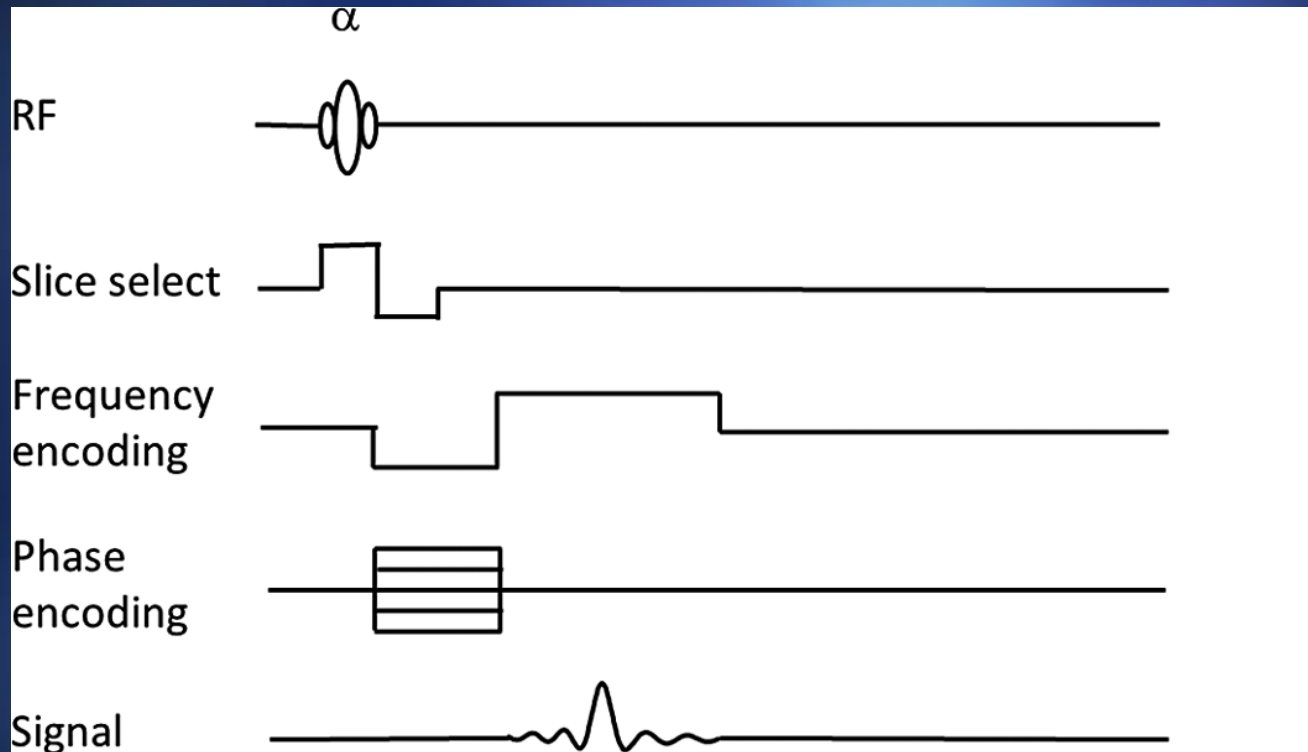


***Speeding Things Up:  
the Gradient-Echo Pulse  
Sequence***

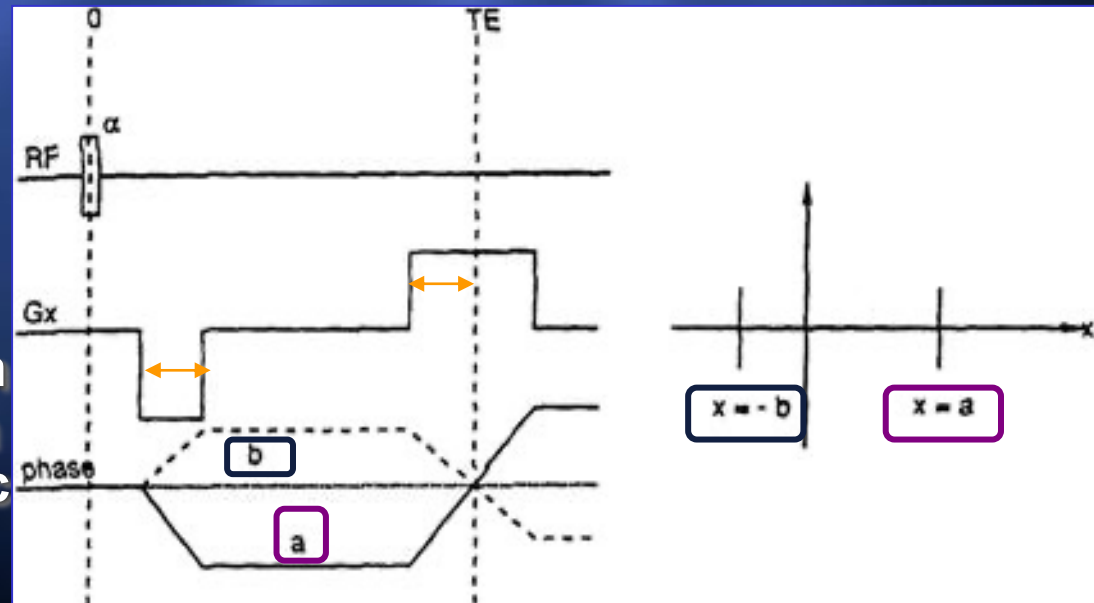
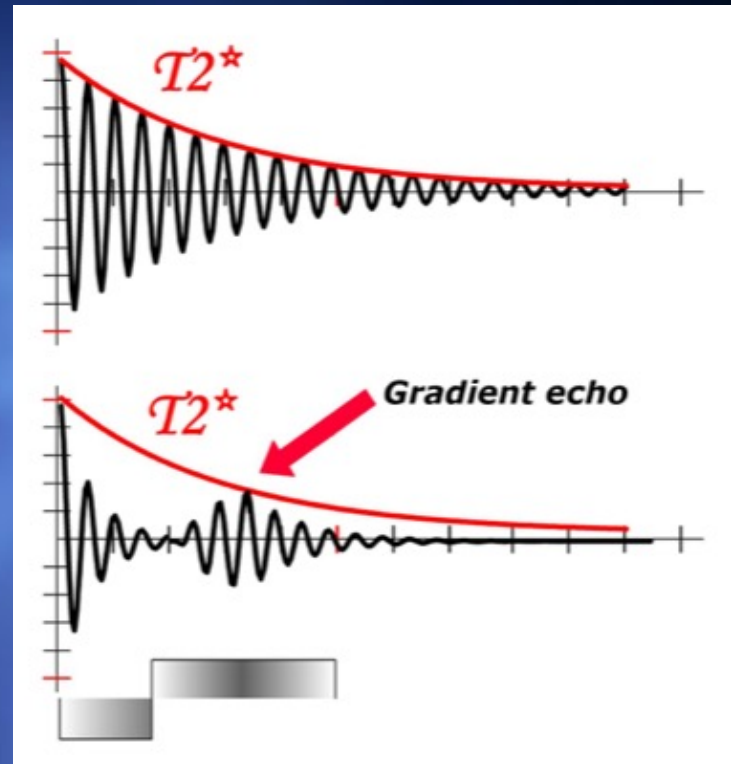
# Gradient-Echo Pulse Sequence



- ✓ the refocusing pulse is omitted the echo is generated only by switching of gradients on the frequency encoding axis
  - ✓ flip angle of the excitation pulse is usually less than  $90^\circ$
- [www.imaios.com/en/e-Courses/e-MRI/MRI-Sequences/gradient-echo](http://www.imaios.com/en/e-Courses/e-MRI/MRI-Sequences/gradient-echo)

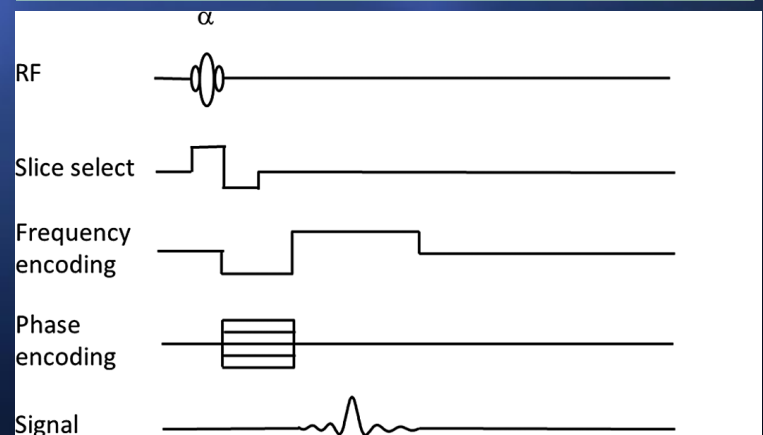
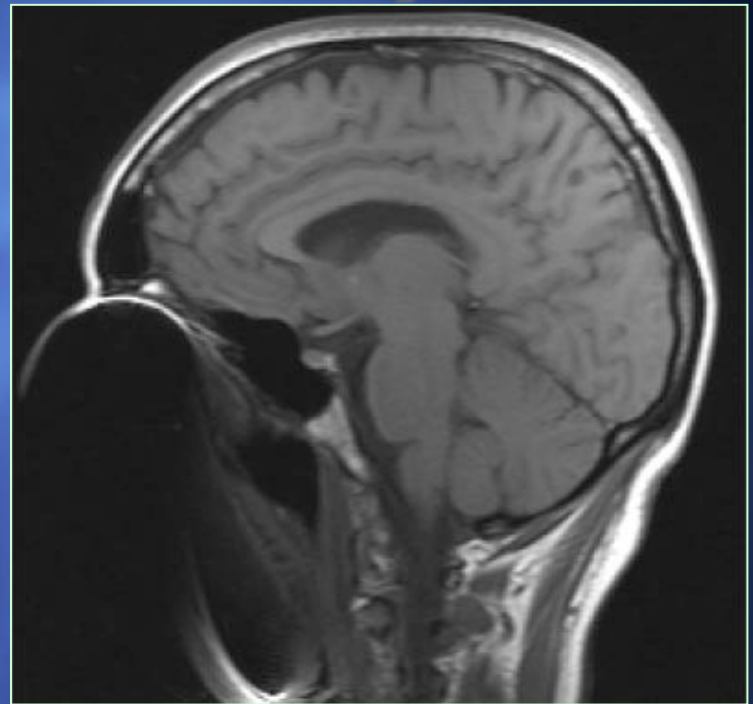
# Gradient Echo

- ✓ No  $180^\circ$  pulse
- ✓  $T_2^*$  contrast
- ✓ No compensation for static field inhomogeneity
  - Magnetic susceptibility
    - ❖ corresponds to the internal magnetization of a tissue resulting from the interactions with an external magnetic field



