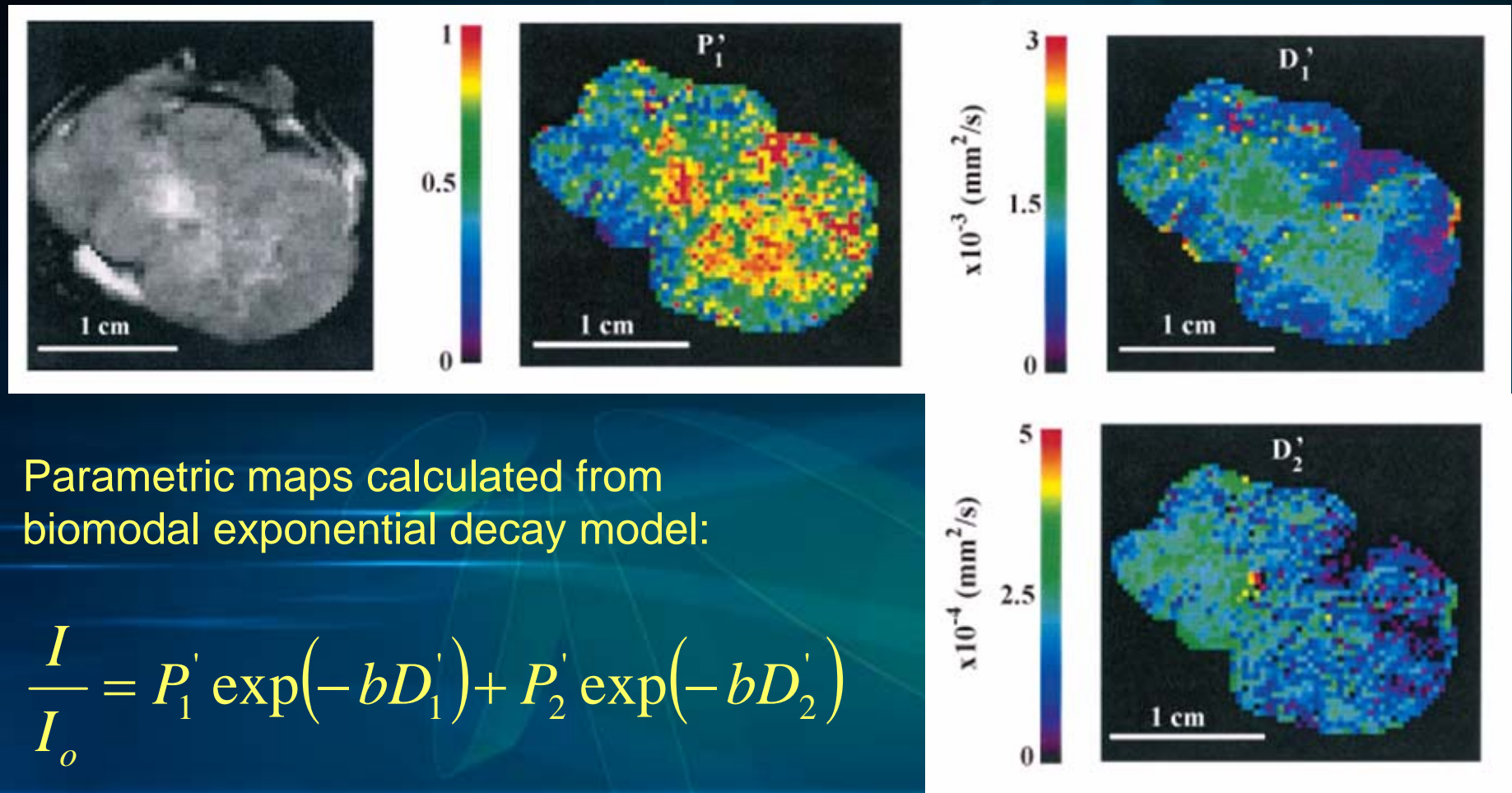


$$b = \gamma^2 G^2 \delta^2 \left(\Delta - \frac{\delta}{3} \right)$$

The b-value

- Controls amount of diffusion weighting in image
- The greater the b-value the greater the area under the diffusion-weighted gradient pulses
 - longer TE
 - stronger and faster ramping the gradients

