# ARTEFACTS: PROBLEMS, SOLUTIONS AND NEW IDEAS

Artefacts are features appearing in an image that are not present in the investigated object

- In MRI, artefacts test our understanding of the underpinning physics
  - if this is understood the artefacts can be avoided, reduced or compensated for

Moreover

 a number of artefacts, or the solutions developed to minimise them, provided the seeds of advanced MRI techniques

Each data point in the k space contains information from the entire selected slice

or selected volume in 3D imaging

- Interleaved multislice imaging is used to cover the whole of an organ or body area of interest
- Movement during acquisition affects all the slices in the investigated volume
  - not a limited number of them
  - The acquisition time of an MRI dataset is typically a few minutes

Movement during acquisition affects all the slices in the investigated volume

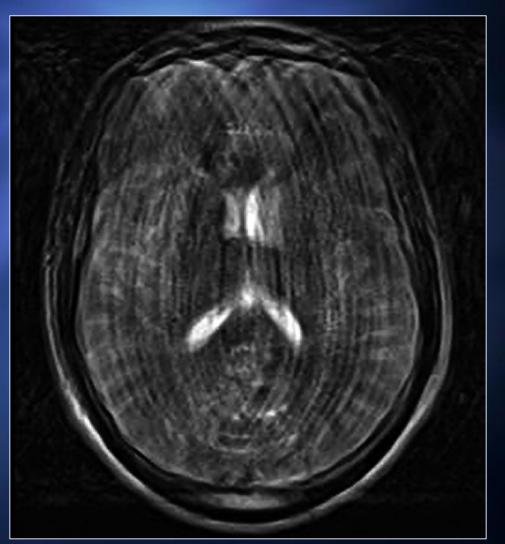
- not a limited number of them
- The acquisition time of an MRI dataset is typically a few minutes

it is of utmost importance to optimise patient positioning

In an uncomfortable position increases the probability of movement during the examination

Patient movement disrupts the relationship between phase and position This can result in faint displaced images

• known as ghosts

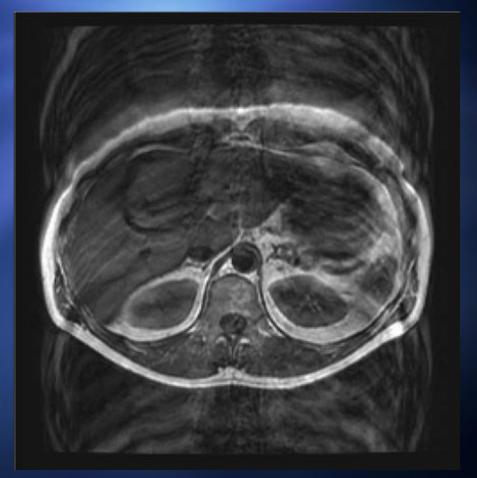


phase encoding gradient in the left-right direction

 In medical imaging, some types of motion cannot be avoided
 breathing, the cardiac

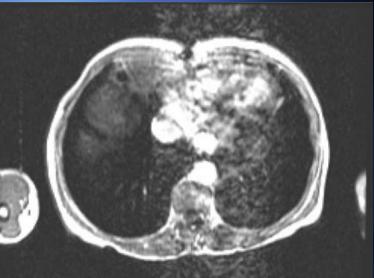
cycle, and blood flow

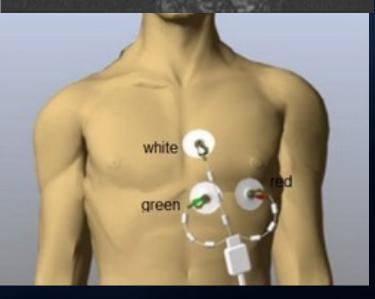
 there are some techniques to minimise the resulting artefacts



