

# Principi fondamentali della produzione di immagini in RM e protocolli di acquisizione

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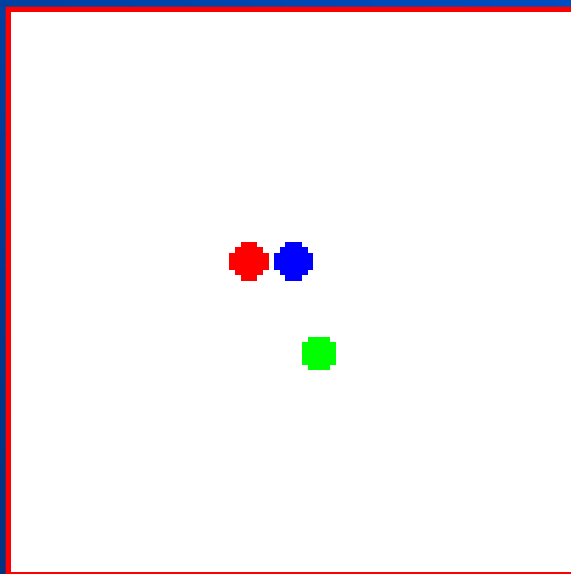


**MRI-LAB**

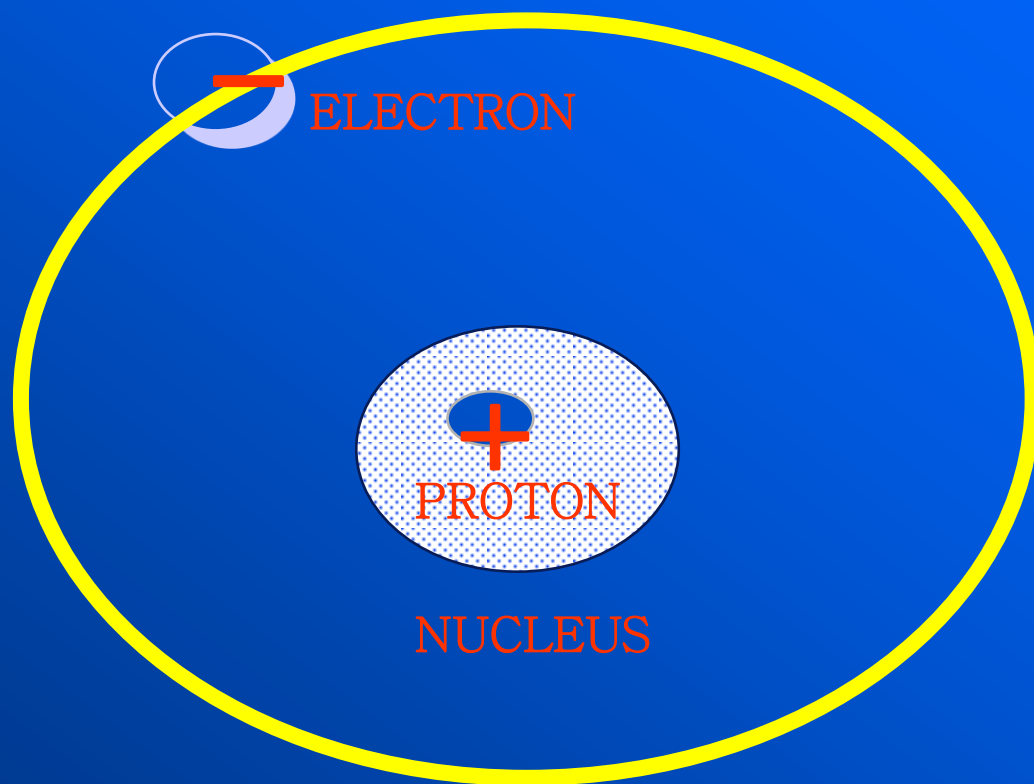


The phenomenon of Magnetic Resonance may be approached using different types of nuclei, however the atom of Hydrogen is generally utilized for creating MR images.

## The atom of Hydrogen

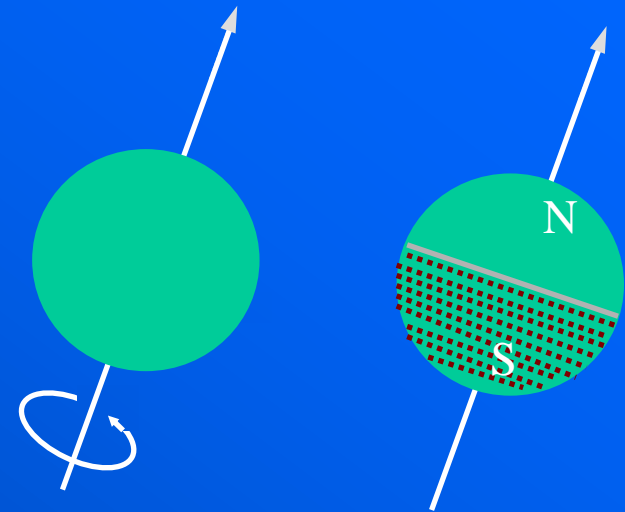


*Deuterium: 1 proton,  
1 neutron, 1 electron*





the nucleus is an electrically charged mass rotating on its own axis that generates a tiny magnetic field with its own direction and orientation. This phenomenon is called "*spin*" and is what gives the magnetic momentum  $\mu$  to the nucleus.