DW MRI on Breast Tumor



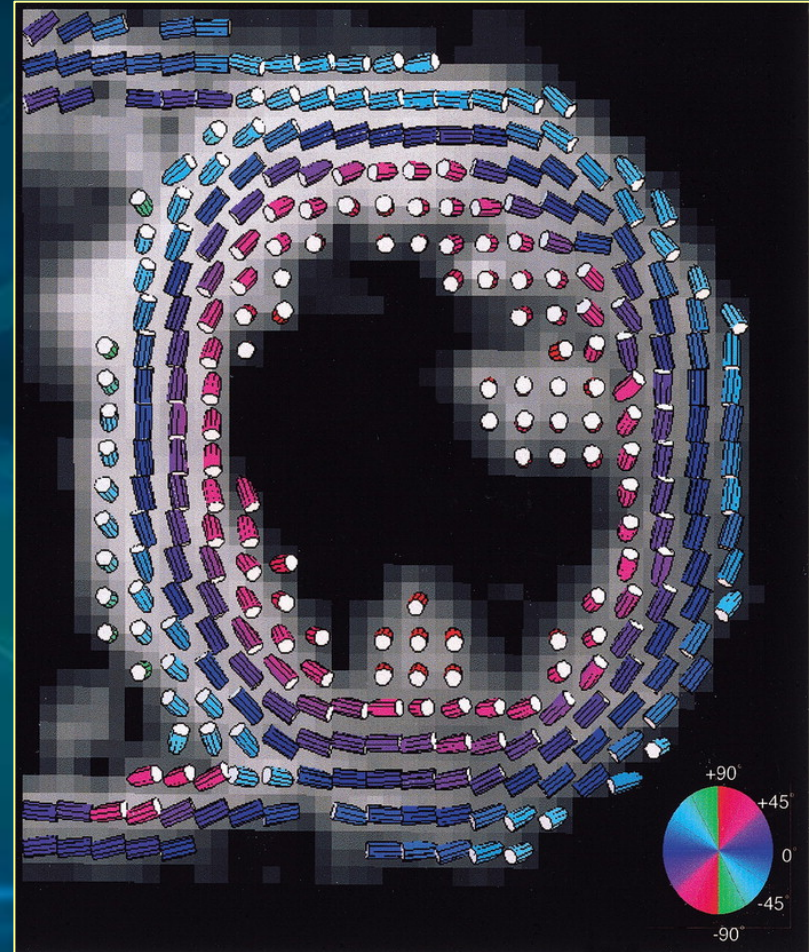
Anisotropic Diffusion



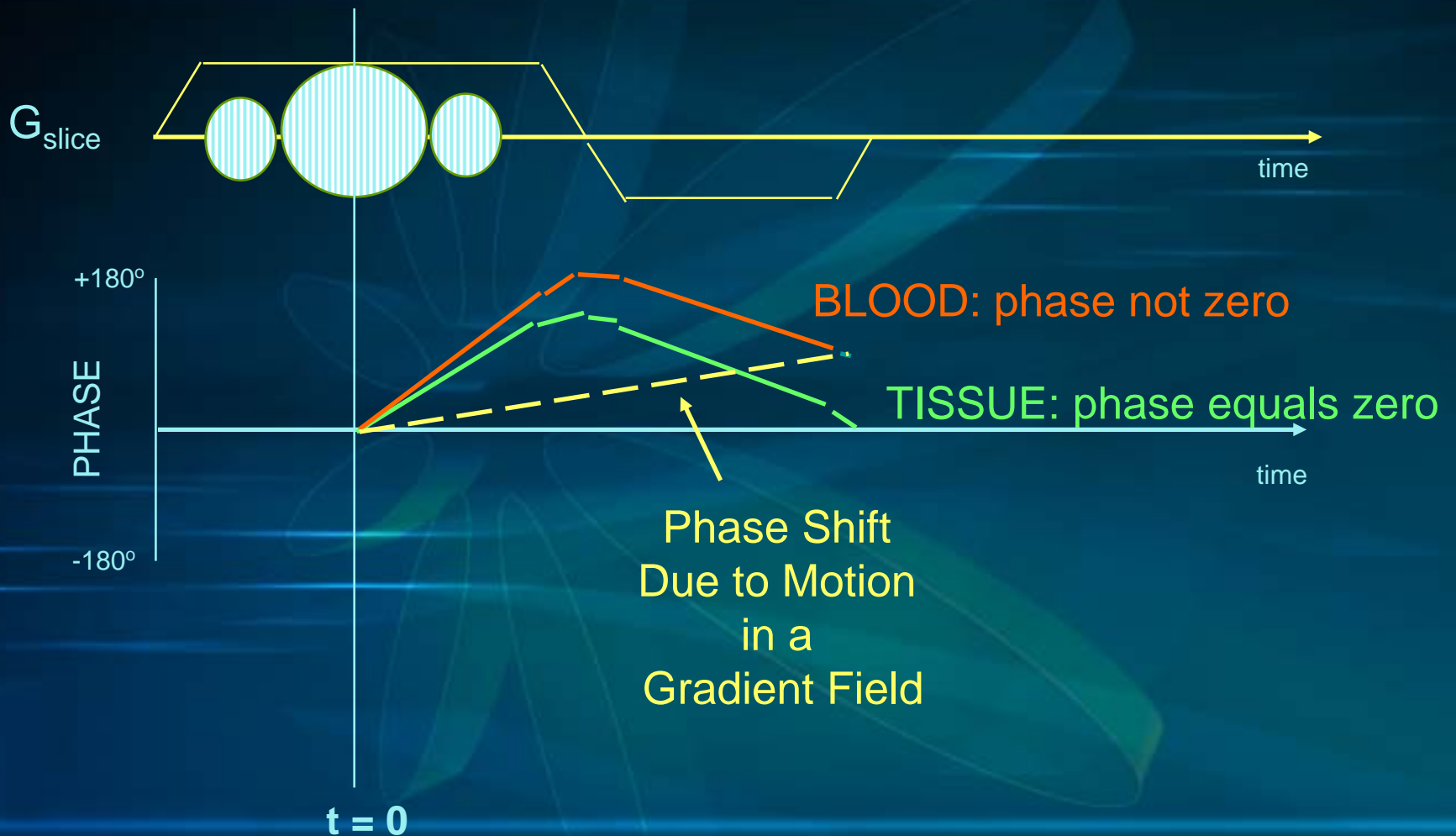
Restricted diffusion along neural fibers