# **Cardiac Motion**

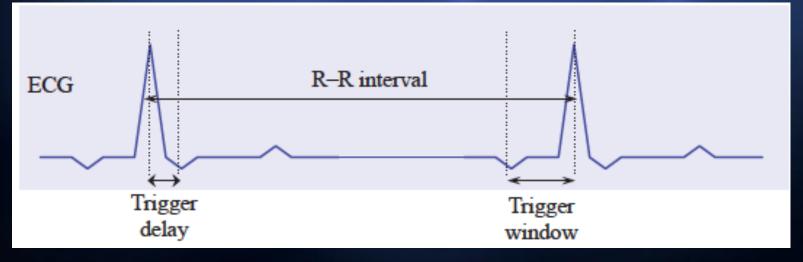
- Synchronizing the sequence to the cardiac cycle
  - known as 'gating'
- Special electrocardiogram
   (ECG) non-metallic electrodes
  - to reduce artefacts on the images
  - to avoid local skin heating
  - The ECG wires typically have high impedance
    - reducing the chance of RF burns for the patient





## **Cardiac Motion**

- The peak of the R wave is used to trigger the next acquisition
- Each line of K space is acquired at the same point in the cardiac cycle
- $\checkmark$  The T<sub>R</sub> is determined by the heart rate (HR)
  - typical HR of 75 beats per minute (bpm)
  - TR ~ 800 ms or multiple of the cardiac cycle

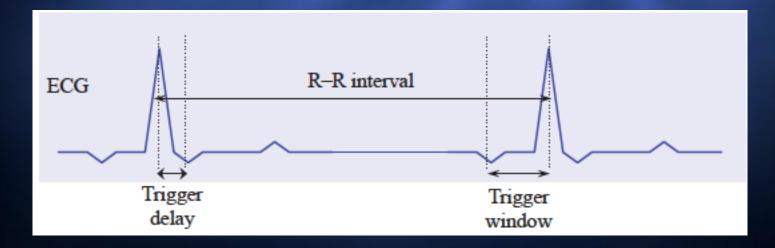


## **Cardiac Motion**

Each line of k space is acquired at the same point in the cardiac cycle

- Ghosting is removed
- Cine imaging of the cardiac movement

<u>https://www.imaios.com/en/e-Courses/e-MRI/Image-guality-and-artifacts/motion-artifact-remedies</u>

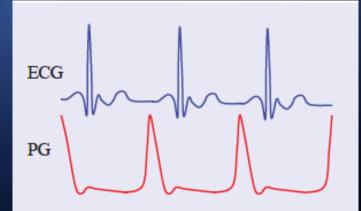


## **Peripheral Gating**

Cardiac gating by detecting the arterial pulse of blood in the patient's finger

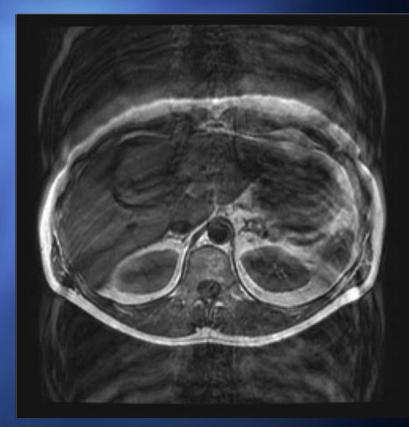
- the blood vessels are very close to the skin
- an infrared light detector pick up the increased volume of blood as the arterial pulse reaches the extremity
   \$sufficient to provide a trigger for the MR sequence
- Peripheral gating is useful for removing pulsatility artefacts in the brain and spine





## **Respiratory Motion**

- Breathing motion causes ghosting thoracic and abdominal imaging
- ✓15 s scan time or less than can be acquired during a single breath-hold
  - The acquisition can be split up so that packages of slices are acquired in separate breathholds
    - The position of the liver and kidneys is more reproducible if the breath is held in expiration

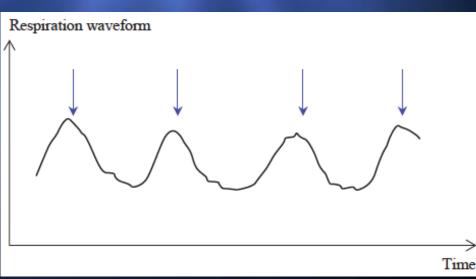


Careful coaching with the patient is needed before the examination

# **Respiratory gating**

- The respiratory cycle can be monitored using a 'breathing belt' strapped around the thorax
- It this generates an electric signal dependent on the pressure of the thorax on the belt
  - allowing breathing to be monitored





https://www.imaios.com/en/e-Courses/e-MRI/Image-quality-and-artifacts/motion-artifact-remedies