# Gradient-Echo Pulse Sequence

- ✓ the echo is T2\* intensity
  - less signal than in a spin-echo sequence at the same echo time
- ✓ particularly sensitive to image distortion due to magnetic field inhomogeneities
  - for example at the interface between different tissues or where there are air-filled sinuses or cavities in the body





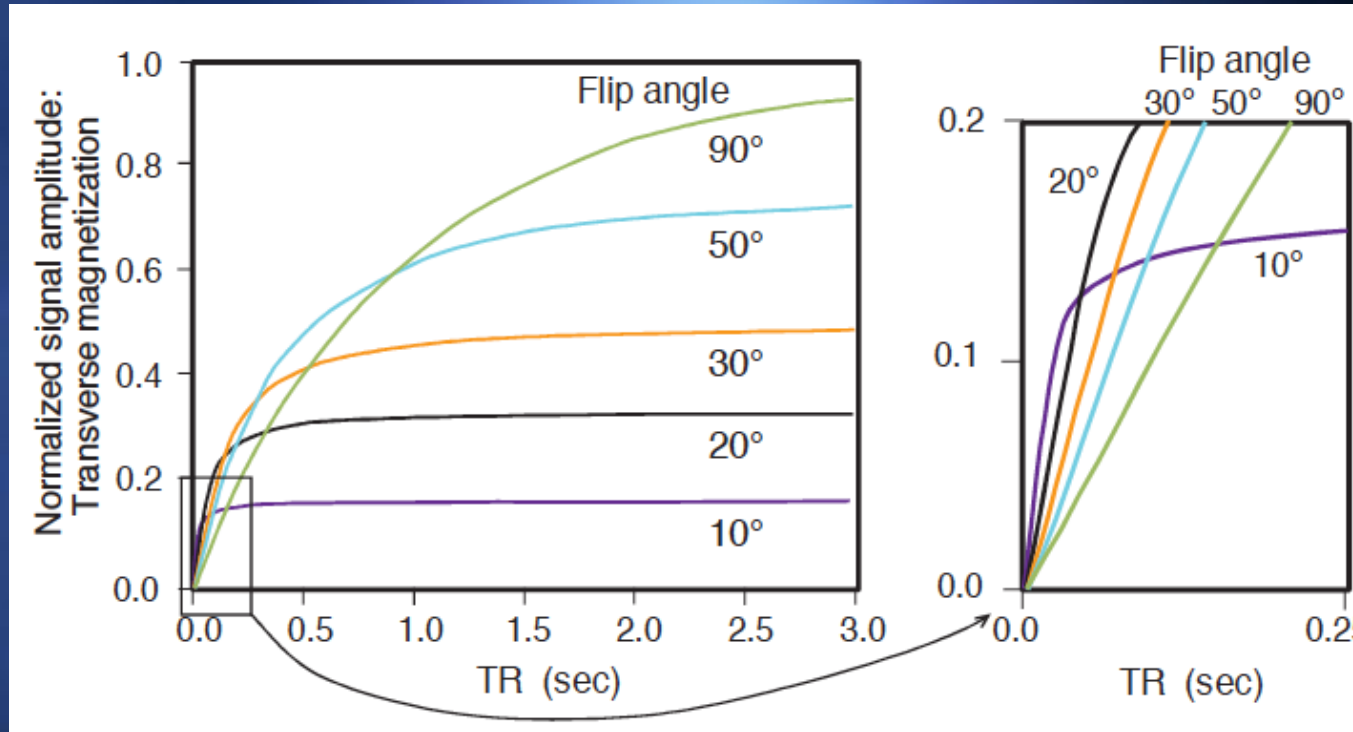
# Flip angle and $T_R$

Given

$$T_R \ll T_1$$

the signal is larger if

$$\alpha < \pi/2$$



$T_R$  defines  
the total acquisition time

# ***Gradient-Echo Pulse Sequence***

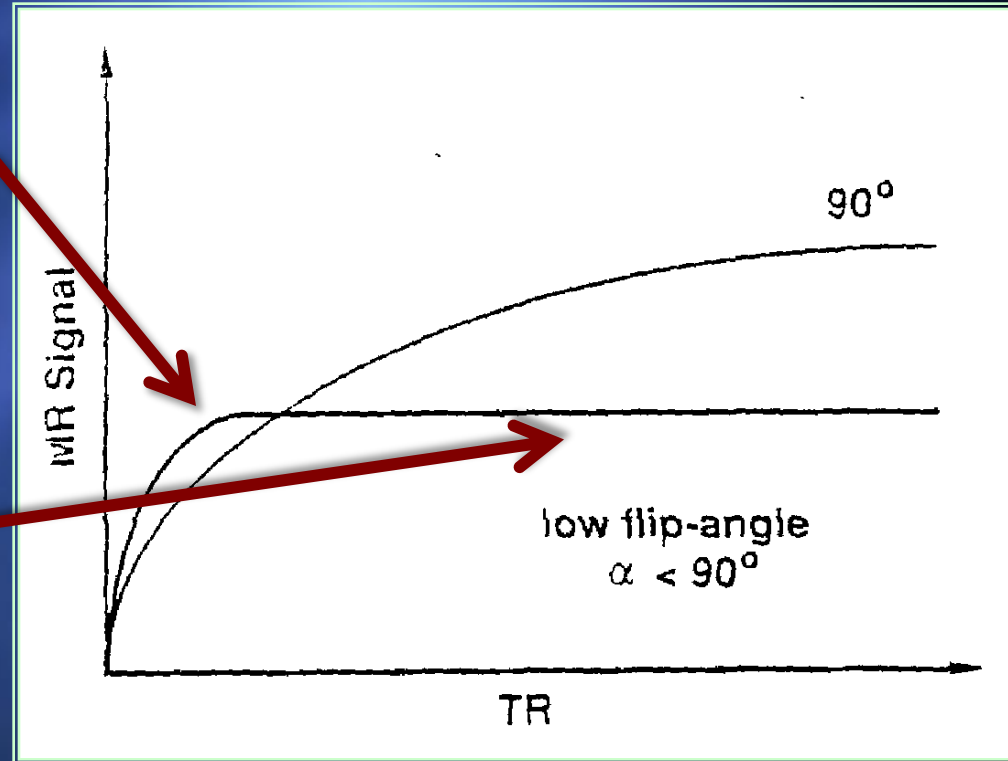
**flip angle  $\alpha = 10^\circ$**

- ✓ **immediately after excitation  $M_z = 0.98|M|$** 
  - $M_z$  reduced by 2% only from its initial value
- ✓  **$T_R$  can be very much shorter**
  - resulting in much quicker image acquisition
- ✓  **$M_{xy} = 0.17 |M|$** 
  - so we have sacrificed most of our signal

***often in MRI, there is a trade-off between image quality and acquisition time***

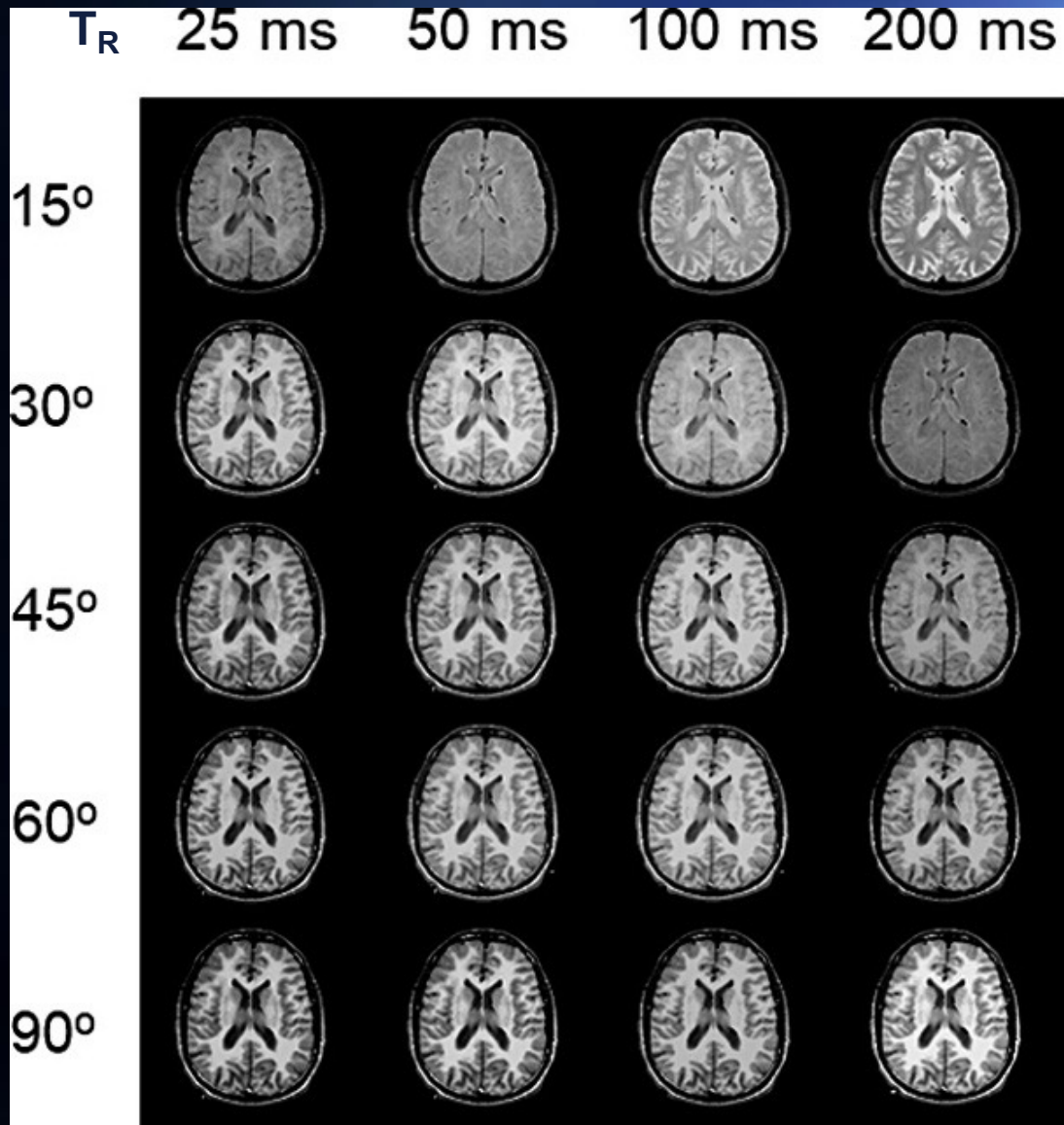
# Flip angle and $T_R$

- ✓ Given a short  $T_R$ , the signal is larger if  $\alpha < \pi/2$
- ✓ Given small  $\alpha$ , the signal recovers faster but it saturates at a lower value