Diffusion Tensor Imaging

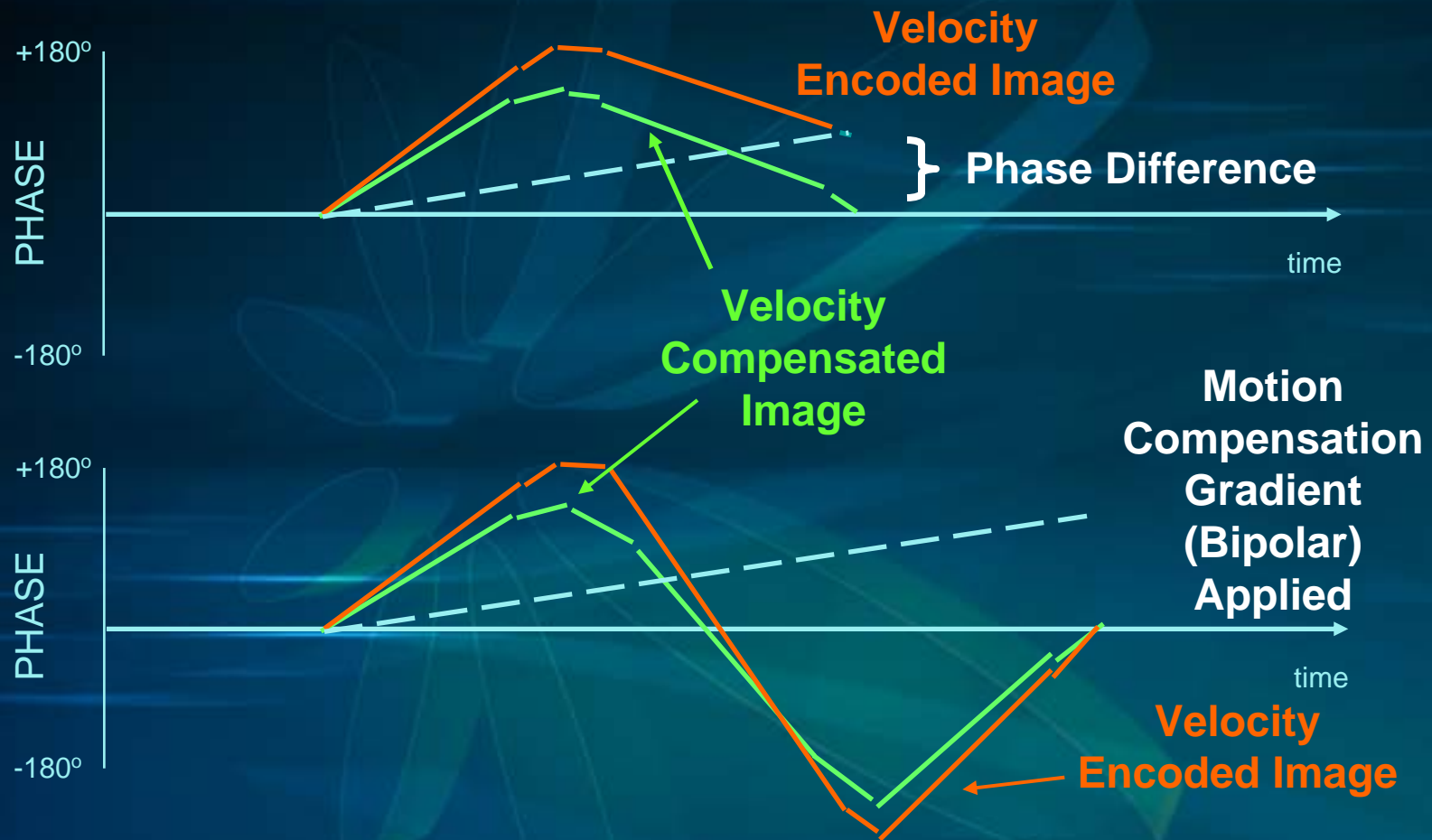
- In anisotropic tissues (neural fibers, muscle fibers) scalar ADC depends on direction of diffusion sensitizing gradient
- Diffusive transport of water can be characterized by an *effective* diffusion tensor
- Direction of diffusion can be used to create a map showing orientation of myocardial fibers.



Dephasing Due to Motion



Phase Contrast Imaging



TISSUE: phase equals zero in BOTH images

BLOOD: phase is DIFFERENT in each image

Phase Contrast Images

Two Signals:

$$S_1 = S_s + S_m ; S_2 = S_s + S_m e(i\varphi_m)$$

where $\varphi_m = \gamma \Delta M_1 v$ ($v = \text{velocity}$)