# **Respiratory gating**

Respiratory gating can then be applied in order to perform acquisition of k-space lines during the end-expiration phase

• when chest motion is minimal

 $\sqrt{T_R}$  is defined by the breathing cycle

- the interval between 2 consecutive triggers
- a normal breathing rate of about 10–15 breaths per minute gives an effective TR of at least 4000 ms

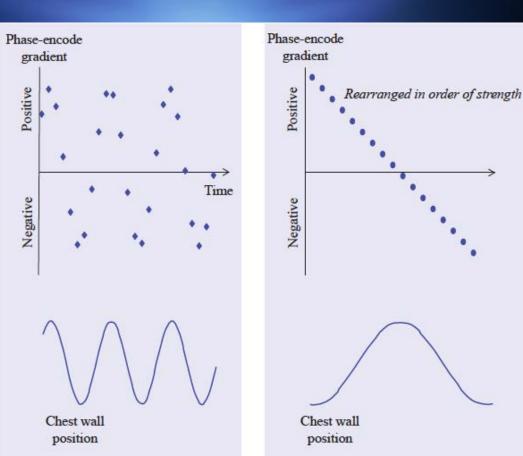
respiratory gating useful for PD or T2-weighted imaging

# **Respiratory compensation**

**Respiratory-Ordered** Phase Encoding (ROPE)

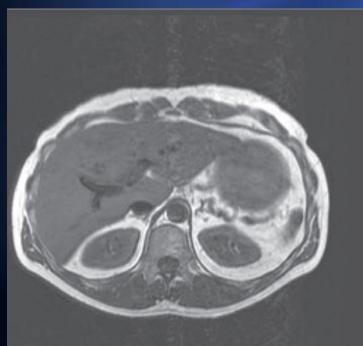
- ✓ k-space lines are acquired during the whole respiratory cycle
- ✓ similar phase encoding gradient amplitudes are used with the thorax at similar positions
- This avoids abrupt variation in the position of chest structures between closely-spaced https://www.imaios.com/en/e-Courses/ek-space lines
  - minimising artefacts

MRI/Image-quality-and-artifacts/motionartifact-remedies

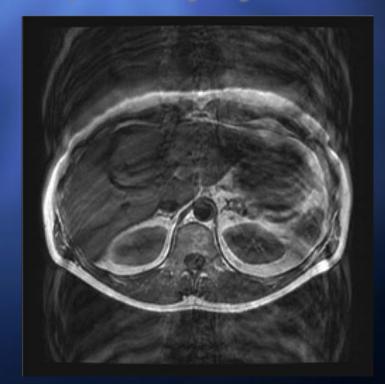


## **Respiratory compensation**

 In respiratory gating only the quasi-stationary phase of the cycle is used for acquisition
 ROPE is more effective because data are acquired throughout the respiratory cycle



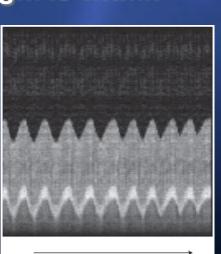
**ROPE** sequence



## **Navigator echoes**

- The navigator 'signal' is from a single column of voxels used to monitor the position of the diaphragm
  - ID signal acquired from 1 echo
  - Strong contrast between the lung and the liver shows up clearly
  - The boundary automatically detected and controls the acquisition: the image data are acquired when the diaphragm is within certain spatial limits
    - typically 2–4 mm
  - ROPE acquisition <u>without</u> respiratory belt

The navigator signal is spatially encoded in one direction: the other direction is time



Time



Tracking volume

## What is the best ?

- a breath-hold is usually the best choice
   breath-hold for morphology and perfusion
- scans
- navigators for coronary arteries and highresolution viability
- ROPE can also be used with classic SE or GE sequences in abdomen imaging
- respiratory gating
  - Too long T<sub>R</sub>

## Flowing blood and cerebrospinal fluid

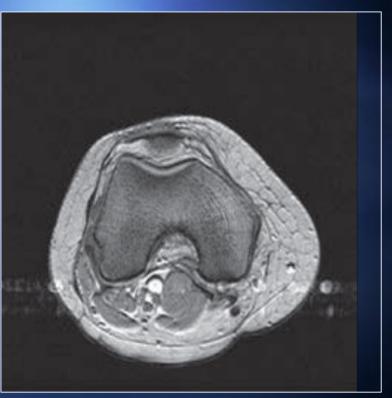
In the flowing blood and CSF represent large numbers of moving spins sometimes they undergo pulsatile flow • more complex than flow at a constant velocity when spins are moving in the time interval between the RF excitation pulse and echo acquisition, they accumulate a phase shift different from the phase expected for a stationary spin