$T_R$  defines  
the total acquisition time

# Gradient-Echo Pulse Sequence

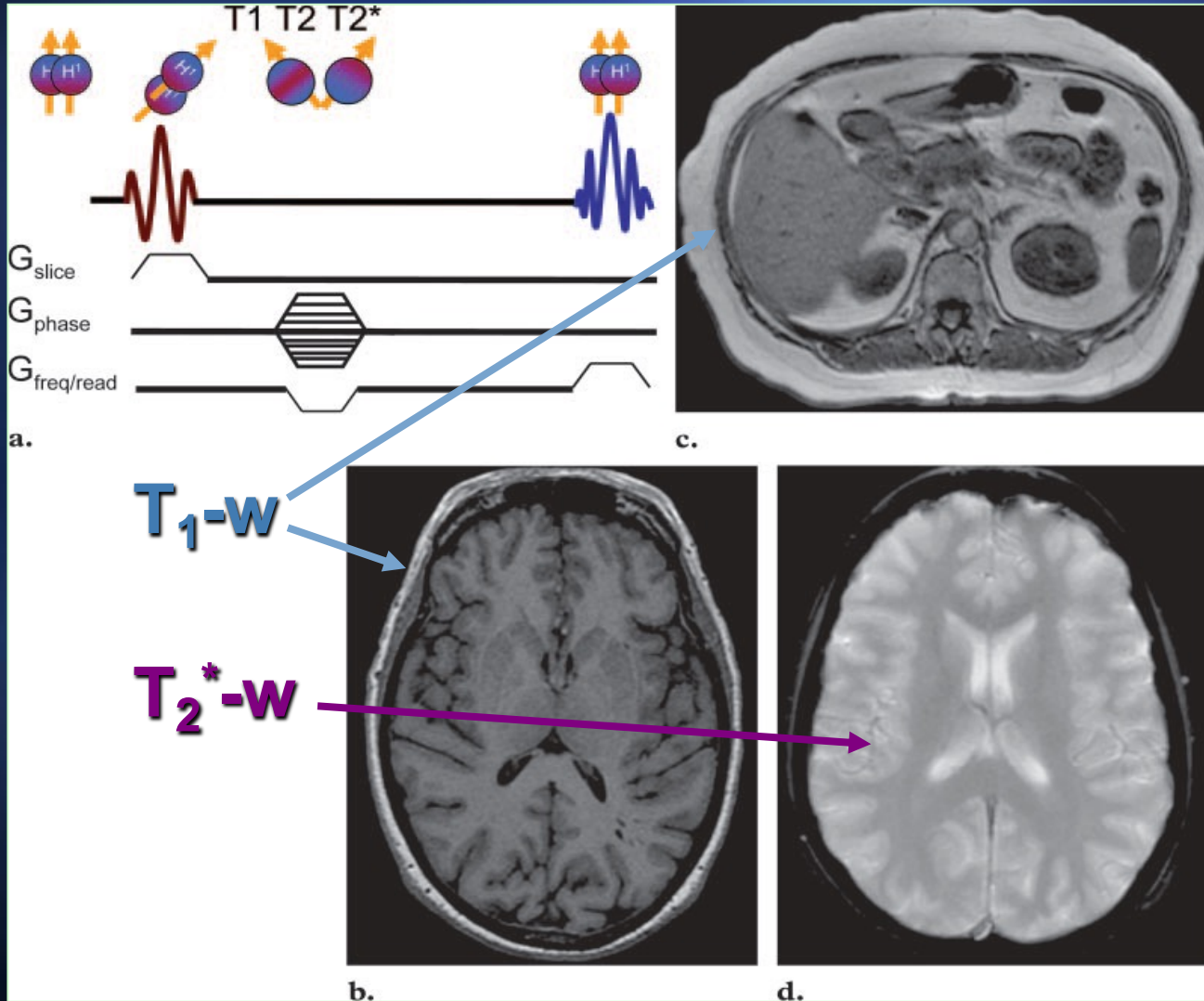


$$I \propto \rho(x,y) \frac{\left(1 - e^{-\frac{T_R}{T_1}}\right)}{1 - \cos\alpha e^{-\frac{T_R}{T_1}}} \sin\alpha e^{-\frac{T_E}{T_2^*}}$$

- ✓ large  $\alpha$  and short  $T_R$ :  $T_1$  weighting
- ✓  $\alpha$  reduced: weighting switches to proton density and then to  $T_2^*$ 
  - even with  $T_R$  as short as 100 ms



# Gradient echo sequence





# K-space

## Path through k-space

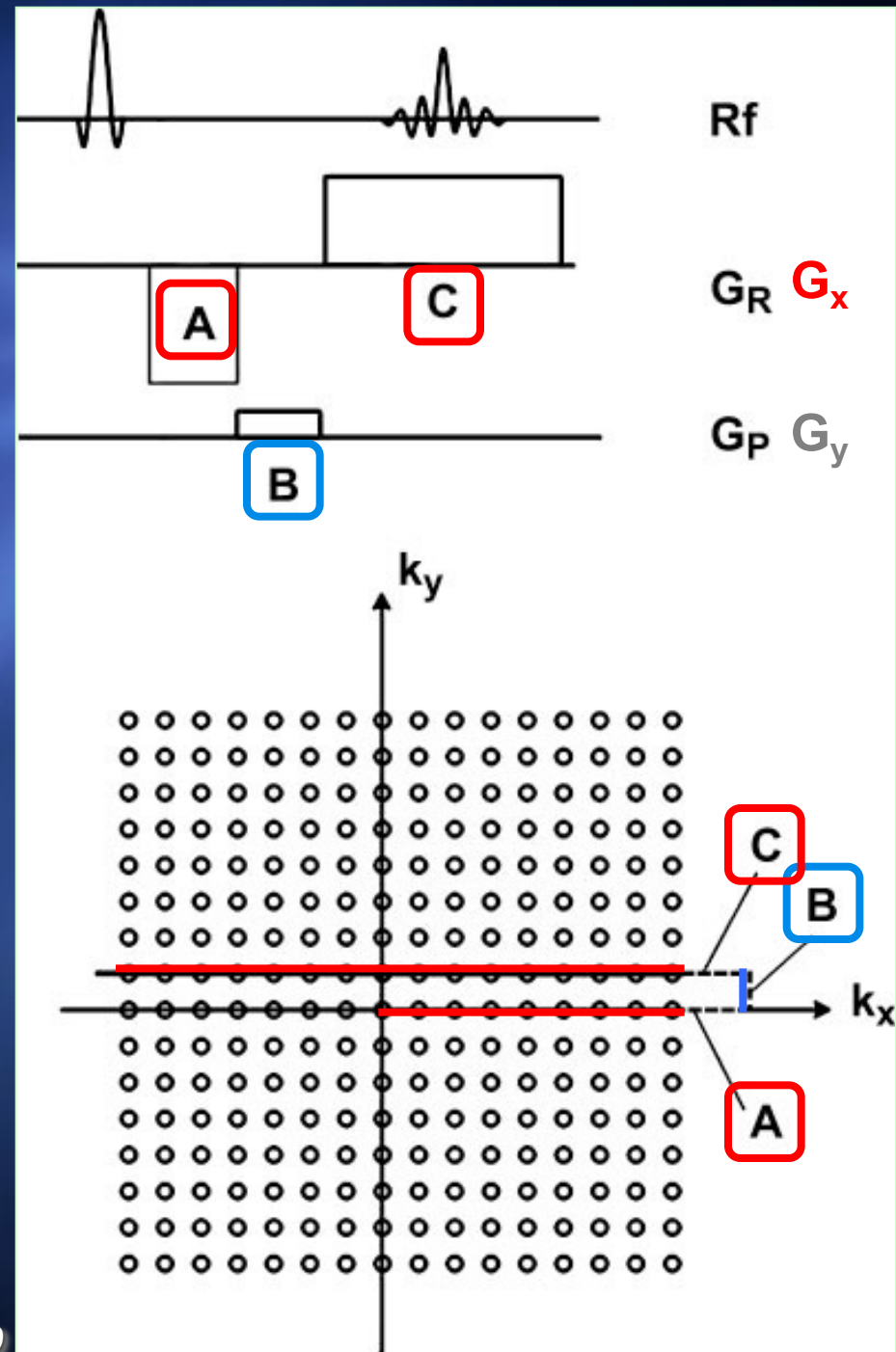
- $k_x = \gamma G_x t$  ;  $k_y = \gamma G_y T$

- ✓ The prewinding gradient **A** carries the k-space trajectory in the  $k_x$  direction out of the sampling area

- ✓ Phase encoding with **B** causes an offset of the trajectory in the  $k_y$  direction, where signal from one k-line is **read out under C**

- Data are acquired only during the last part (**red line**), and signal during A and B is discarded (**dashed line**)

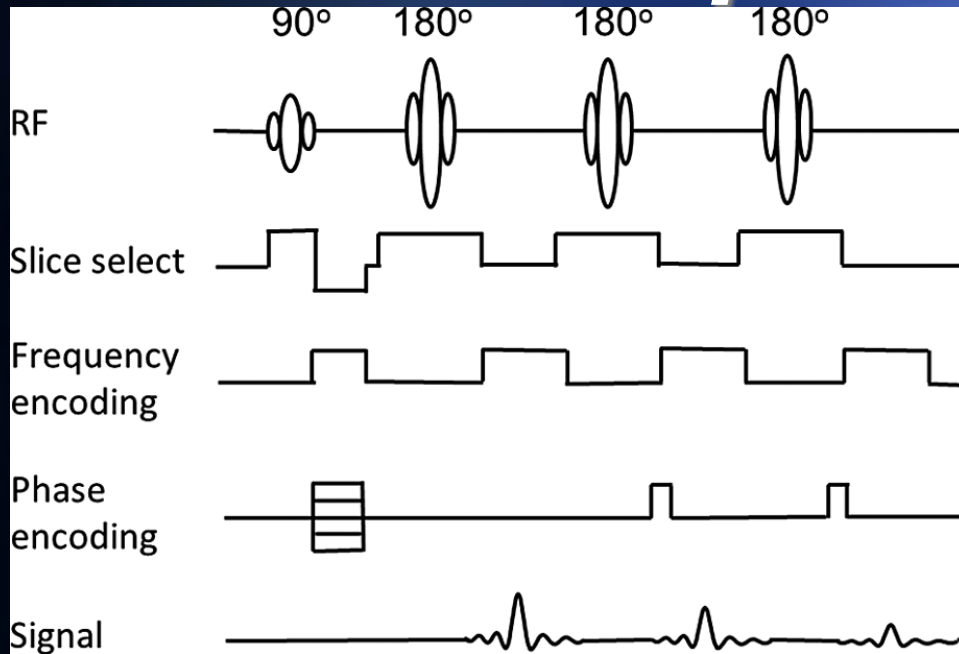
- ✓ The experiment is repeated with different **B**