Complex difference -

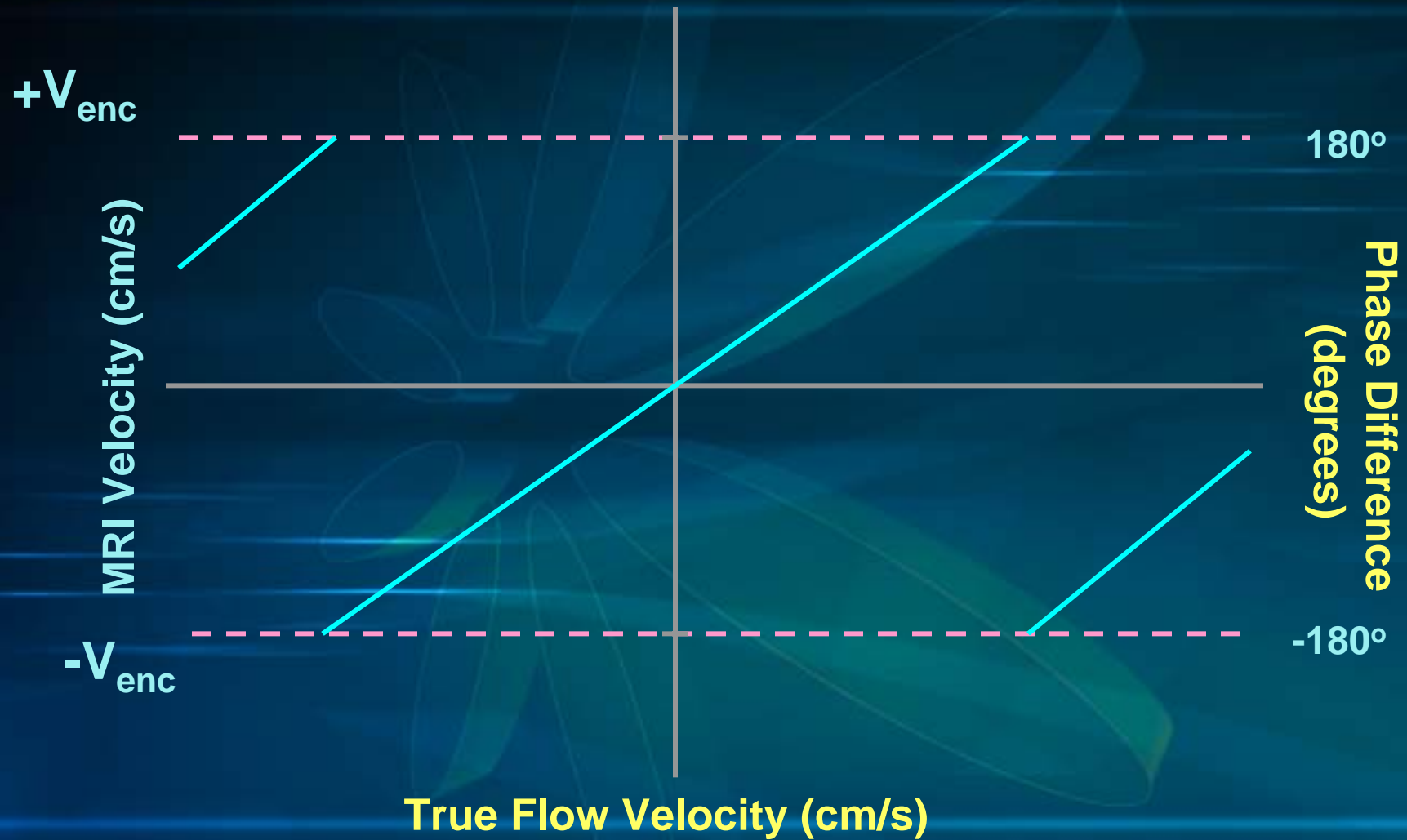
$$\Delta S = S_2 - S_1 = S_m [e(i\varphi_m) - 1] = 2iS_m \sin(\varphi_m/2)$$

Phase difference -

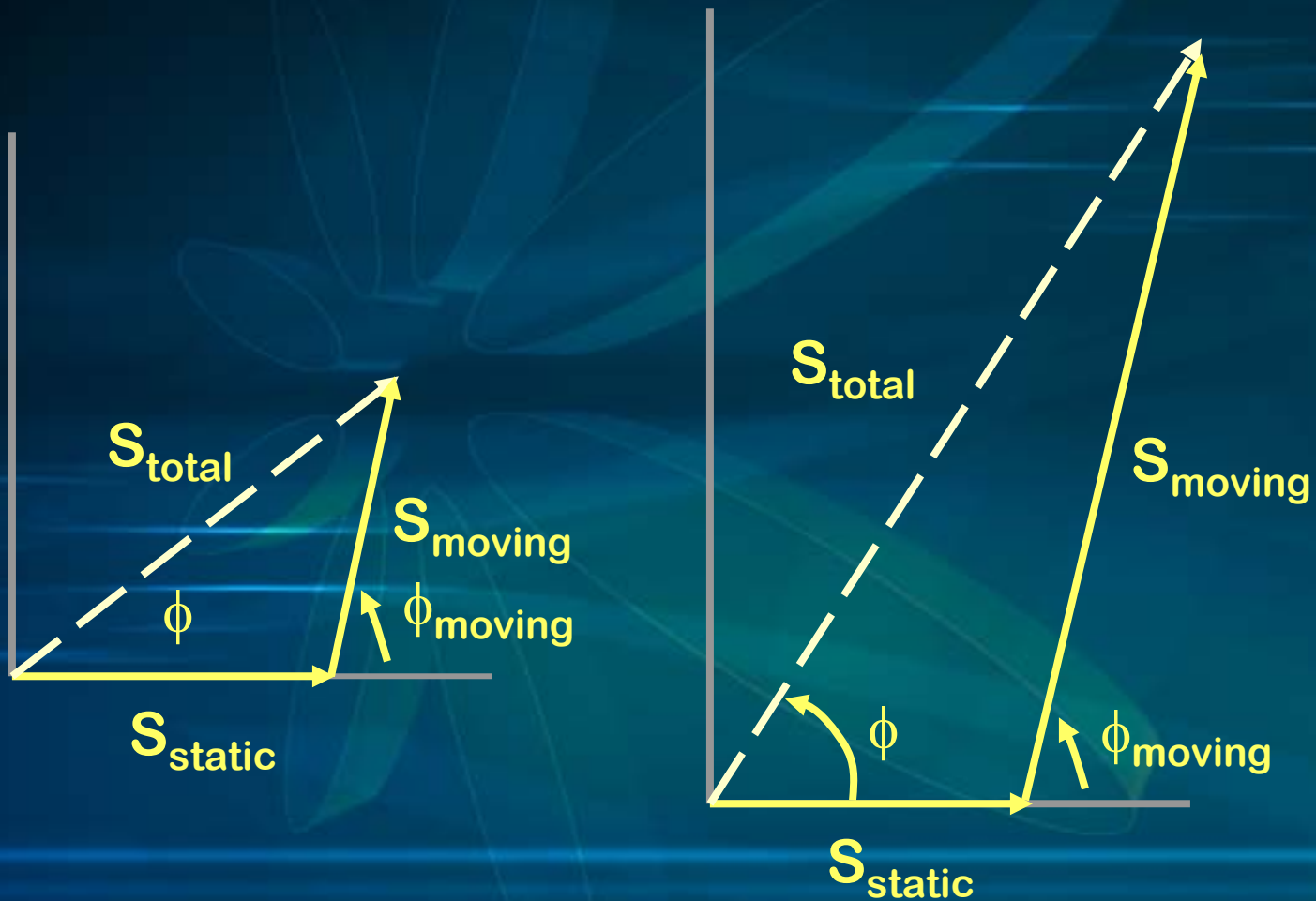
$$\Delta\varphi = \arg S_2 - \arg S_1, \text{ then}$$