Gyromagnetic ratio  $\rightarrow \gamma = \frac{\mu}{p}$

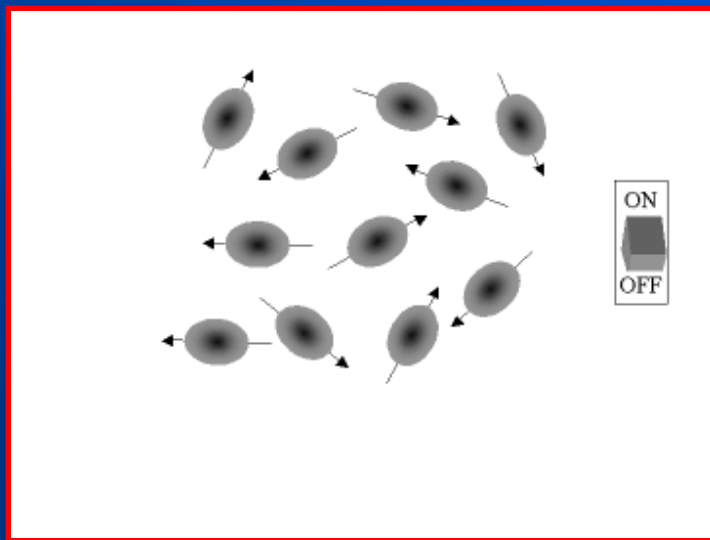
Magnetic momentum (due to a rotating electric charge)

Angular momentum (due to a rotating mass)

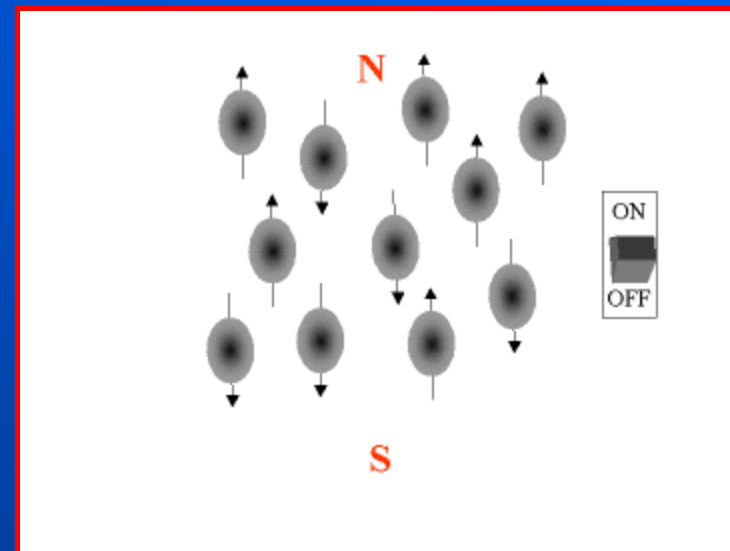
# SPINS ORIENTATION



The spins in normal conditions are randomly oriented.  
When we turn on a static magnetic field  $B_0$  the spins align with that external field in parallel or anti-parallel position.



Without  $B_0$



With  $B_0$

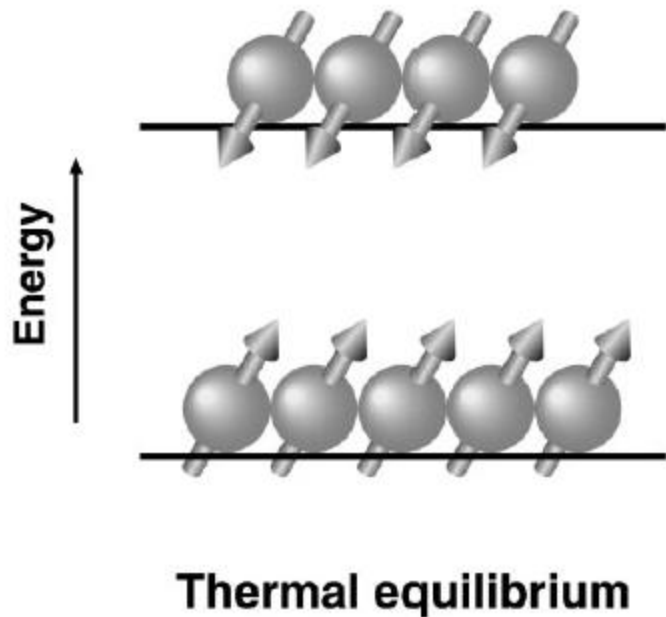
$B_0$





# Polarizzazione

In condizioni di equilibrio termico c'è un lieve vantaggio energetico in direzione up=>down quindi  $N^-$  saranno lievemente minori di  $N^+$



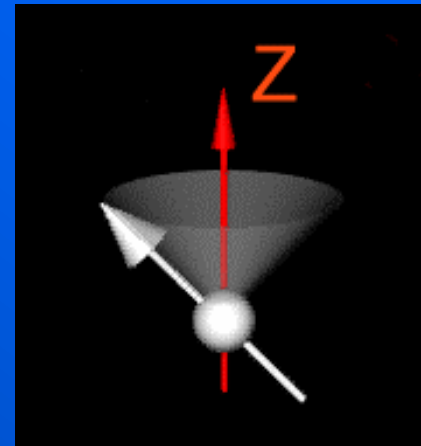
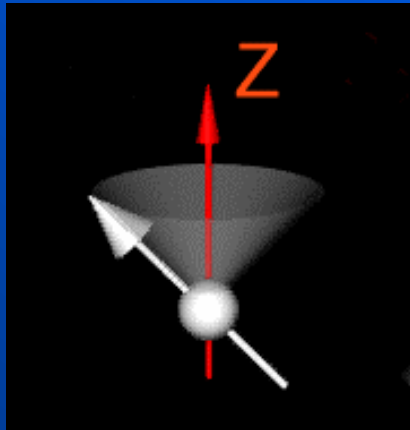
$$P = \tanh\left(\frac{\gamma \hbar B_0}{2k_B T}\right)$$