# *Spoiler gradient*

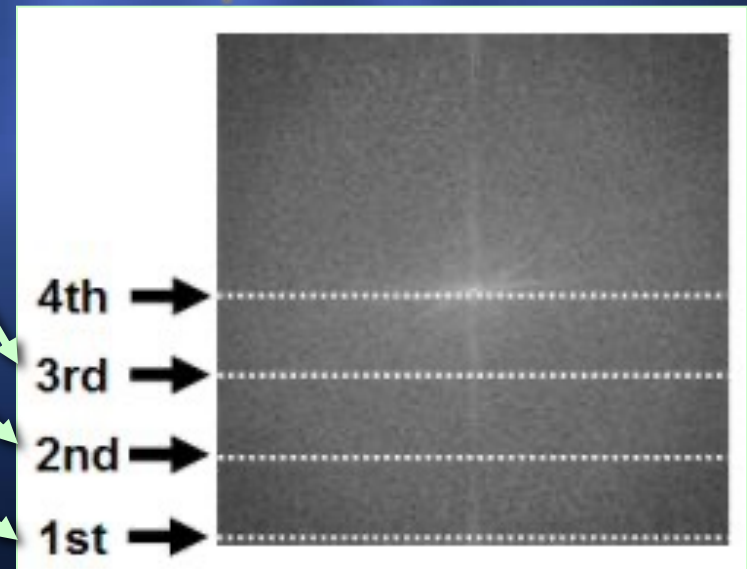
- ✓ If  $T_R$  is so short to be comparable to the  $T_2$  values of tissues the effect of subsequent excitation pulses on residual  $M_{xy}$  as well as on recovering  $M_z$  must be considered
- ✓ the simplest approach is to add a strong gradient pulse after acquisition of each echo signal
  - before the next excitation pulse
- ✓ This so-called 'spoiler gradient' completely dephases the residual  $M_{xy}$ 
  - so that we only have to consider  $M_z$

# *fast spin-echo (FSE) or turbo spin-echo (TSE)*



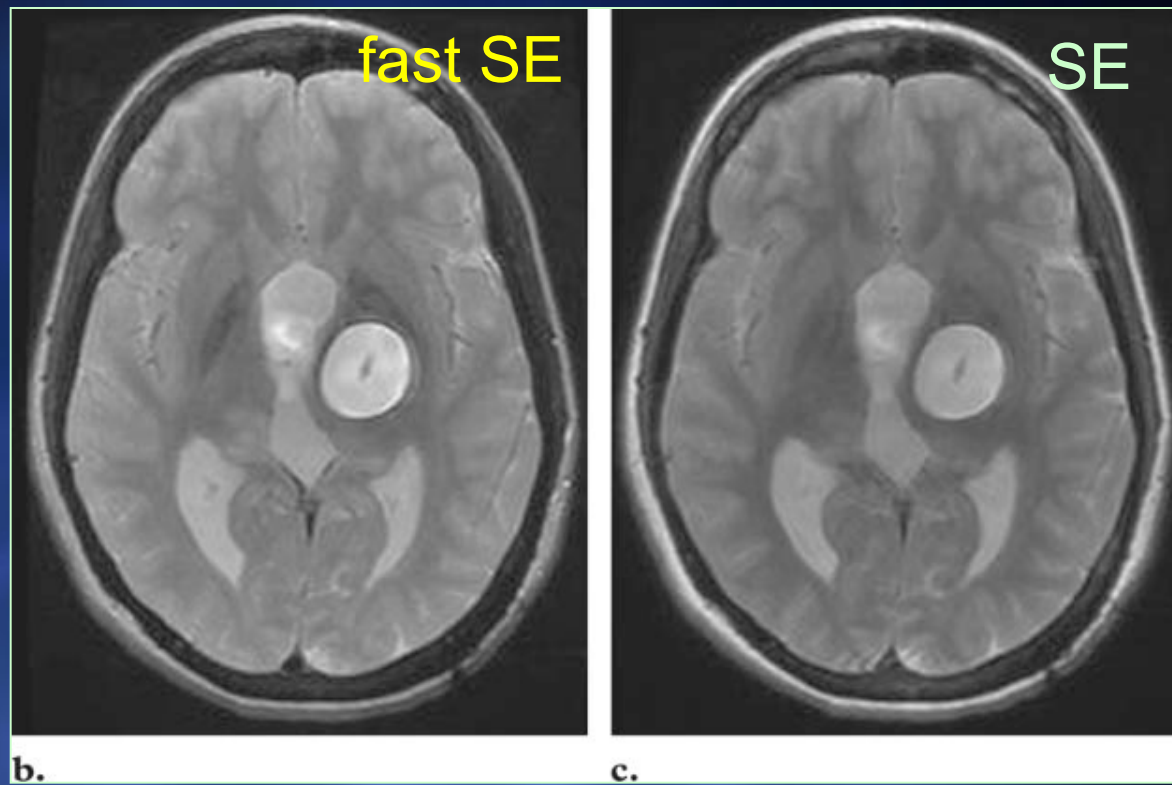
✓ before each echo there is an additional phase encoding gradient

- each echo is collected from a different line in k-space





# *fast SE*



b.

c.

b) Axial T2-weighted fast SE image

c) conventional SE image

provide comparable depiction of a brain tumor

✓ The acquisition time for conventional SE imaging was 7 minutes 17 seconds, whereas that for fast SE imaging with an echo train length (ETL) of 16 was 34 seconds

# *Fast SE*

- ✓ echo train length (ETL) or turbo factor (TF) is typically 4–32
- ✓ ETL is limited by the repetition time
- ✓ there is a trade-off with the number of slices imaged
  - interleaved excitation of multiple slices during the  $T_R$
- ✓ limit to the achievable ETL since echoes acquired at very long  $T_E$  contain very little signal

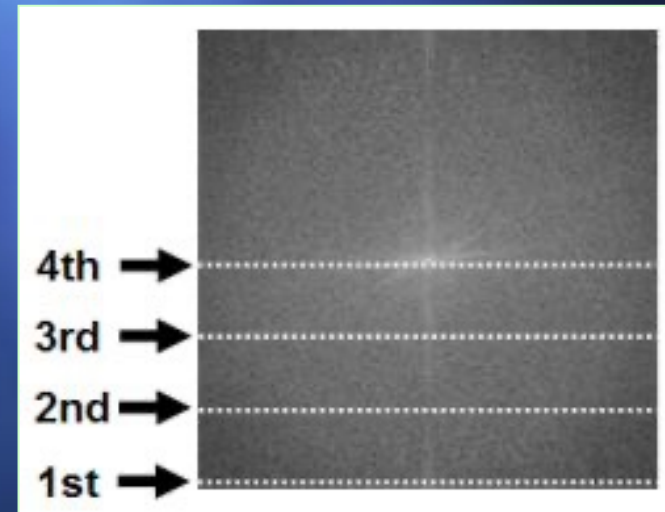


# Fast SE

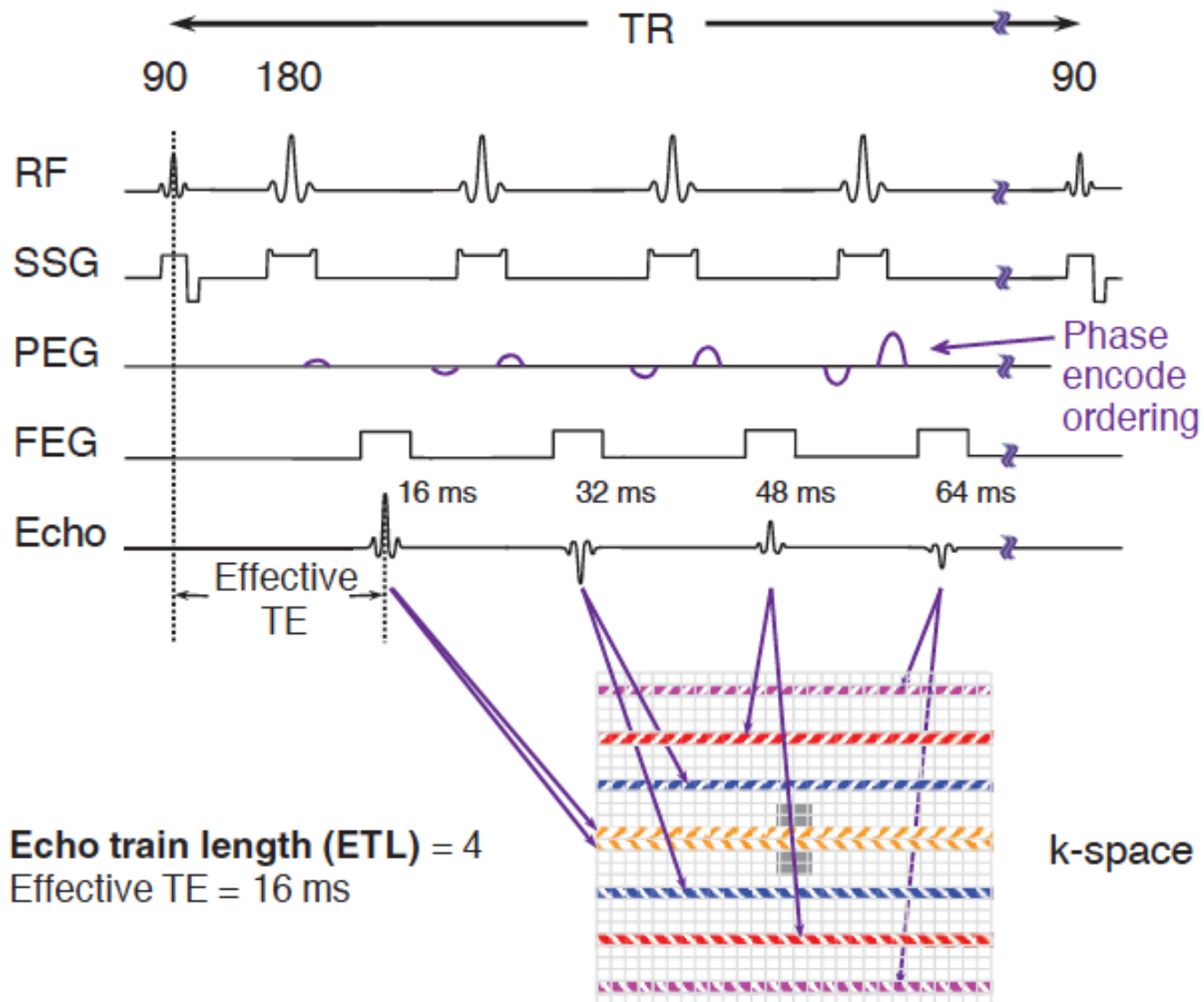
How much weighting is there when different lines in k-space have been acquired at different echo times?