$$v = \Delta\varphi / (\gamma \Delta M_1)$$

Velocity Encoding (V_{enc})



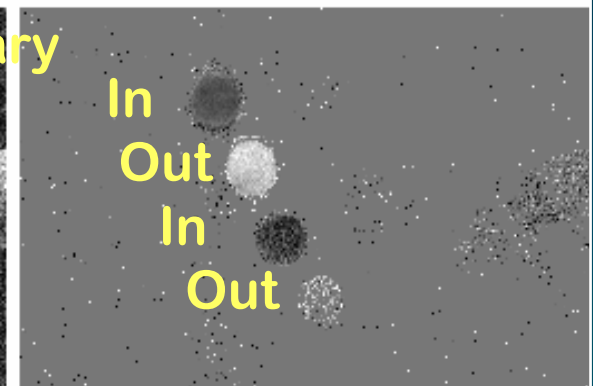
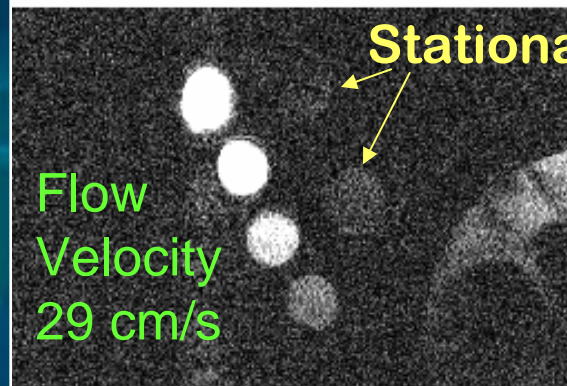
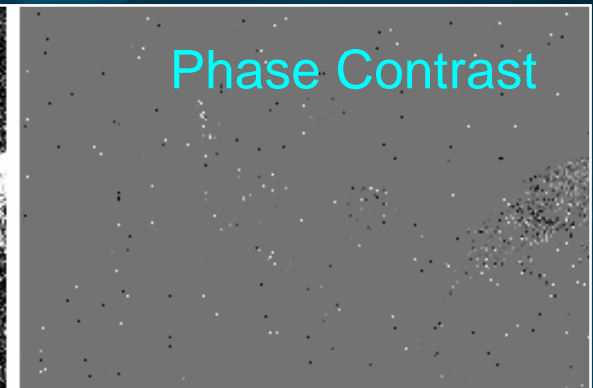
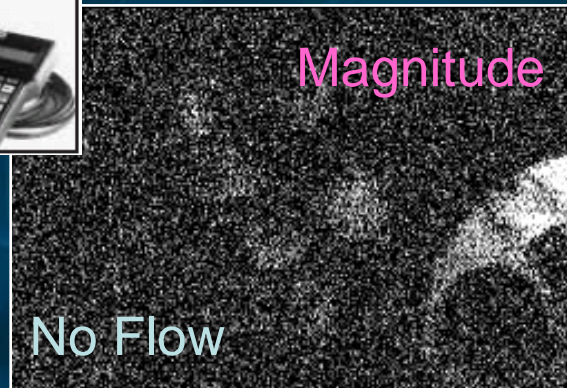
SNR in Phase Contrast