- $\gamma$  costante giromagnetica dell'atomo
- $\hbar$  costante di Planck
- $B_0$  campo magnetico
- $k_B$  costante di Boltzmann
- $T$  temperatura

a 1.5 T 7/1000000 per  $^1\text{H}$  (SNR di una singola ripetizione 0.0007%)



# MRI: PRECESSION FREQUENCY



$\uparrow B_0$

$\uparrow B_0$

LARMOR FREQUENCY:

$$\omega = \gamma B_0$$

Frequency of precession

Static magnetic field

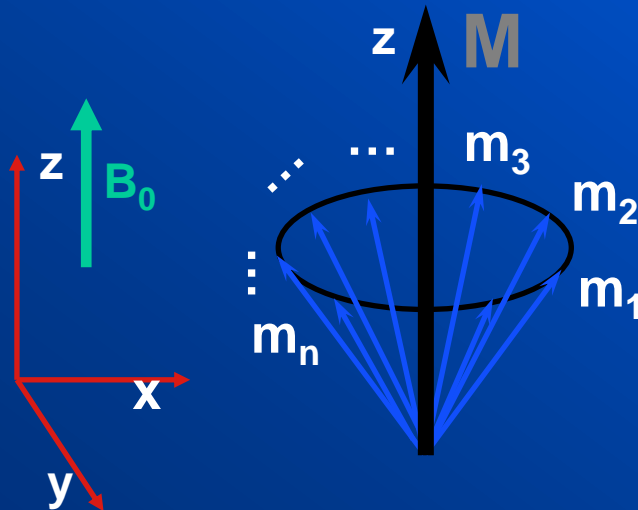
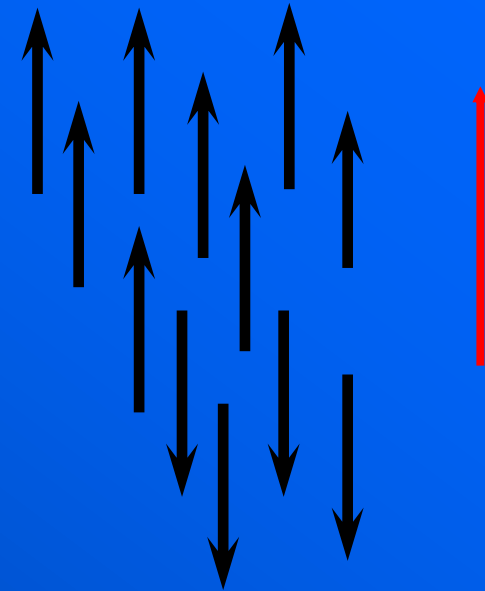
Gyromagnetic ratio

$\omega = 42.58 \times 1.5 = 63.87$  MHz:  
we are in the radiofrequency domain (RF)

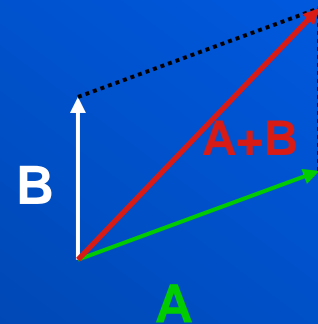
1 Tesla is = 10.000 Gauss.  
(1 Gauss = intensity of the earth magnetic field)



It is given by the vectorial summation of the single magnetic moments of the spins.



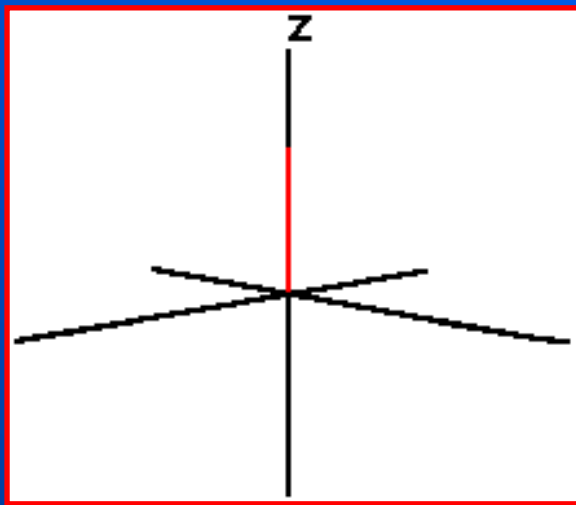
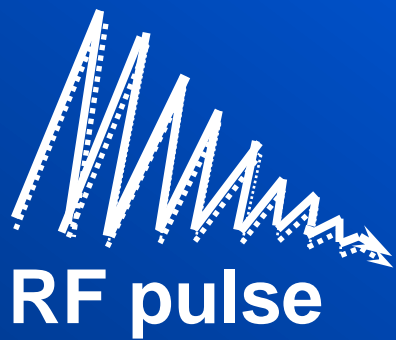
Sum of two vectors  $A+B$ :





# MAGNETIC RESONANCE

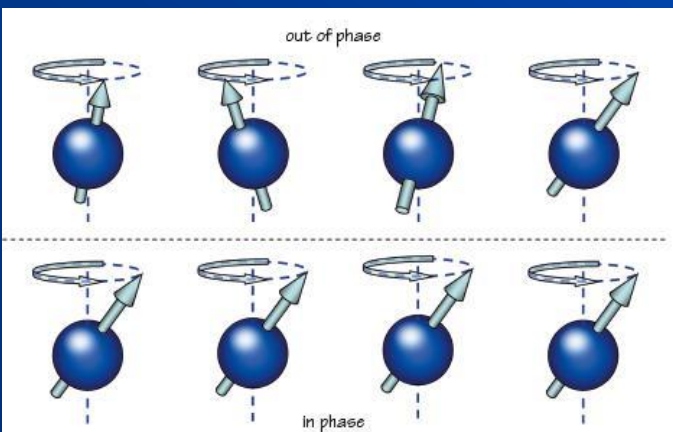
If the protons aligned in a static magnetic field  $B_0$  are excited by a RF pulse at the frequency of precession (Larmour Frequency) we have the phenomenon of nuclear magnetic resonance.