# Flowing blood and cerebrospinal fluid

 when spins are moving they accumulate a phase shift different from the phase expected for a stationary spin

 in the phase encoding direction we observe ghosts of the vessel

✓ if the motion is pulsatile the intensities of the ghosts have a periodic modulation



## Flow compensation gradient

The phase-encoding gradient waveform can be modified to cancel the additional phase shift associated with flow

 The drawback is that the duration of a flowcompensated phase-encoding gradient is longer limiting the minimum echo time

https://www.imaios.com/en/e-Courses/e-MRI/MR-Angiography-Flow-imaging/ilowcompensation

Sometimes the flow artefacts overlie the region of clinical interest : the simplest solution is to swap frequency and phase encoding directions and repeat the acquisition

artefacts are still present but the direction will be rotated by 90°

https://www.imaios.com/en/e-mri/image-guality-and-artifacts/motion-and-ghostingartifacts-remedies

# **Motion Artefacts from Flowing Blood**

### Two effects

- ✓ an in-flow effect which produces high signal within blood vessels on GE images
  - new protons flow into the imaging slice during the T<sub>R</sub>
- velocity-induced phase effects
  - reduce the blood signals
  - create ghost images of vessels in the phase encode direction



# **Motion Artefacts from Flowing Blood**

### SE sequence

### Signal loss within the blood vessel

### Characteristic dark-blood appearance of spinecho images

https://www.imaios.com/ en/e-Courses/e-MRI/MR-Angiography-Flowimaging/flowphenomena-mri

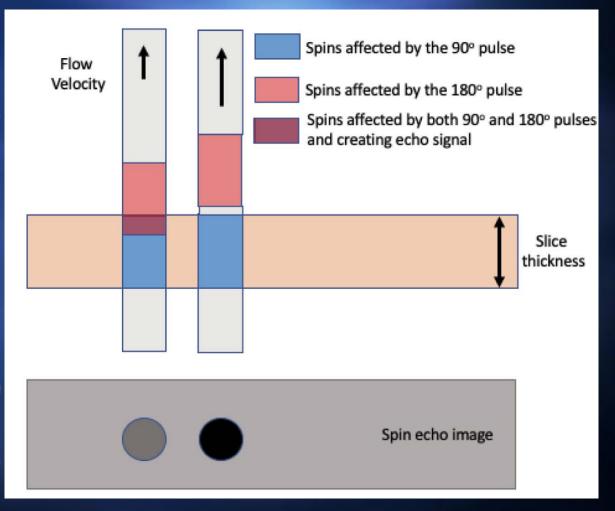


# **Motion Artefacts from Flowing Blood**

## SE sequence

The bolus of blood within the imaging slice is replaced by a second bolus of fresh unsaturated blood during the gap between the 2 pulses

- The first bolus receives a 90° pulse but not the 180°
- The second receives only the 180° pulse
- No bolus produces an echo



# Motion Artefacts from Flowing Blood GE sequence

on GE images blood vessels have high intensity

 easily produce ghost images in the phase encode (PE) direction

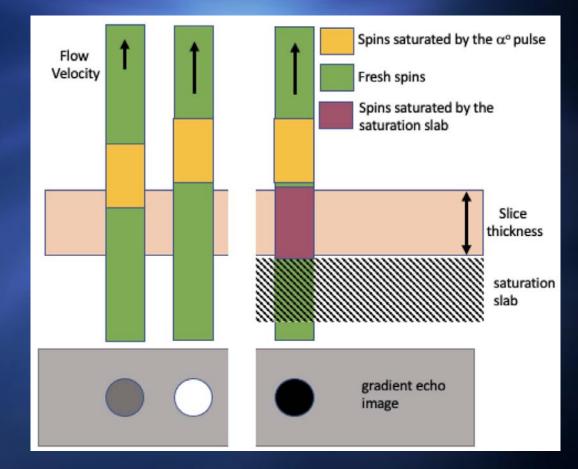


https://www.imaios.com/en/e-Courses/e-MRI/MR-Angiography-Flow-imaging/time-of-flight-mra