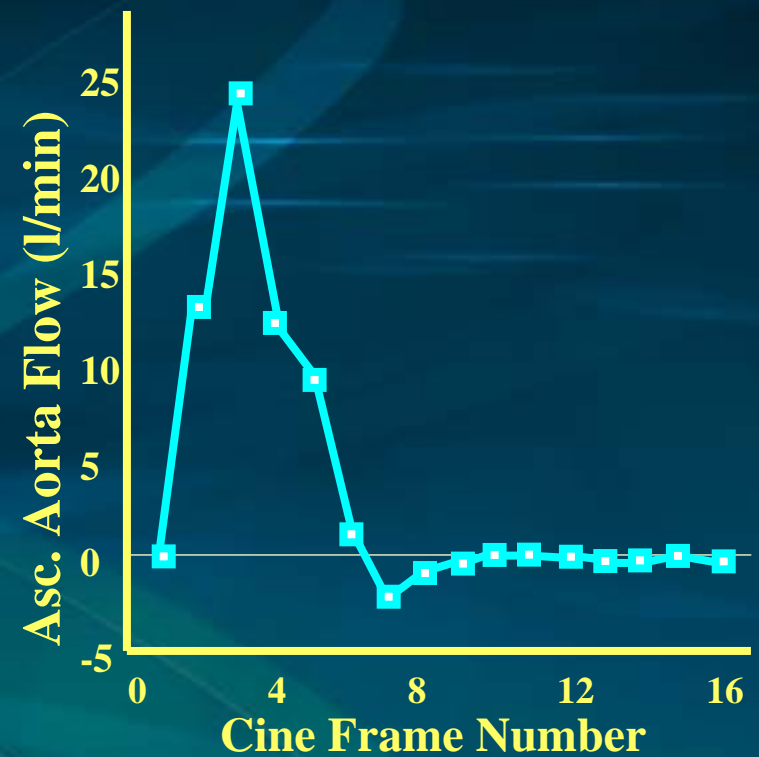
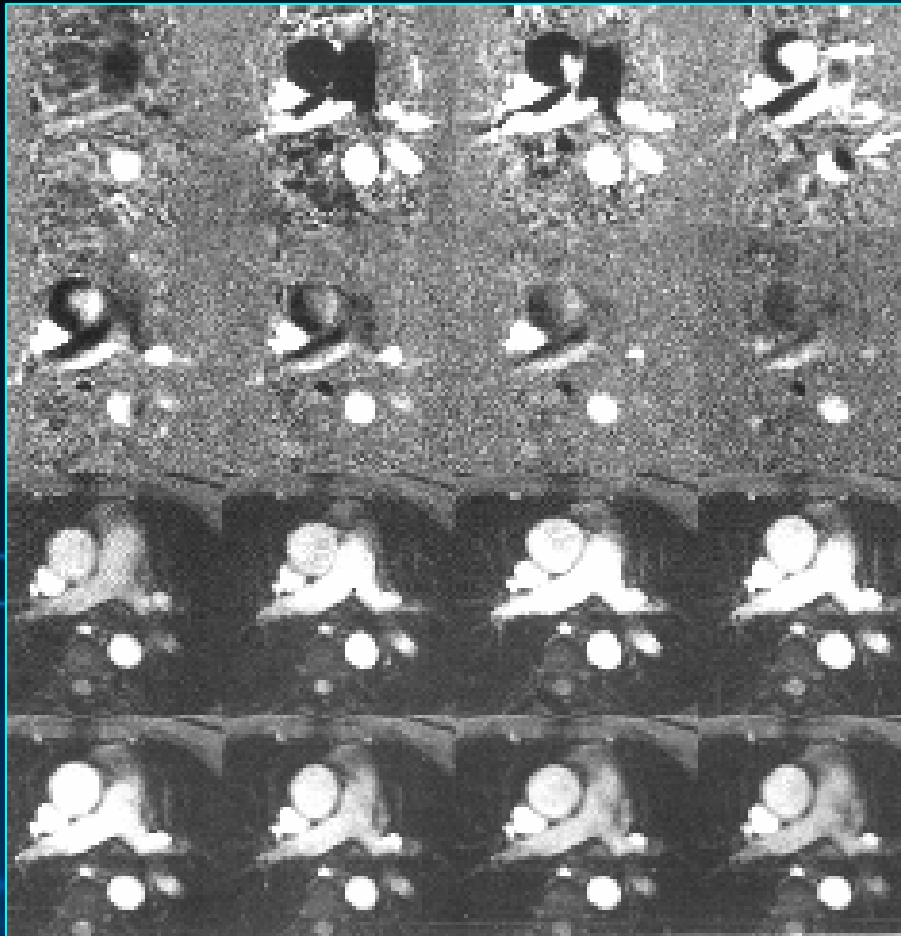
Flow Phantom Calibration



Can use commercially available flow pumps to accurately simulate blood flow in various vessel in the body.

