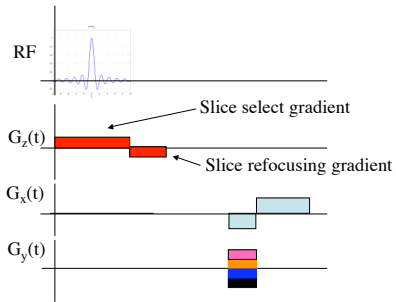


Bioengineering 280A
Principles of Biomedical Imaging

Fall Quarter 2005
MRI Lecture 5

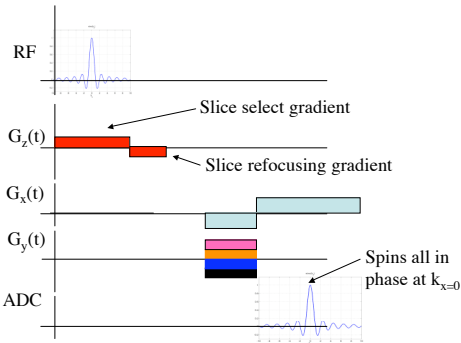
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Slice Selection



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Gradient Echo



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Static Inhomogeneities

In the ideal situation, the static magnetic field is totally uniform and the reconstructed object is determined solely by the applied gradient fields. In reality, the magnet is not perfect and will not be totally uniform. Part of this can be addressed by additional coils called "shim" coils, and the process of making the field more uniform is called "shimming". In the old days this was done manually, but modern magnets can do this automatically.

In addition to magnet imperfections, most biological samples are inhomogeneous and this will lead to inhomogeneity in the field. This is because, each tissue has different magnetic properties and will distort the field.

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Static Inhomogeneities

The spatial nonuniformity in the field can be modeled by adding an additional term to our signal equation.

$$s_r(t) = \int_V M(\vec{r}, t) dV$$

$$= \int_x \int_y \int_z M(x, y, z, 0) e^{-t/T_2(\vec{r})} e^{-j\omega_0 t} e^{-j\omega_e(\vec{r})t} \exp\left(-j\gamma \int_0^t \vec{G}(\tau) \cdot \vec{r} d\tau\right) dx dy dz$$

The effect of this nonuniformity is to cause the spins to dephase with time and thus for the signal to decrease more rapidly. To first order this can be modeled as an additional decay term of the form

$$s_r(t) = \int_x \int_y \int_z M(x, y, z, 0) e^{-t/T_2(\vec{r})} e^{-t/T_2'(\vec{r})} e^{-j\omega_0 t} \exp\left(-j\gamma \int_0^t \vec{G}(\tau) \cdot \vec{r} d\tau\right) dx dy dz$$

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T₂^{*} decay

The overall decay has the form.

$$\exp\left(-t/T_2^*(\vec{r})\right)$$

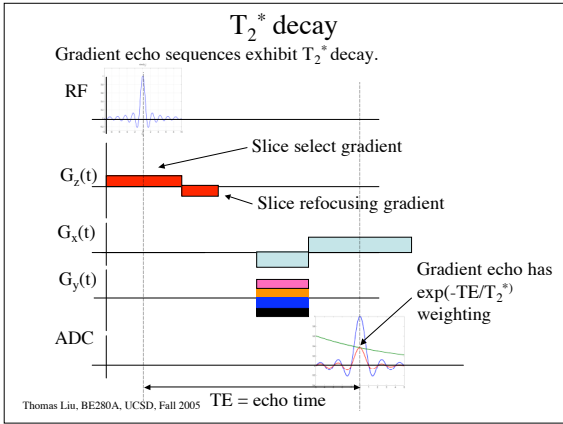
where

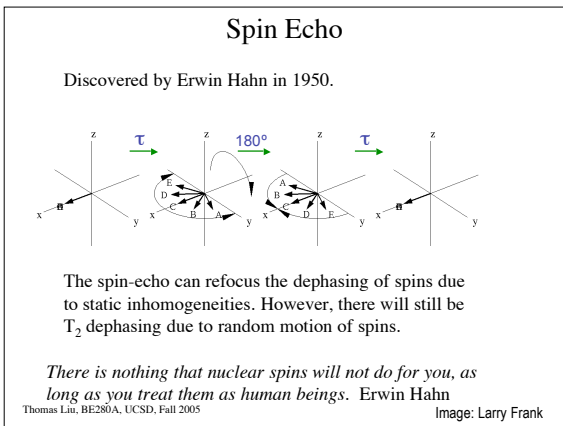
$$\frac{1}{T_2^*} = \frac{1}{T_2} + \frac{1}{T_2'}$$

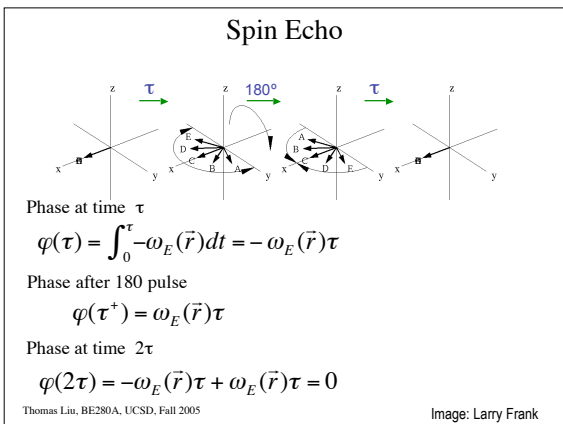
Due to random motions of spins.
Not reversible.