The centre of k-space dominates the contrast properties of the image

- where magnetisation is in phase throughout the imaged object
- ✓ The degree of  $T_2$ -weighting in a fast SE image is dominated by echoes close to the centre of the  $k_y$ -axis
  - effective echo time or pseudo echo time



# Fast SE

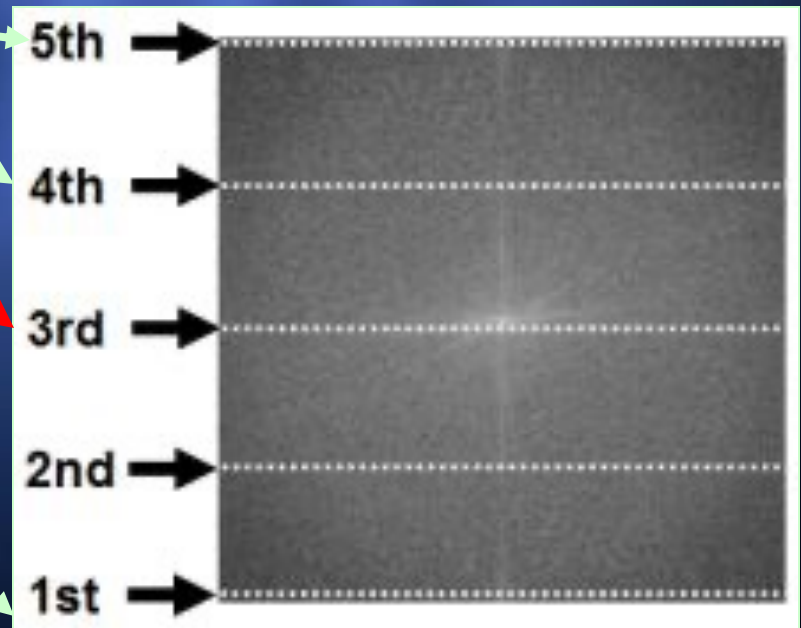
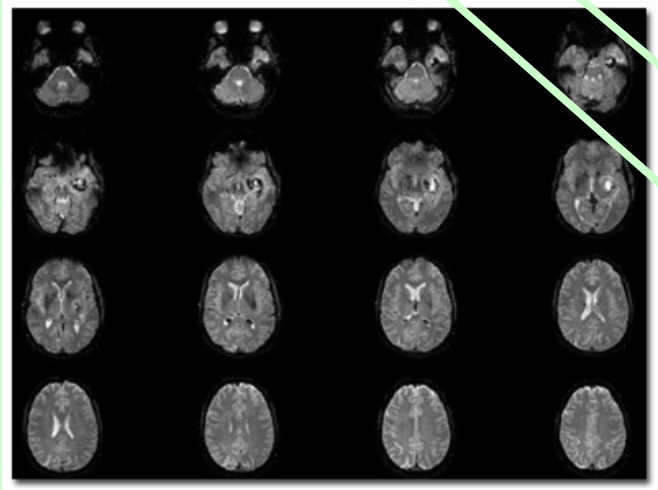
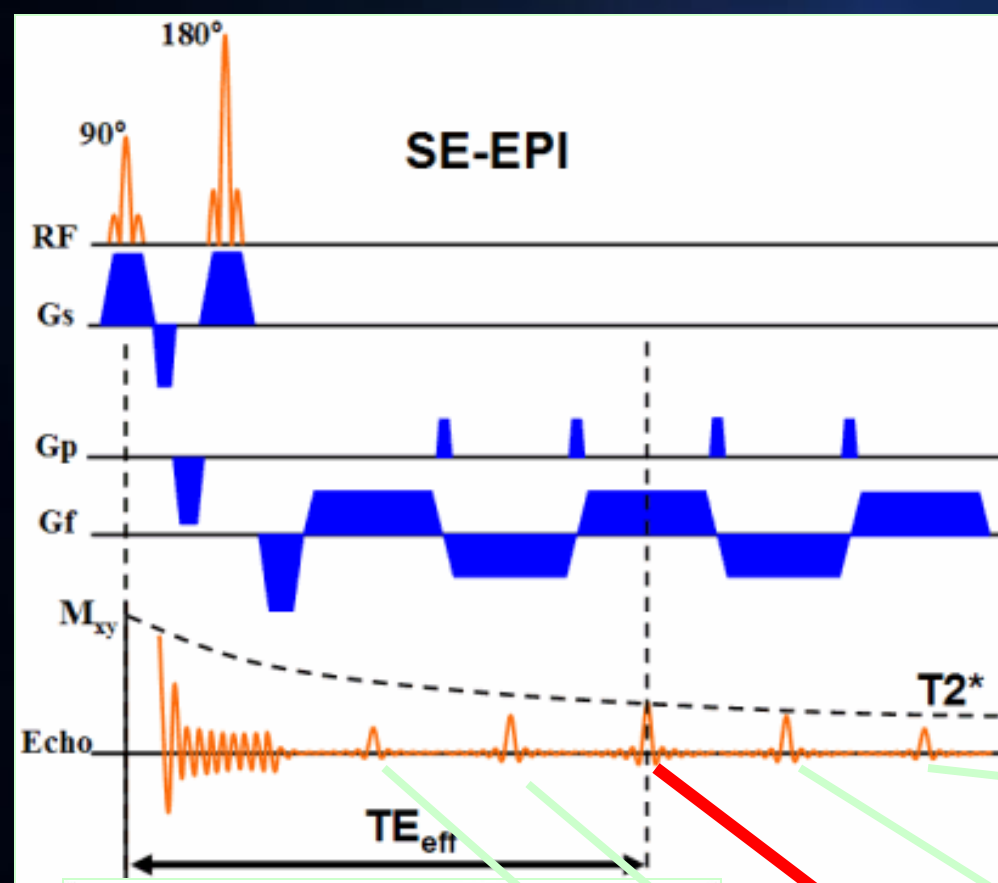


# *Fast SE*

- ✓ echoes acquired at long TE are typically positioned towards the edges of k-space
  - very long  $T_E$  contain very little signal
  - at high ETL this can become an issue because of the loss of high spatial resolution data and consequent image blurring
- ✓ The appearance of the image is determined by the k-space trajectory
  - identical sequences with different phase encoding patterns can produce very different images!

# Echo planar imaging

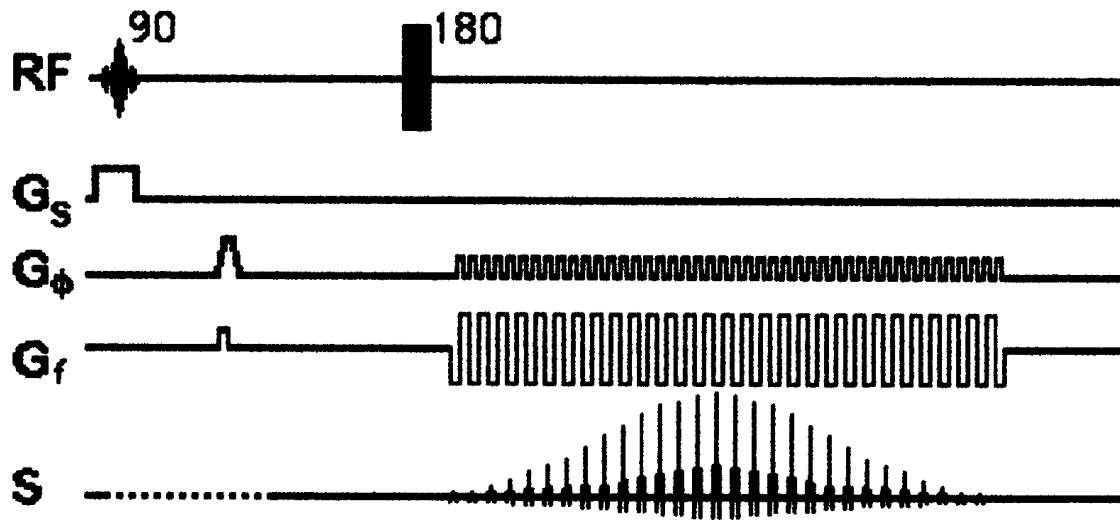
- ✓ “Single shot” acquisition
  - A mixture of SE and GRE
- ✓ A brain in 3 s
  - 16 slices





# Echo Planar Imaging

- ✓ Oscillating gradients to explore all the k-space
- ✓ The phase encoding gradients are summed after each echo acquisition
- ✓ Gradient frequencies ~ KHz
- ✓  $\Delta T_e \sim \text{ms}$
- ✓ Artifacts are visible if the gradients are not perfectly repeated