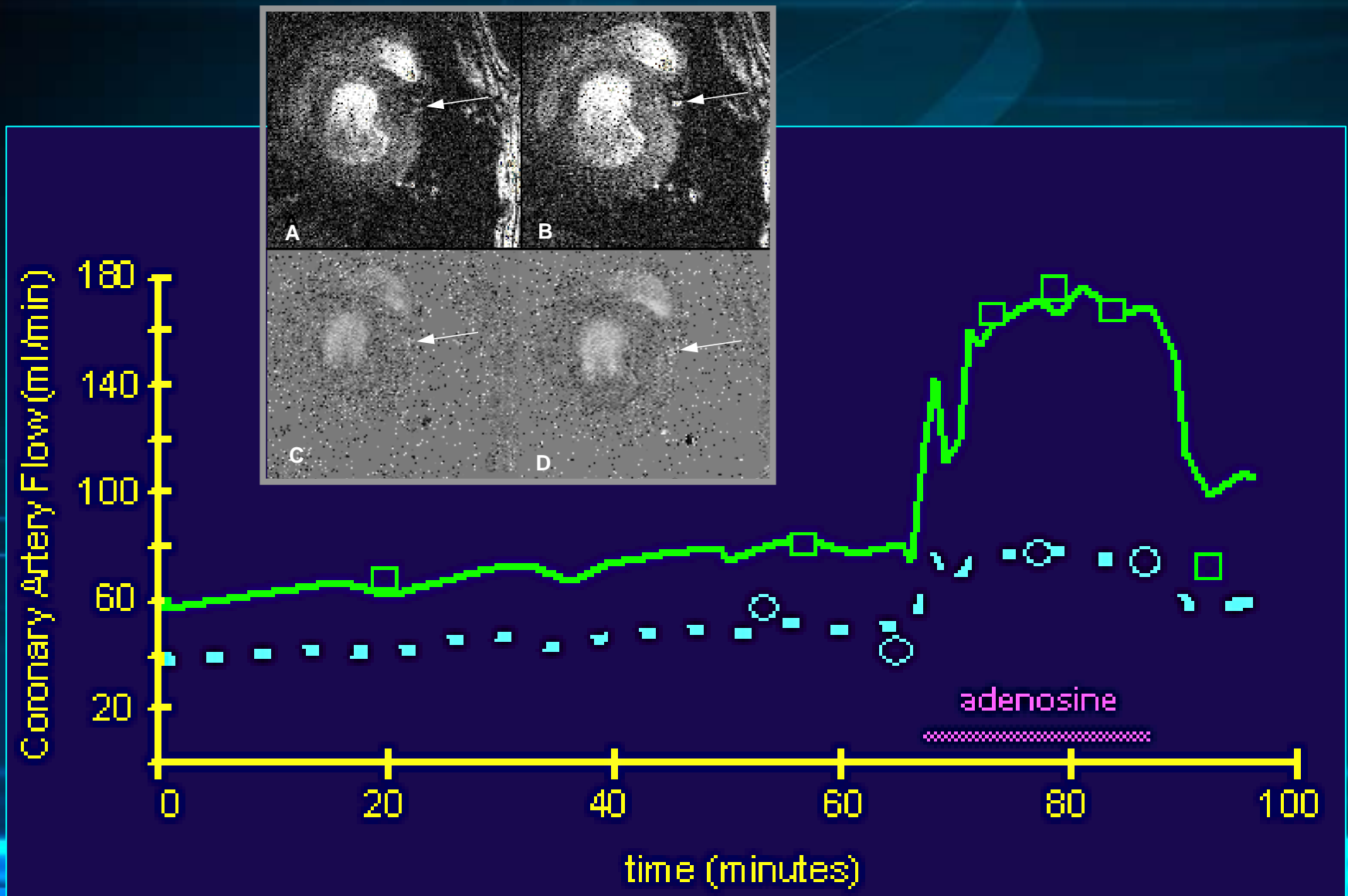


Cardiac Output

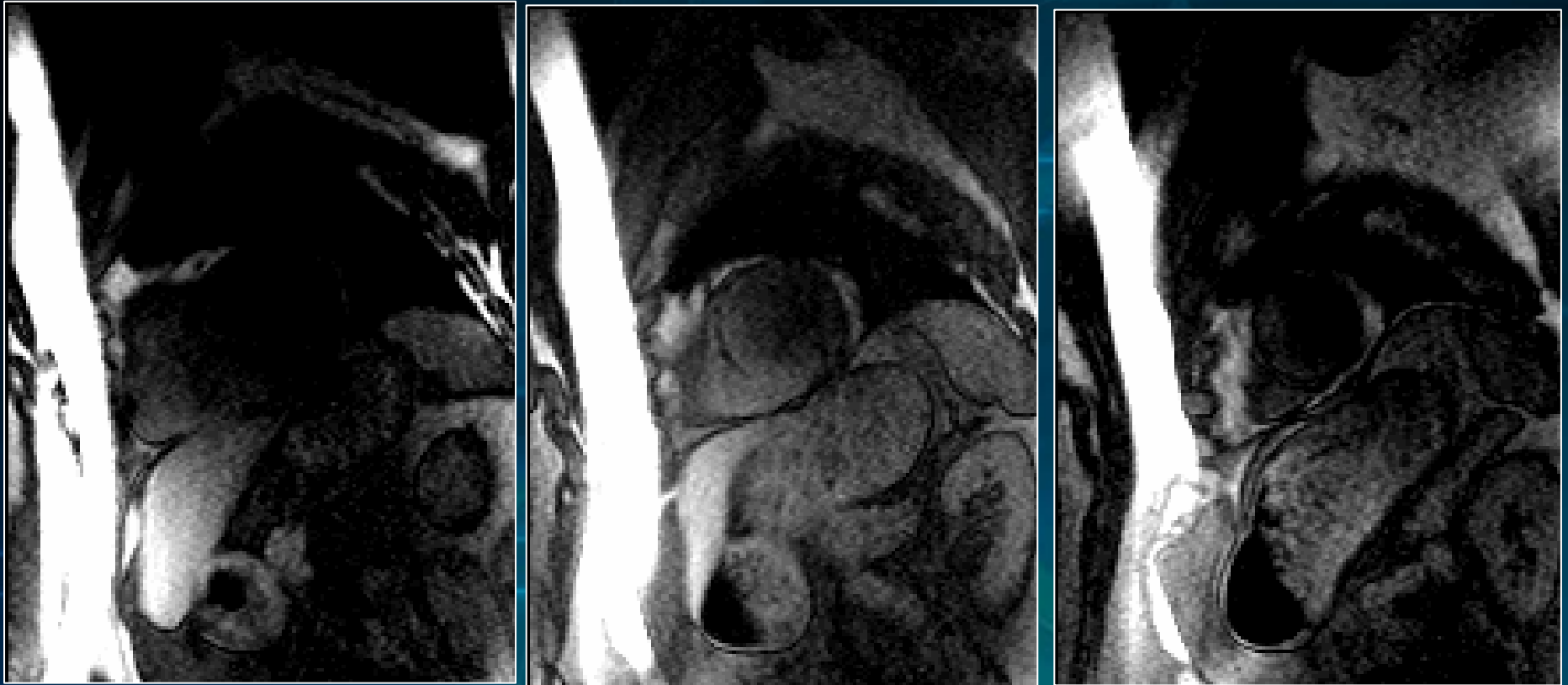


Cardiac Output = 4.01 L/min
Stroke Volume = 68 ml

Coronary Artery Flow



Myocardial Perfusion



Short-axis views of patient's heart showing Gd-DPTA Uptake into RV, LV and then myocardium