### Motion Artefacts from Flowing Blood **GE** sequence ✓ an excitation pulse and Flow Velocity the echo is formed using the gradients Excited bolus of blood always contributes a signal, provided it is still within the gradient volume For every subsequent slice there is a fresh bolus of blood within the slice fully relaxed magnetization

To avoid flow or movement artefacts

 spatial saturation slabs just outside the field of view or in the slice direction

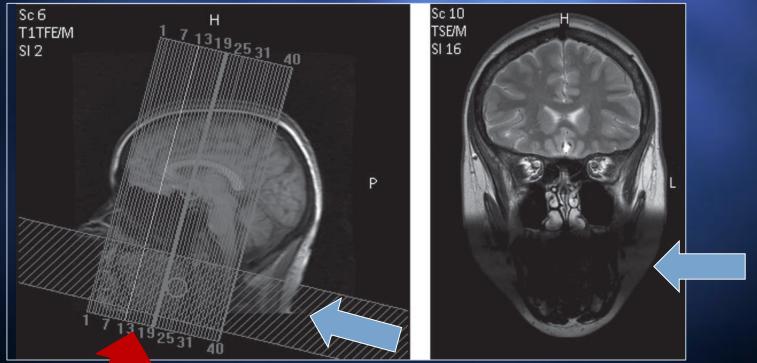


To avoid flow or movement artefacts

- Spatial saturation slabs just outside the field of view or in the slice direction
  - A 90° pulse to all the tissues within the slab immediately before the RF excitation pulse for the imaging sequence
  - After the saturation pulse a large gradient pulse is applied to diphase the protons
     \*leaving no signals from the tissues in the sat band
  - <u>https://www.imaios.com/en/e-Courses/e-</u> <u>MRI/Image-quality-and-artifacts/motion-artifact-</u> <u>remedies</u>

To avoid flow or movement artefacts

spatial saturation slabs just outside the field of view or in the slice direction



### Planned Coronal slices

### Saturation band

To avoid flow artifacts

spatial saturation slabs in the slice direction

- Placed above and/or below the image FOV
- sat bands remove arterial and/or venous blood flow

Pay attention to the flow direction to make sure you put the sat band in the right place!

 REST is used if a moving organ in the FOV prevents correct visualisation of the region of diagnostic interest

## Water and Fat Frequency Shift

✓ The resonance frequencies of water and fat protons are slightly different

chemical shift

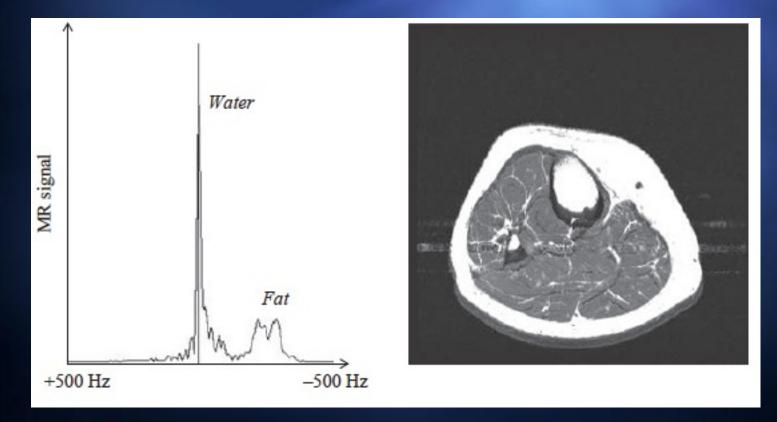
$$\omega_0 = \gamma B_0$$

The cloud of moving electrons surrounding each molecule creates a magnetic field of its own

- Depending on the structure of the molecule
- ✓ as a result the local magnetic field experienced by the H nucleus differs from B<sub>0</sub>
- The frequency shift between different molecules is proportional to magnetic field strength
  - at 1.5 T the water—fat shift is about 220 Hz

# Water and Fat Frequency Shift

<u>https://www.imaios.com/en/e-Courses/e-</u>
<u>MRI/Image-quality-and-artifacts/chemical-shift</u>