Loss of longitudinal magnetization

$B_0$

## Phase coherence

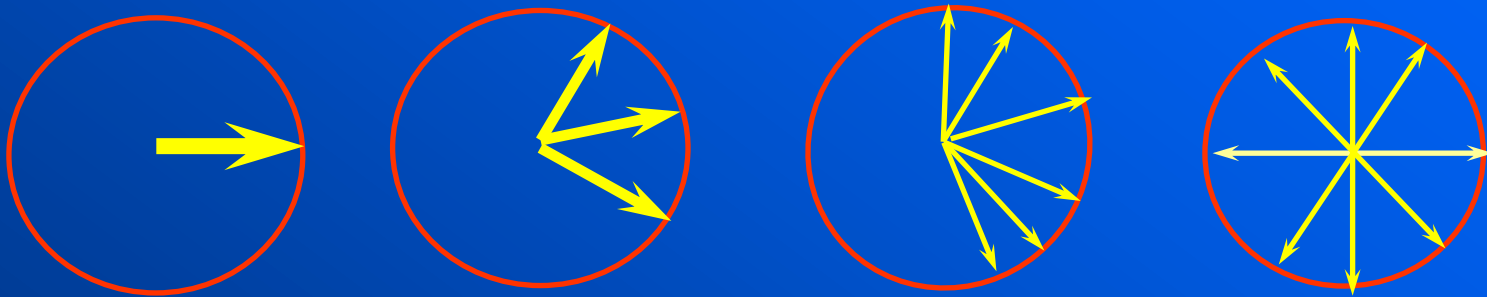




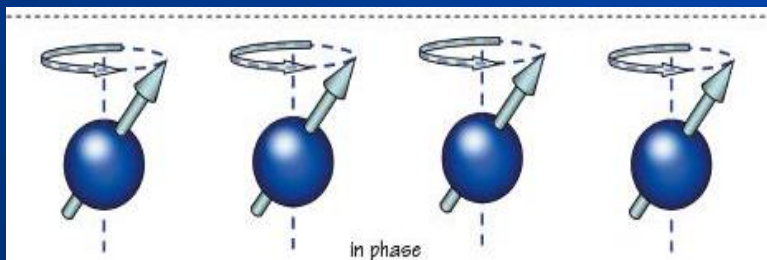


## $T_2$ = spin-spin relaxation

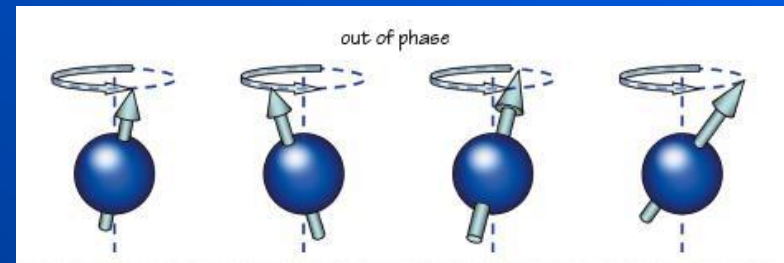
The spin-spin relaxation is caused by the interaction between nuclear magnetic moments.



These spin-spin interactions cause a change in the precessing frequencies of each nucleus. The result is a loss in phase coherence.



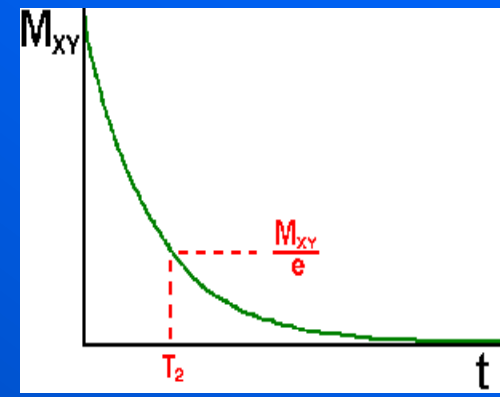
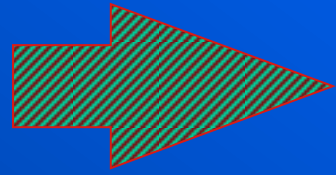
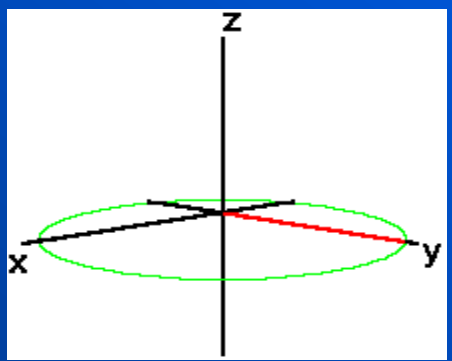
in phase



out of phase



## TRANSVERSE RELAXATION TIME T2

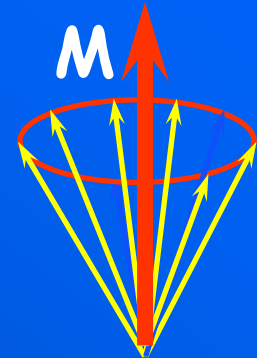
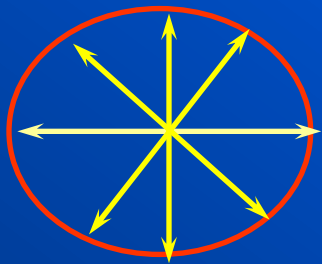


The constant of the transverse relaxation time is given by  $T_2$ , that is the time necessary to reduce the value of  $M_{xy}$  by 63%.