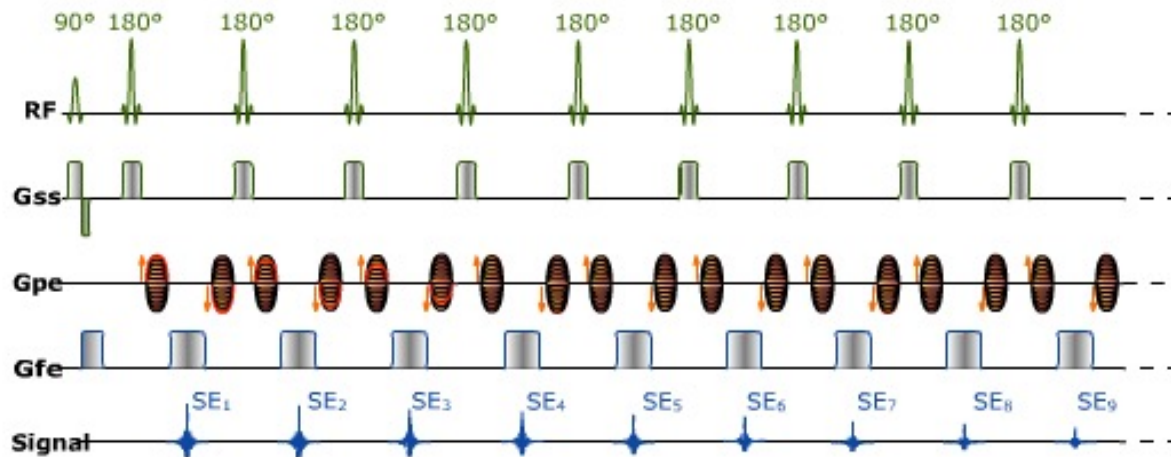


$$k_x = \gamma \int_0^t ds G_x(s)$$
$$k_y = \gamma \int_0^t ds G_y(s)$$

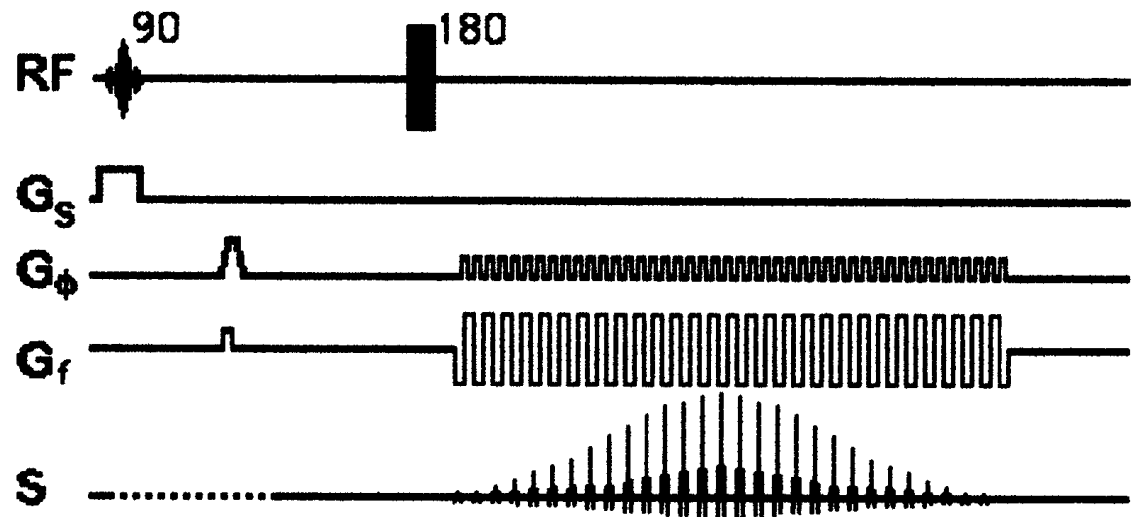


# Ultrafast Spin Echo & EPI

Multiple echos  
in order  
to fill the whole  
k-space



© IMAIOS 2015

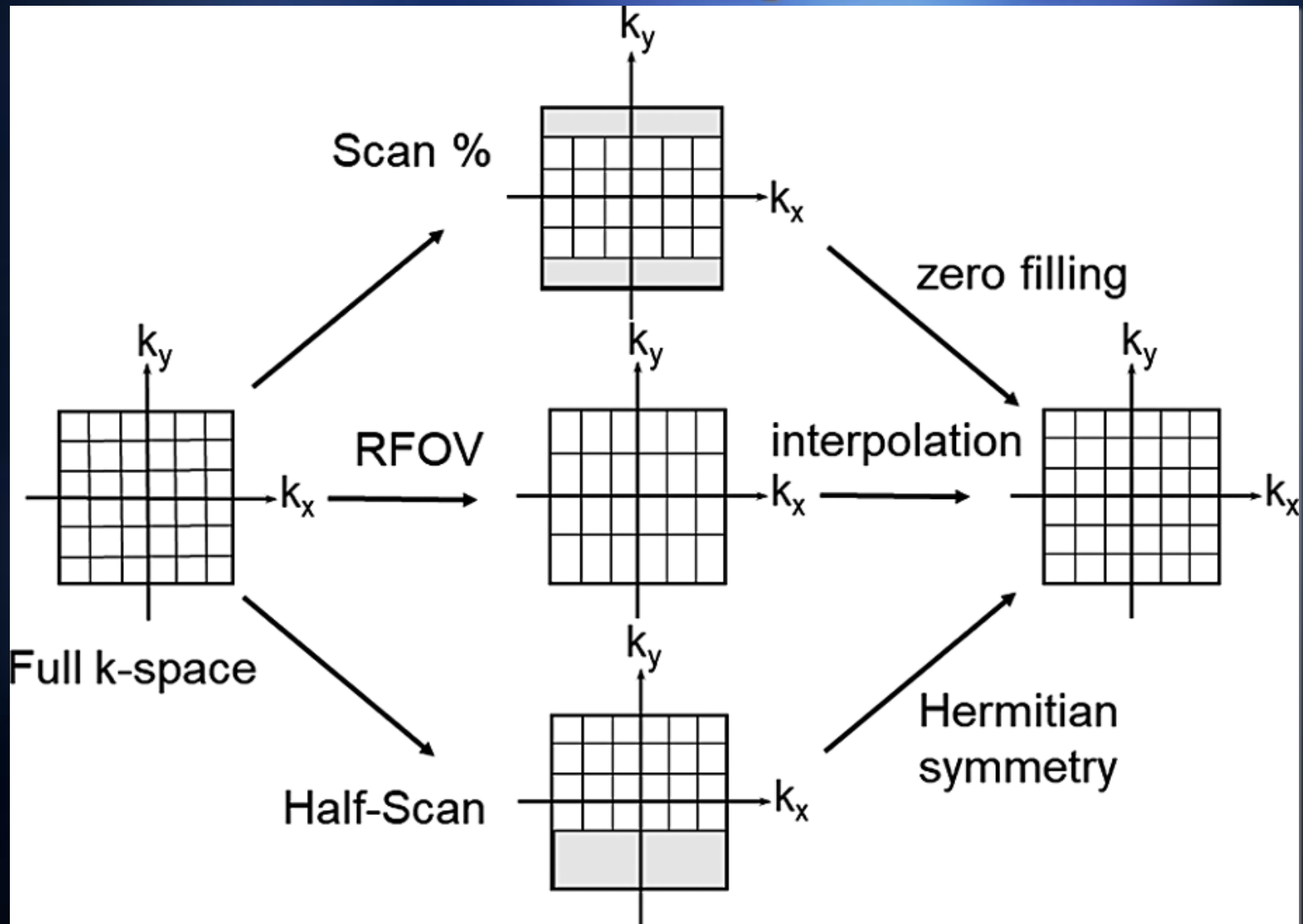


# ***Acquisition time reduction strategies***

How to reduce the acquisition time with minimum effects on the image quality !?!?

- ✓ **Multislice sequence**
- ✓ **Gradient echo sequences**
  - Reduced flip angle and reduced Tr and TE
- ✓ **Multiple rows filling from 1 excitation pulse**
  - Fast and EPI imaging
- ✓ **Incomplete filling of the k-space**
- ✓ **Parallel imaging**
  - How an artifacts becomes a resource !