# water—fat shift and fat suppression

**Chemical Shift Selective Saturation (CHESS)** 

- an alternative to STIR for fat suppression
- immediately before the imaging pulse sequence a selective 90° pulse is applied at the fat proton frequency only
- ✓ the resulting transverse magnetisation is completely dephased using a spoiler gradient
- Therefore there is no fat magnetisation for the subsequent excitation pulse to nutate
  - hence no fat signal

## Water and Fat Frequency Shift artefacts

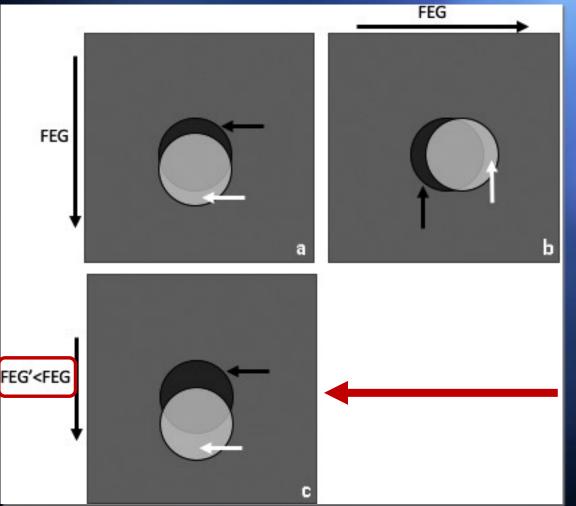
The water-fat shift is a source of an artefact in the frequency encoding direction at the interface between water and fat

- e.g. kidney and surrounding adipose tissue
- Fat and water signals are displaced from each other along the frequency encoding direction

Resulting in a hyperintense edge where the signals from water and fat overlap and a dark edge on the opposite side of the organ

Due to a signal void

# Water and Fat Frequency Shift This artefact is known as the chemical shift artefact of the first kind



larger displacement with a weaker gradient and hence smaller pixel bandwidth

## Water and Fat Frequency Shift

- The displacement may be expressed in pixels
- It is calculated as the ratio between the water-fat shift and the pixel bandwidth
  - the range of frequencies contained in each pixel

which depends on the gradient strength used

The pixel bandwidth influences the signal to noise ratio (SNR) of the image

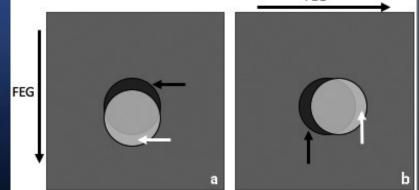
 Using a smaller bandwidth improves the SNR, but also results in a larger water-fat shift artefact

# Water and Fat Frequency Shift

The balance between these effects must be optimised by the user taking into account the anatomical region

 for example, water—fat shift is crucial in abdominal imaging but negligible in brain studies

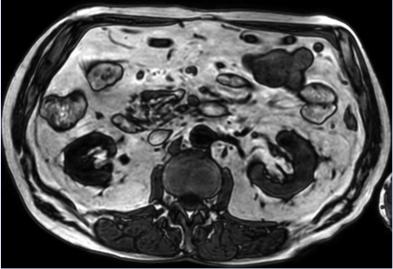
If there is doubt as to whether a feature is a water—fat shift artefact, the simple way to solve the impasse is to re-acquire the image switching phase and frequency encoding directions



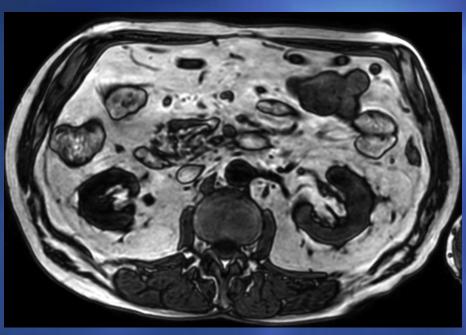
Chemical shift artefact of the second kind When both water and fat are present in the same voxel, the signal acquired depends on the relative phase of water and fat signals at the echo time

 if the relative phase is zero the signal is maximum

✓ if the relative phase is 180° the signal is the difference between water and fat signal intensities



## **Quantification of fat content**



The echo time needed to obtain in-phase or opposedphase signals depends on B<sub>0</sub>