# LONGITUDINAL RELAXATION TIME T1

The spin-lattice relaxation is caused by the exchange of energy between spins and the surrounding environment.

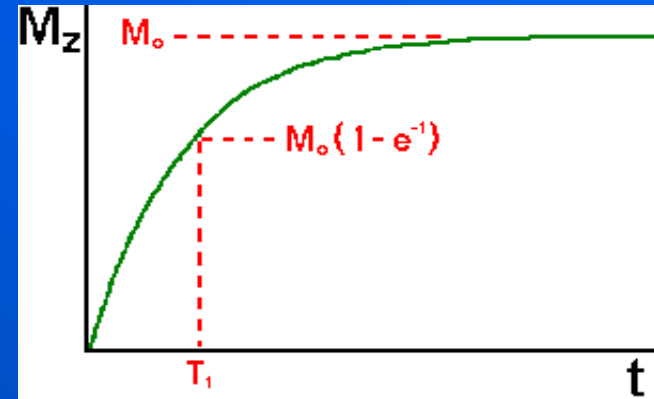
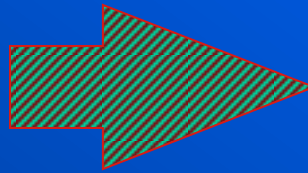
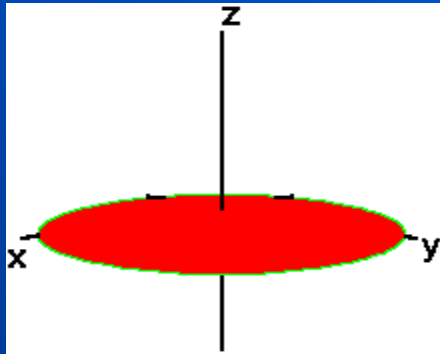


The magnetic moments gradually realign with  $B_0$





# LONGITUDINAL RELAXATION TIME T1



$T_1$  is the time needed for 63% of  $M_z$  to return to equilibrium  $M_0$  after a  $90^\circ$  RF pulse.

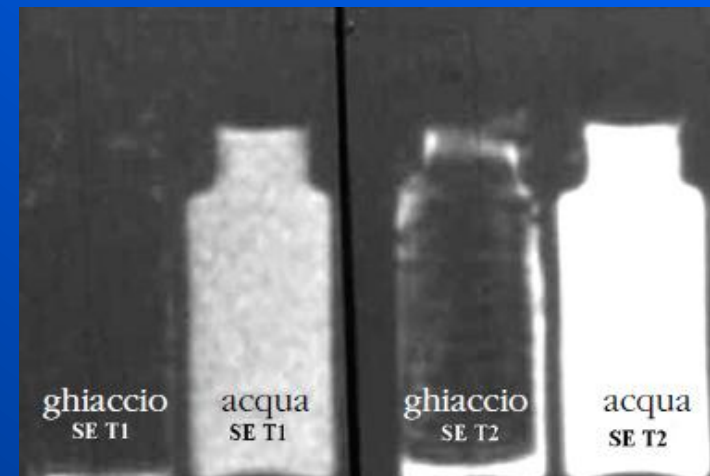


# MRI - parameters

Longitudinal relaxation time, or spin-lattice relaxation time:  $T_1$

Transverse relaxation time, or spin-spin relaxation time:  $T_2$  ( $T_2^*$ )

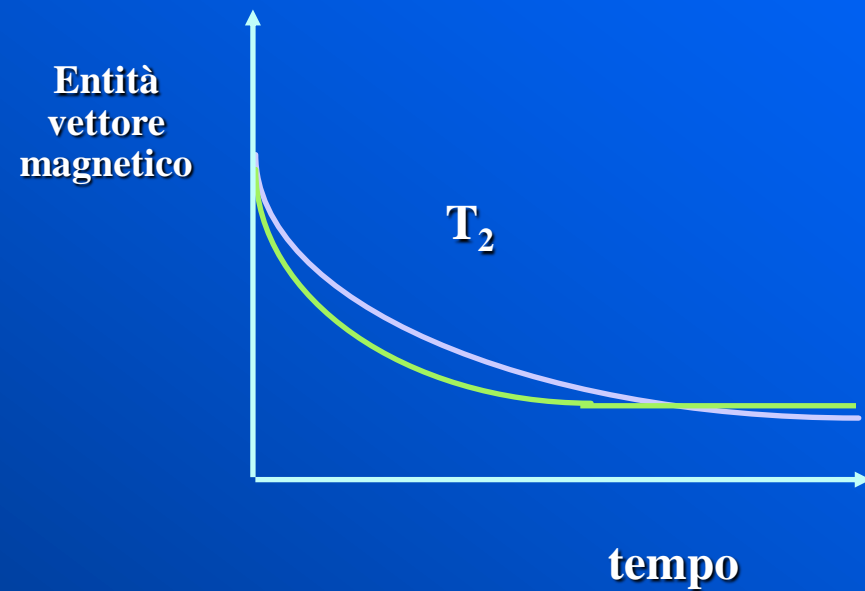
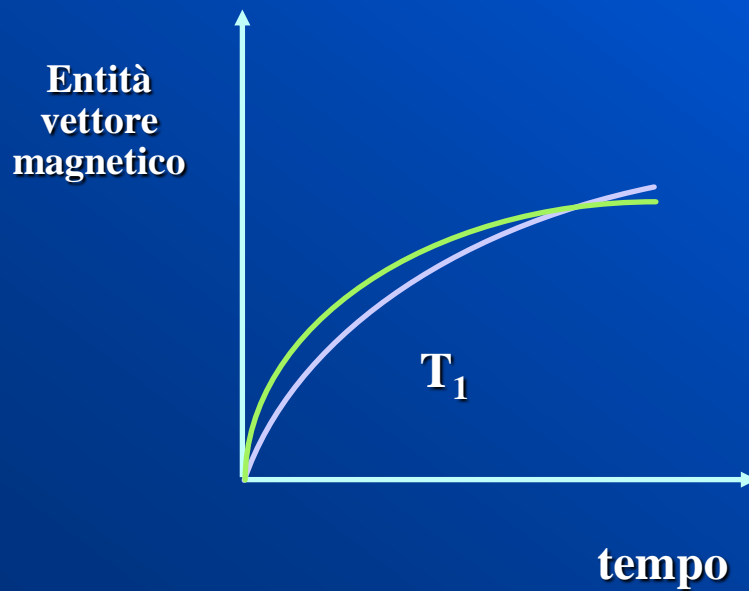
Proton density: PD