Due to static inhomogeneities. Reversible with a spin-echo sequence.

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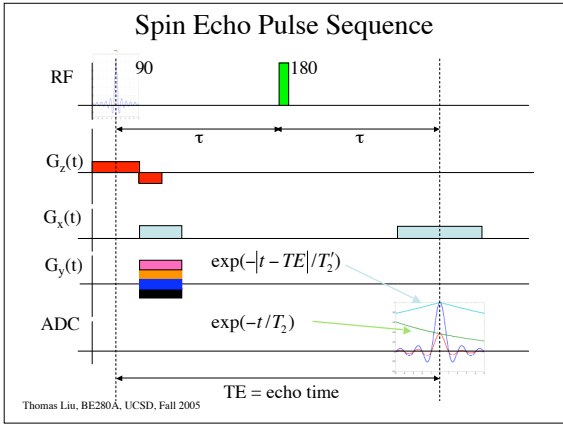
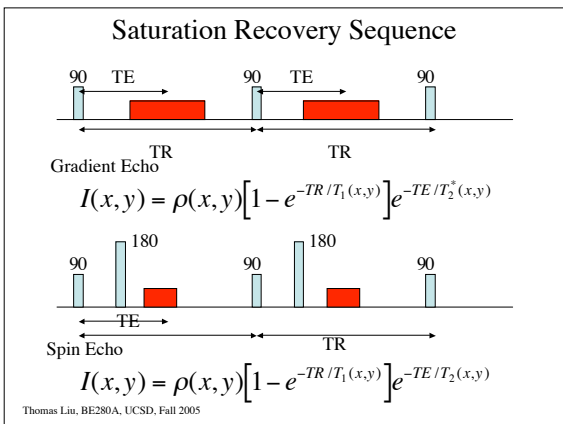


Image Contrast

Different tissues exhibit different relaxation rates, T_1 , T_2 , and T_2^* . In addition different tissues can have different densities of protons. By adjusting the pulse sequence, we can create contrast between the tissues. The most basic way of creating contrast is adjusting the two sequence parameters: TE (echo time) and TR (repetition time).

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T1-Weighted Scans

Make TE very short compared to either T_2 or T_2^* . The resultant image has both proton and T_1 weighting.

$$I(x, y) \approx \rho(x, y) \left[1 - e^{-TR/T_1(x, y)} \right]$$

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T2-Weighted Scans

Make TR very long compared to T_1 and use a spin-echo pulse sequence. The resultant image has both proton and T_2 weighting.

$$I(x, y) \approx \rho(x, y) e^{-TE/T_2}$$

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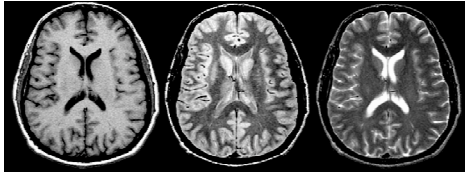
Proton Density Weighted Scans

Make TR very long compared to T_1 and use a very short TE. The resultant image is proton density weighted.

$$I(x, y) \approx \rho(x, y)$$

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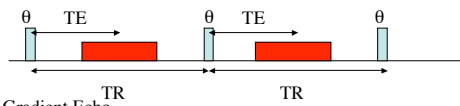
Example



T₁-weighted Density-weighted T₂-weighted

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FLASH sequence



Gradient Echo

$$I(x,y) = \rho(x,y) \frac{[1 - e^{-TR/T_1(x,y)}] \sin \theta}{[1 - e^{-TR/T_1(x,y)} \cos \theta]} \exp(-TE/T_2^*)$$

Signal intensity is maximized at the Ernst Angle

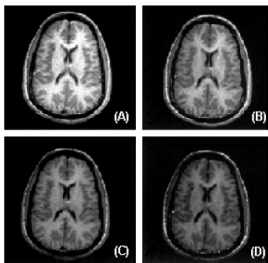
$$\theta_E = \cos^{-1}(\exp(-TR/T_1))$$

FLASH equation assumes no coherence from shot to shot. In practice this is achieved with RF spoiling.

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FLASH/SPGR

Figure 22-7 SPGR Images with Different TRs



- (A) 24 TR and a 30° flip
- (B) 24 TR and a 45° flip
- (C) 50 TR and a 30° flip
- (D) 50 TR and 60° flip

NOTE: At the 24 TR time, the 30° flip is better than the same flip angle at 50 TR.

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Inversion Recovery

$$I(x,y) = \rho(x,y) \left[1 - 2e^{-TI/T_1(x,y)} + e^{-TR/T_1(x,y)} \right] e^{-TE/T_2(x,y)}$$

Intensity is zero when inversion time is

$$TI = -T_1 \ln \left[\frac{1 + \exp(-TR/T_1)}{2} \right]$$

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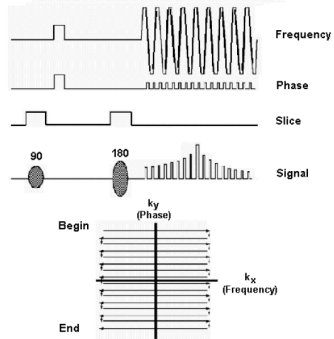
Inversion Recovery

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Fast Spin Echo

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Echoplanar Imaging



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