 At 1.5 T, in phase signals occur at TE = 4.55n ms and opposed phase at TE = (2.3 + 4.55n) ms n is an integer

Comparing in-phase and opposed-phase imaging allows quantification of the fat component in tissue

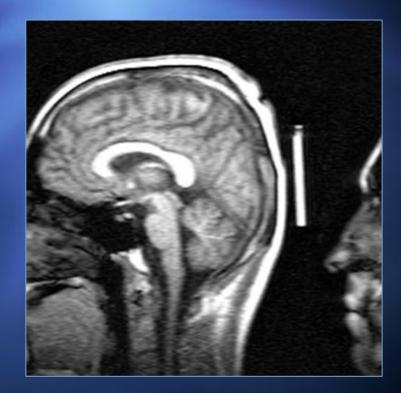
• known as the Dixon method

## Wrap-Around Artefacts

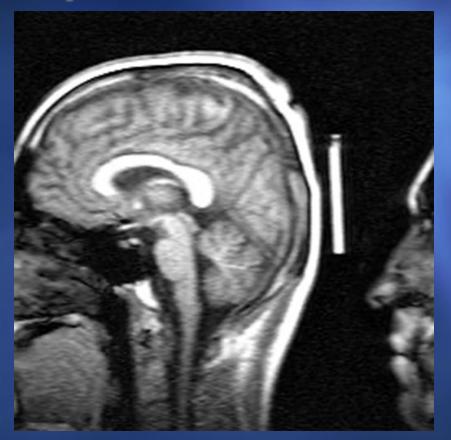
The spatial encoding process in MRI is known as frequency and phase encoding

Phase is a periodic variable between 0° and 360°

When the phase value reaches 360° it starts again from 0°, and in general phase \$\$\phi\$>360° maps to a value \$\$\$\$\$\$\$\$\$\$\$\$\$\$\$-360°



## Wrap-Around Artefacts

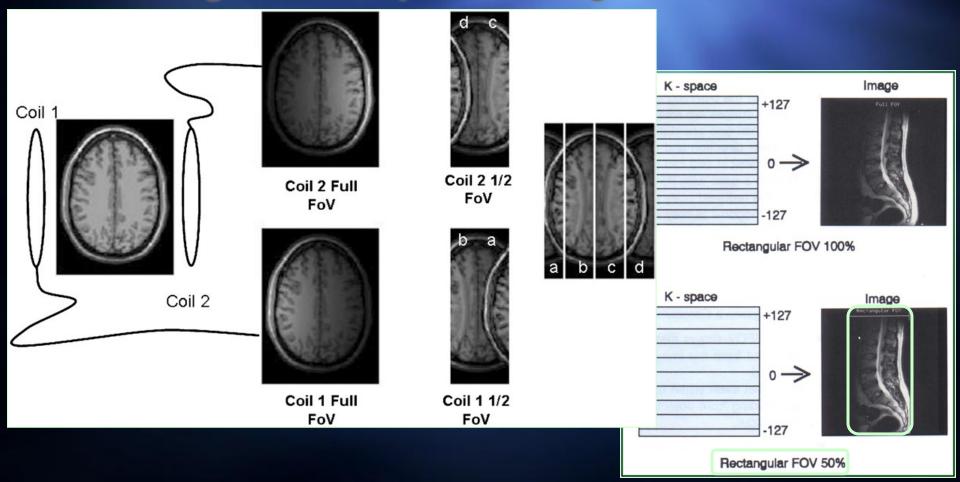


 in parallel imaging the acquisition of 'folded' images using a reduced FOV is done intentionally to reduce acquisition time