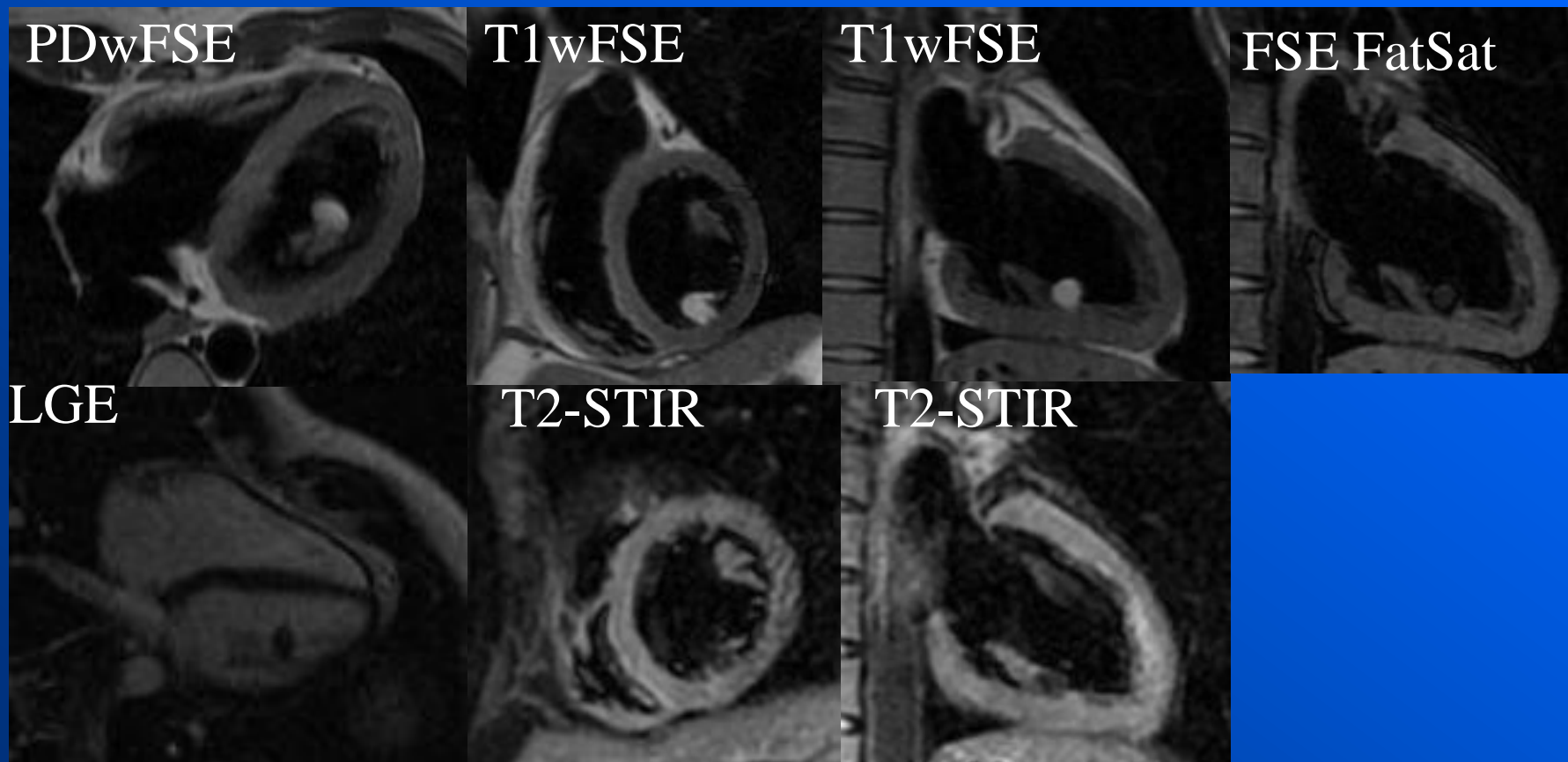
# Longitudinal and transversal Relaxation