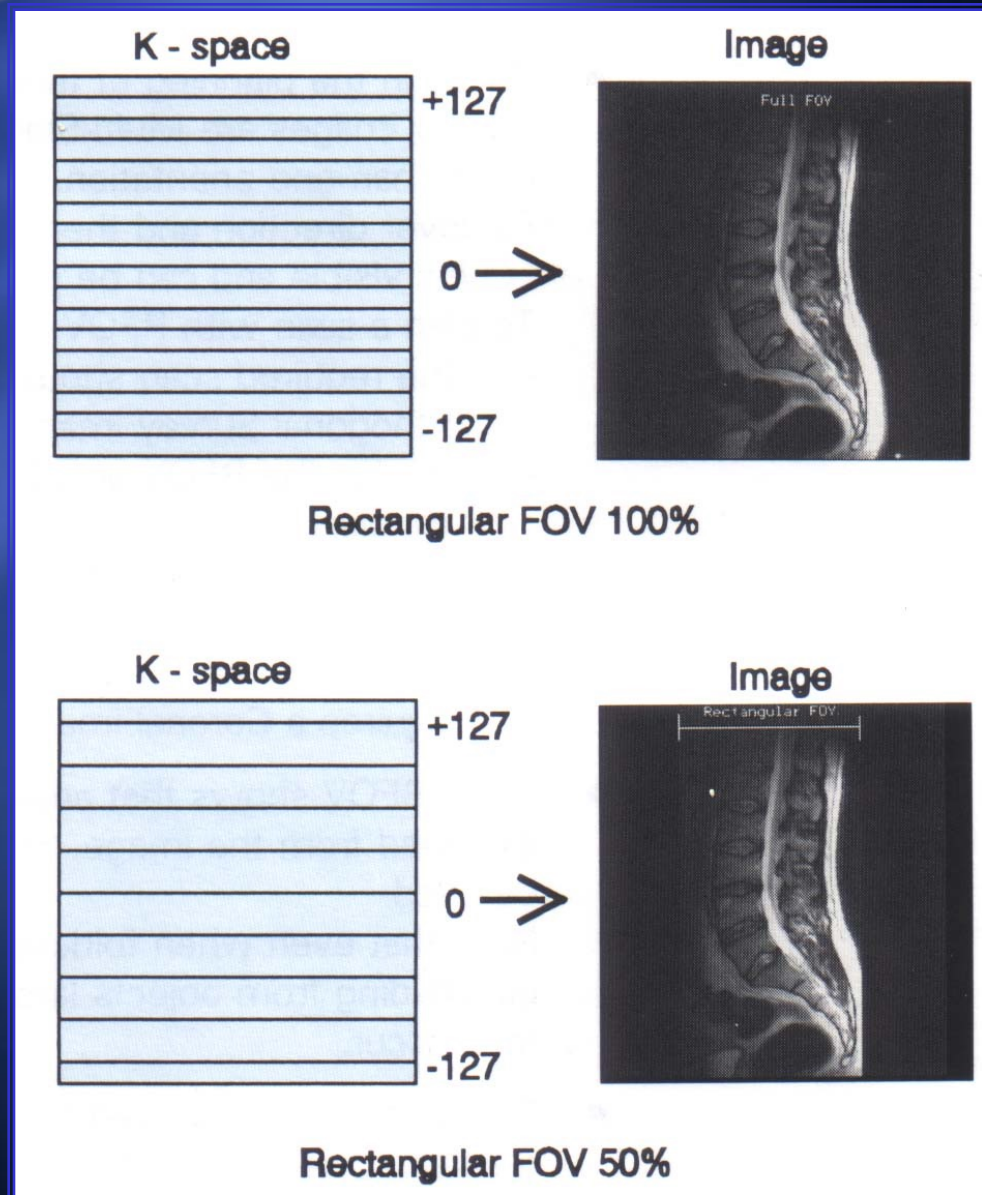
# Reduced acquisition of the $k$ -space



# rectangular FOV

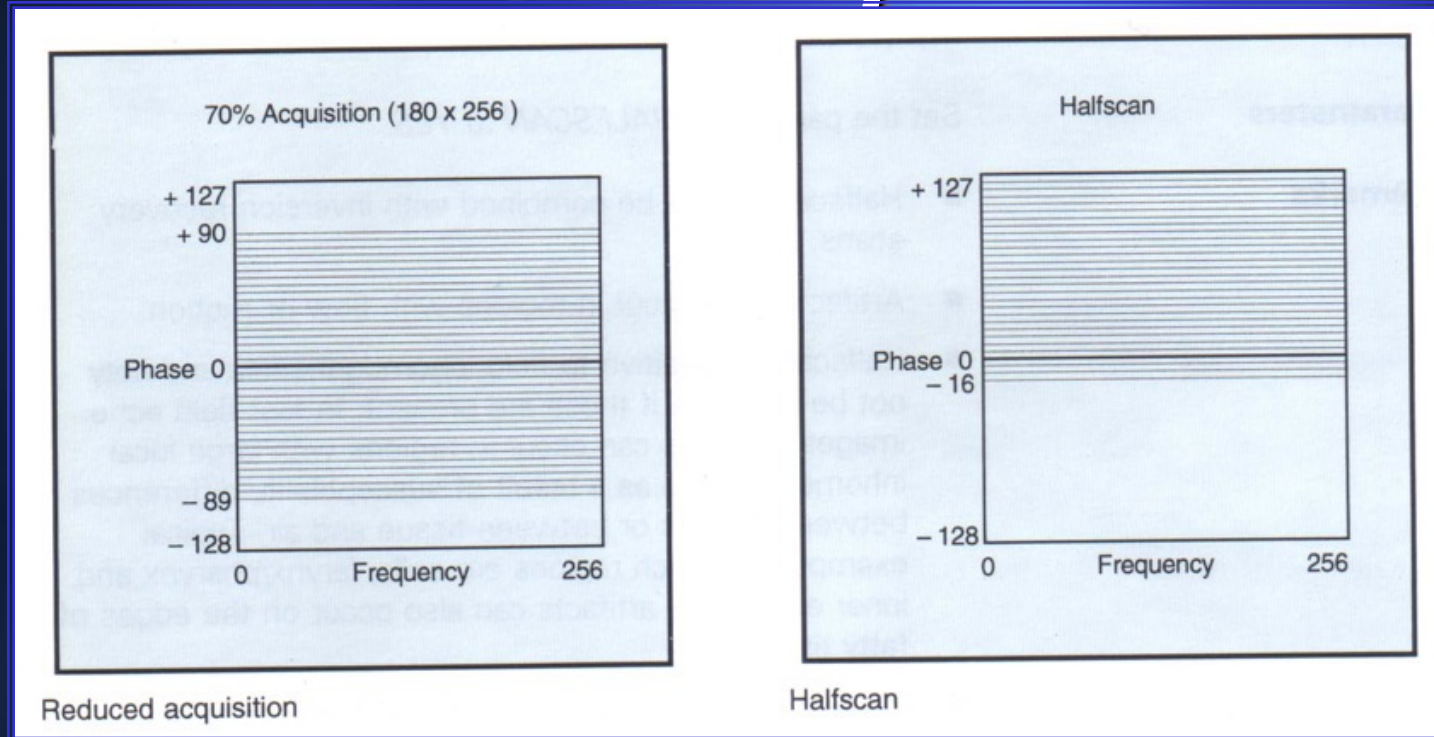
✓  $\Delta k_x$   $\Delta k_y$  may be not equal

- FOV rectangular in the image space
- $\Delta k_x$  sampling step in the K-space





# Reduced acquisition of the $k$ -space



- ✓ Acquiring the central part of the  $k$ -space the spatial resolution is affected
- ✓ In halfscan approach the spatial resolution is preserved but the SNR is decreased

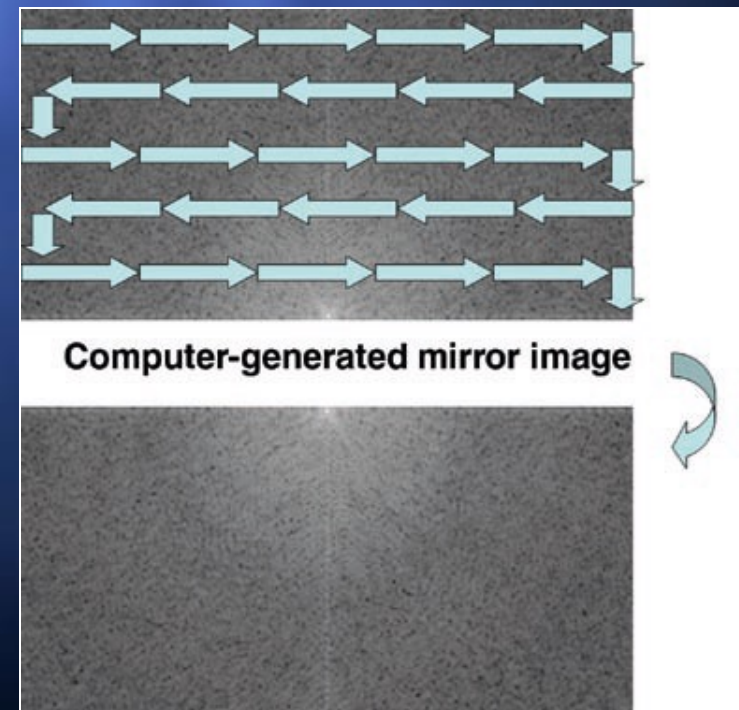
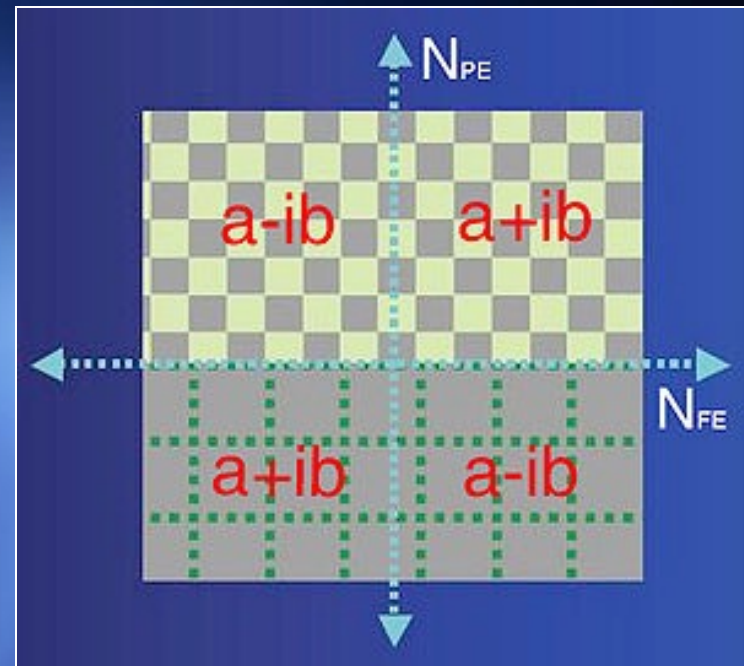


# Half scan

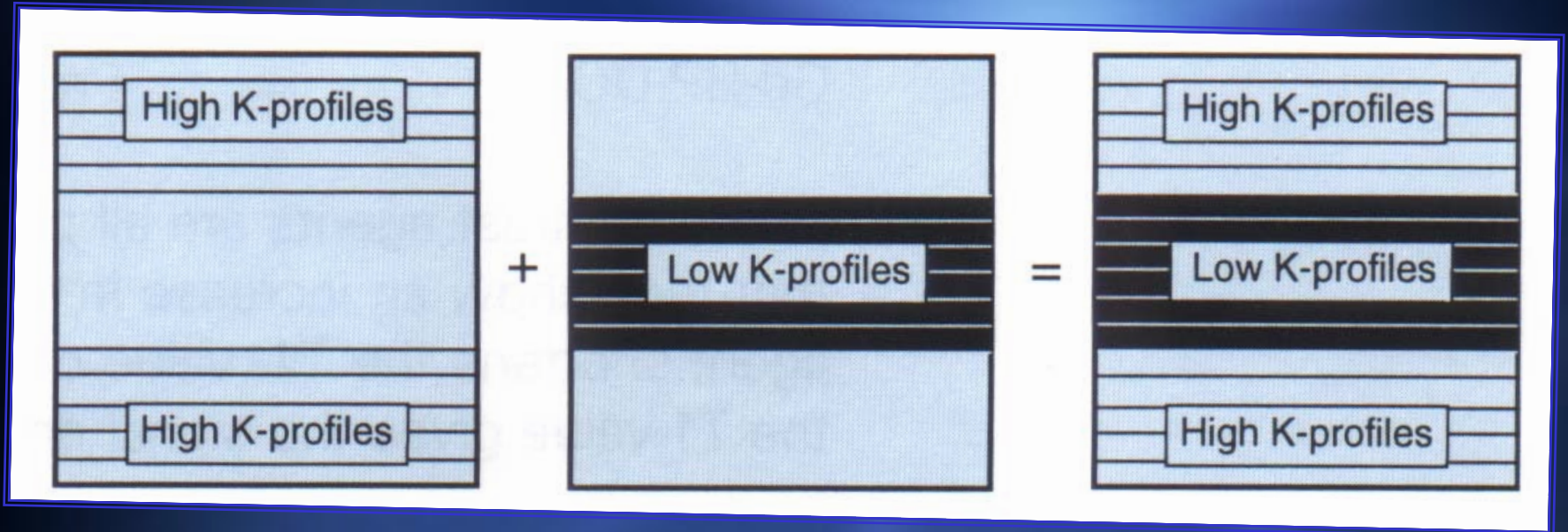
- ✓ K-space: complex conjugate symmetry

$$S(-k_{FE}, -k_{PE}) = S^*(k_{FE}, k_{PE})$$

- ✓ Time-saving mechanism at the expense of the SNR
- ✓ In single-shot techniques (EPI) it is a powerful technique
  - Due to the very low signal of the late echos ..