fewer k-space lines



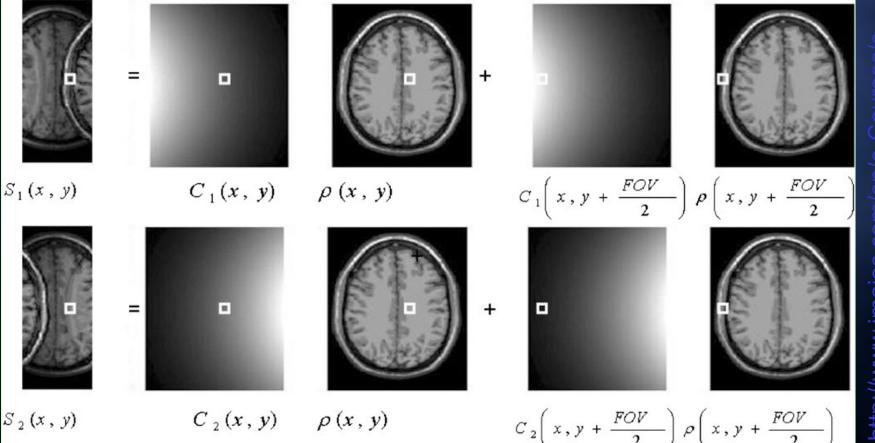
 In this example two local receiver coils are used for spatial discrimination
 But images are acquired using ½ FOV



# Parallel imaging (image domain)

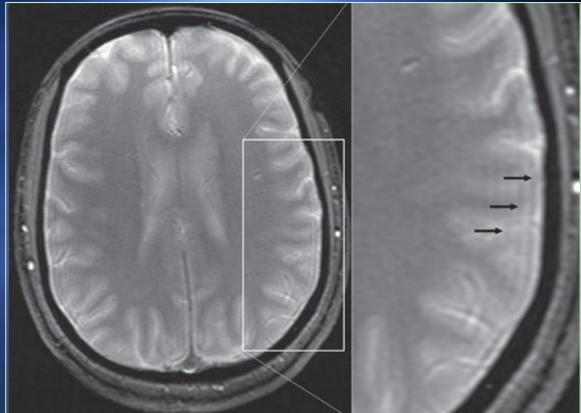
The aliased pixel in the image from coil 1, marked by a white square, S1 contains the sum of the spin density at location 1 multiplied by the coil sensitivity at location 1 and the spin density at location 2 multiplied by the coil sensitivity at location 2, likewise for coil 2.

If the coil sensitivities are known (the maps shown) then these equations are soluble for the spin density at both locations.



# **Truncation Artefacts**

**Truncation or Gibbs** artifacts: ✓ appear as parallel lines adjacent to high-contrast interfaces due to the Fourier transform from a finite sampled signal



Gallager et al AJR 2008; 190:1396–1405

http://www.imaios.com/en/e-Courses/e-MRI/Image-quality-and-artifacts/truncation

## Zipper and 'Corduroy' Artefacts

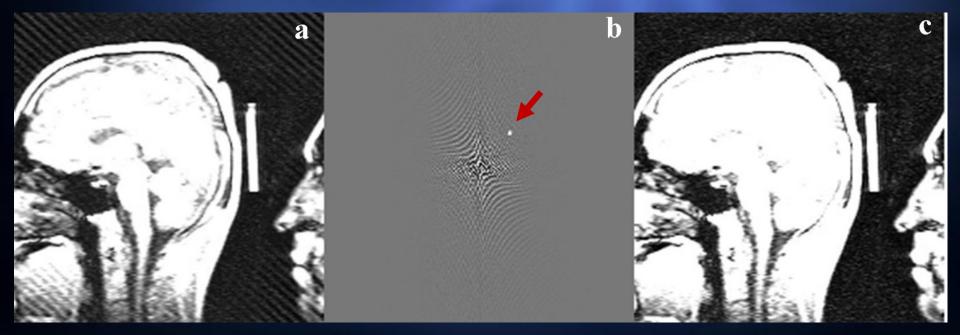
 Data errors in k-space, or detection of unwanted RF signals due to external interference, result in characteristic artefacts related to the Fourier transform

A single abnormal bright pixel in k-space is transformed into sinusoidal noise in image space: this is the source of the corduroy (or herring-bone) artefact, often caused by static electrical discharge

 One possible cause the humidity in the scanner room being too low

## **Zipper and 'Corduroy' Artefacts**

a zipper artefact due to an abnormal bright pixel in k-space visible in (b)(c) is the image reconstructed with this bright pixel removed



## **Equipment-Related Artefacts**

- There is another family of artefacts that arise when some component of the scanner is not working optimally
- If these artefacts are very evident, clinical use of the scanner may have to be stopped until it is repaired

A good programme of quality assurance (QA) may prevent this through early detection of small deterioration in image quality which can be addressed without interruption of clinical activity