— Tessuto A  
— Tessuto B





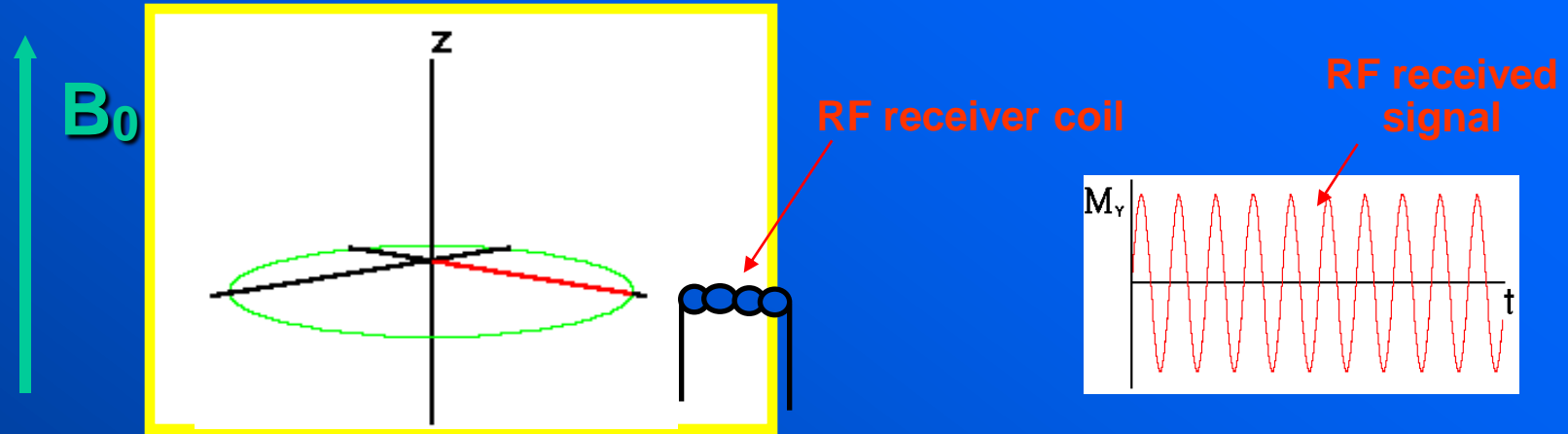
# WEIGHTED IMAGES



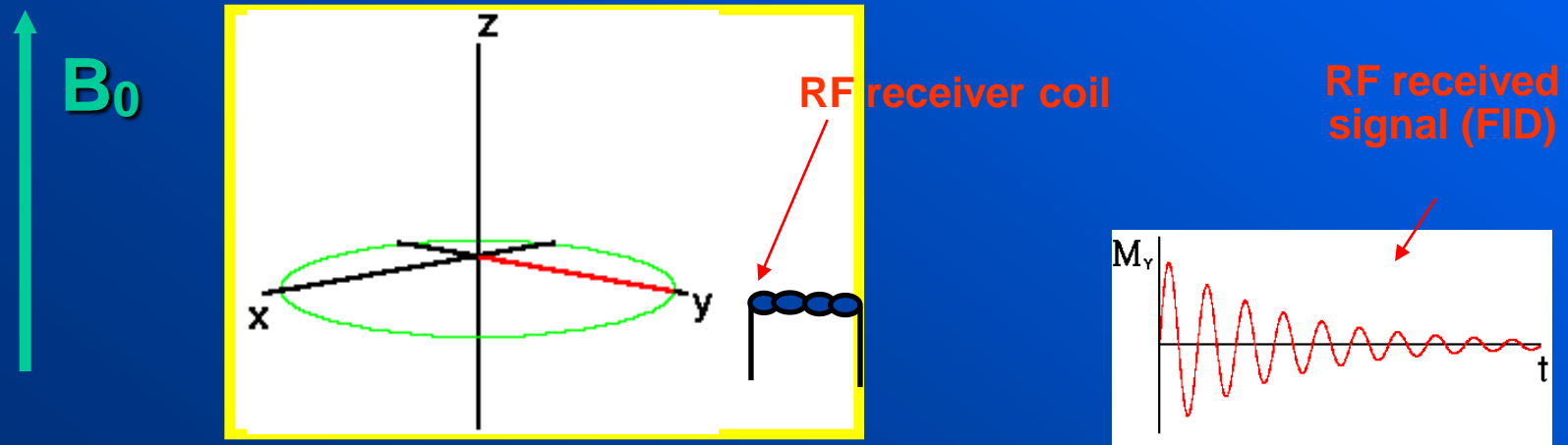
# THE SIGNAL



In a perfect world....



.....in the real one FID Signal: Free Induction Decay







# PARAMETERS OF ACQUISITION:

 FA - Flip Angle: the angle between  $B_0$  e  $M$ ; it is proportional with the duration of the RF pulse.

 TR - Time of Repetition: the time between an RF pulse and the next.

 TE - Time of Echo: the time between emission of the RF pulse and reception of the signal.

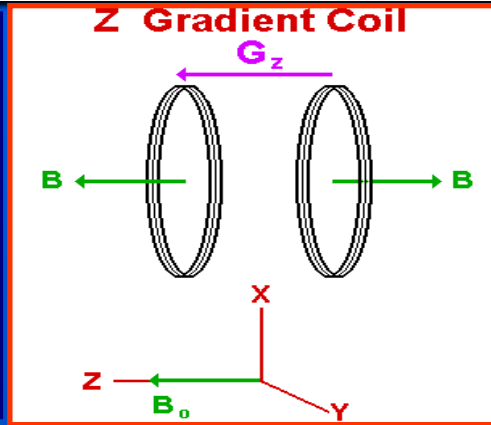
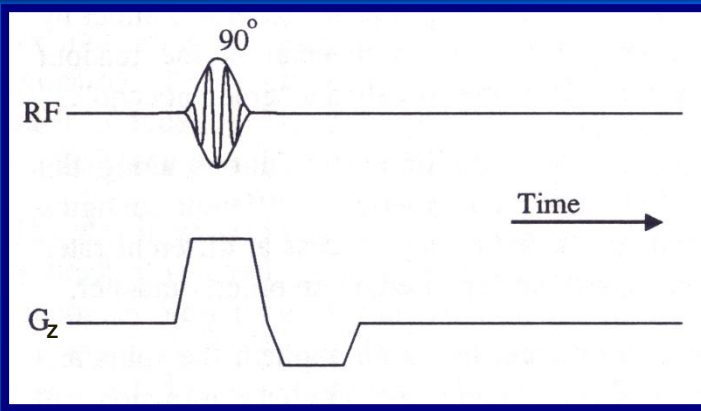
T1w = short TR (<800 ms) - short TE (<10 msec)