Nonuniformity Correction

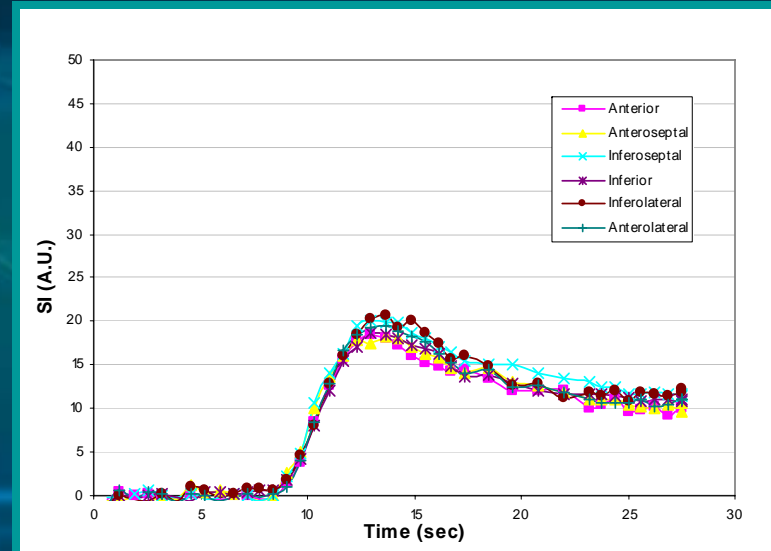
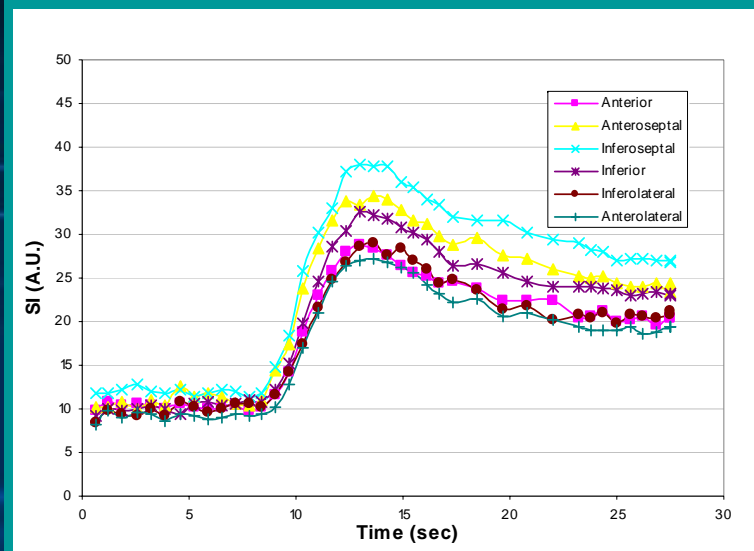
Real-time TrueFISP (GRAPPA=2) cine study was performed between two perfusion scans on the heart of a normal volunteer

$$SI_{\text{normalization}} = SI_{\text{perf}} / SI_{\text{cine}}$$

$$SI_{\text{correction}} = SI_{\text{normalization}} - SI_{\text{base}}$$

Correct for overall effects

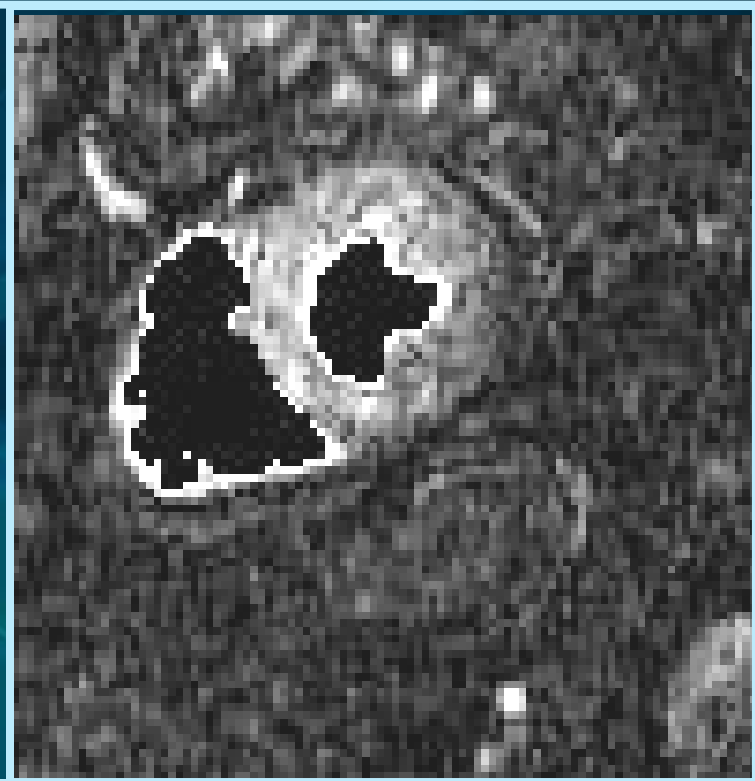
Correct for baseline signal



Max Upslope Parametric Map



MRI perfusion image



Maximum upslope
parametric map

SUMMARY

Before undertaking qMRI:

- Check gradient calibrations
- Understand gradient non-linearities
- Evaluate eddy currents
- Measure RF pulse changes in space
- Determine RF receive nonuniformities

Quantitative MRI Methods

- Measuring things with MRI:
 - Diffusion Coefficients
 - Frequency of a signal
 - Relaxation rates (T1, T2 , MTC)
 - Velocity of motion
 - Volumes of tissues

Learn More Details ...