# *Reduced acquisition of the k-space (II)*

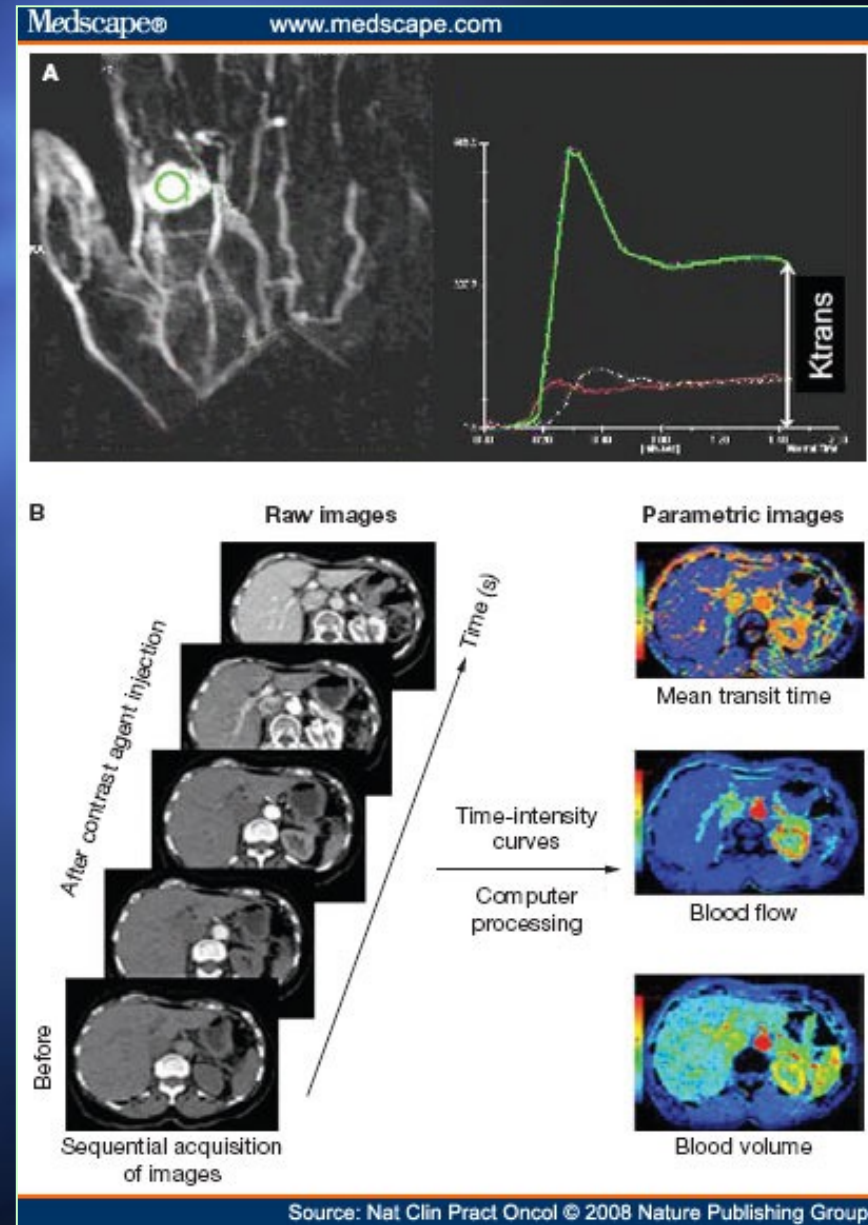


- ✓ **KeyHole technique:**
  - Fast dynamic acquisition



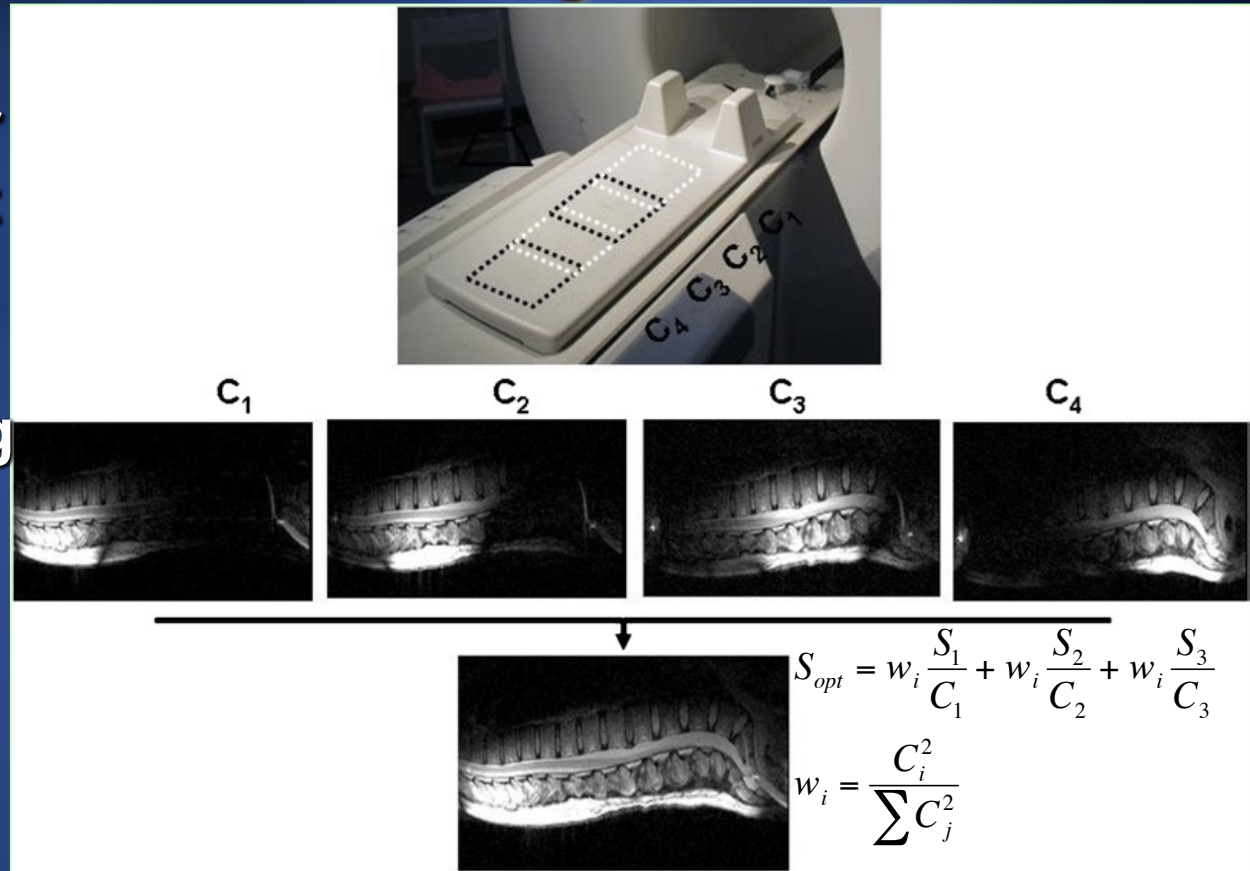
# Dynamic contrast-enhanced MRI

- ✓ DCE-MRI enables analysis of blood vessels generated by a tumor
- ✓ The T1w images are taken after an injection of the contrast agent
- ✓ The concentration of the contrast agent is measured, when it passes from the blood vessels to the extra-cellular space of the tissue and whether it goes back to the blood vessels
  - it cannot enter inside the cells



# Phased array coils

The use of multiple receiver coils to augment the time-consuming Fourier encoding has reduced acquisition times significantly



- ✓ A typical spine coil which uses an array of four coils arranged linearly which reflects the extended anatomy of the spine
- ✓ Each coil element  $C$ , produces a separate image shown as  $C1-C4$ , the bottom image is an optimal combination of these images performed on a pixel by pixel basis using equation



# *Artificial Contrast Agents*

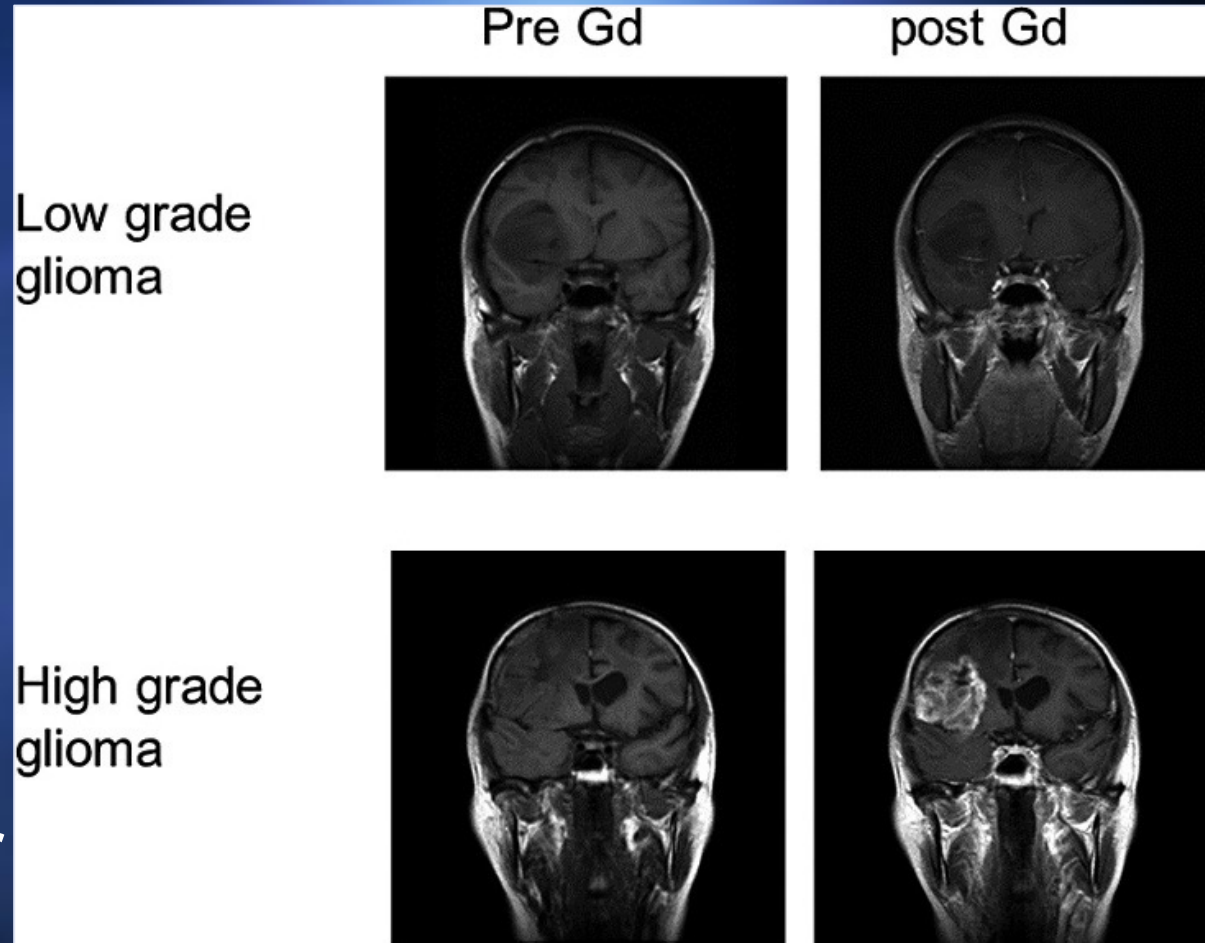
- ✓ it may be advantageous to introduce exogenous agents which affect contrast in a way that reflects some pathological or physiological process
- ✓ MRI agents generally work by reducing the T1 and/or T2 of tissue in which they reside
  - resulting in local changes in signal intensity

# *Artificial Contrast Agents*

- ✓ the contrast agents in clinical use are intravascularly injected agents based on the gadolinium ion (Gd)
  - gadolinium-based contrast agents
    - ❖ GBCAs
- ✓ Gadolinium is a paramagnetic ion, which affects signal primarily by shortening  $T_1$ 
  - Because it is a toxic heavy metal, it is used in a chelated form, in which each gadolinium ion is embedded in a molecular 'cage' to reduce toxicity

# Artificial Contrast Agents

- ✓ In a high-grade glioma, where the blood-brain barrier is compromised, GBCA crosses into the tumour and causes signal enhancement
  - due to  $T_1$  shortening
- ✓ Being able to differentiate tumour grade makes a great difference to the patient's



# ***Artificial Contrast Agents***

## **applications categories**

- ✓ **To change signal from selected tissues or areas of pathology**
  - usually on the basis of vascularity
  - usually signal increases
- ✓ **To enhance signal from flowing blood**
  - contrast-enhanced MR angiography
- ✓ **To eliminate signal from organs that might otherwise obscure structures of interest or cause artefacts**
  - such as the bowel
- ✓ **For dynamic studies of perfusion or function**
- ✓ **In emerging areas such as molecular imaging**



# ***GBCA safety***

- ✓ a very favourable safety profile compared to agents used in x-ray imaging

However

- ✓ guidance limiting the use of GBCAs in renal failure patients
- ✓ in 2014 it emerged that there are permanent changes in signal intensity in some parts of the brain following repeated GBCA administration, believed to result from retention of Gd
  - there are no known clinical consequences

**one-off administration of GBCA as part of an MRI examination appears to be safe**