T2w = long TR (>1800) - long TE (>60 msec)

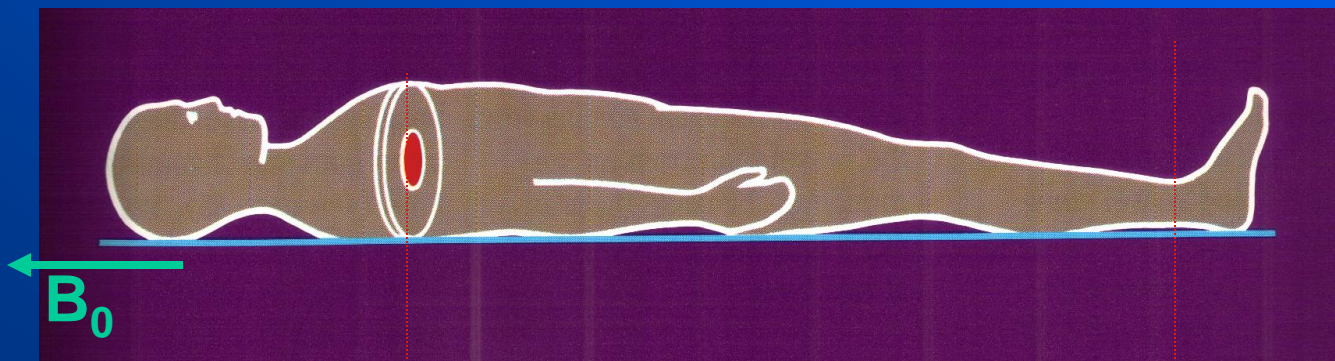
PDw = long TR - short TE



# GRADIENT FOR THE SLICE SELECTION



Gradient  
along  $B_0 - z$

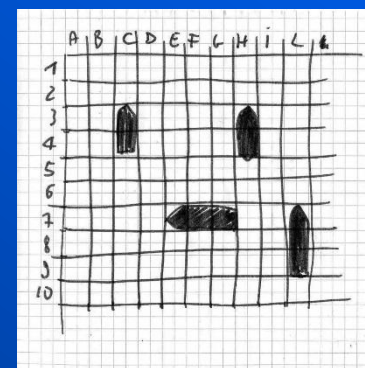
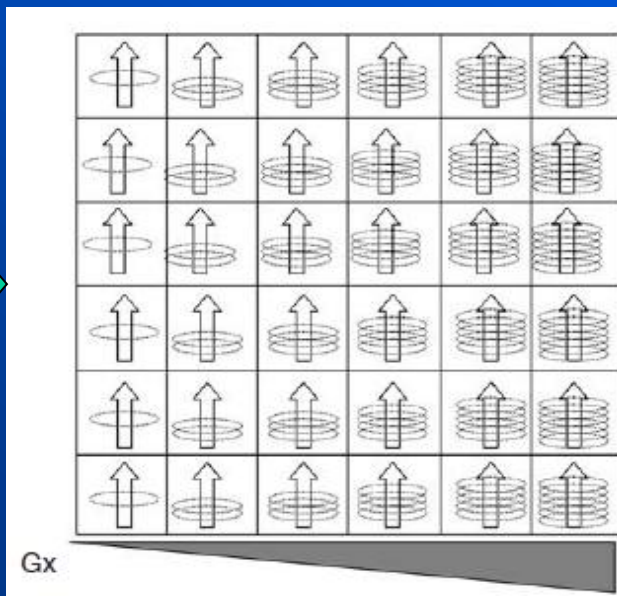
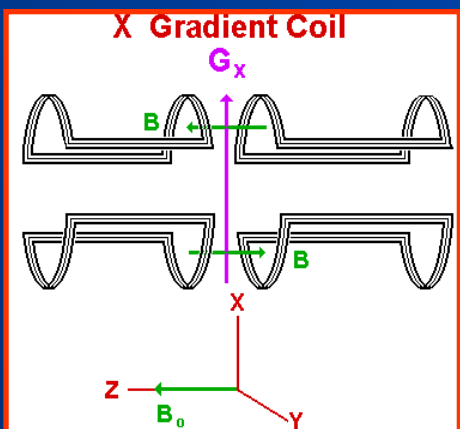
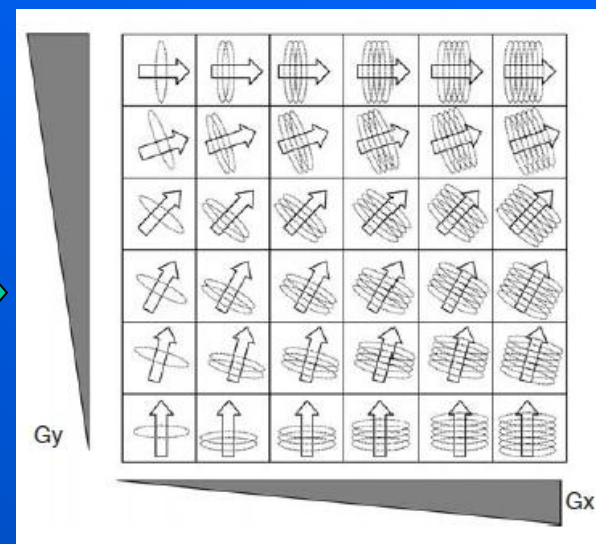
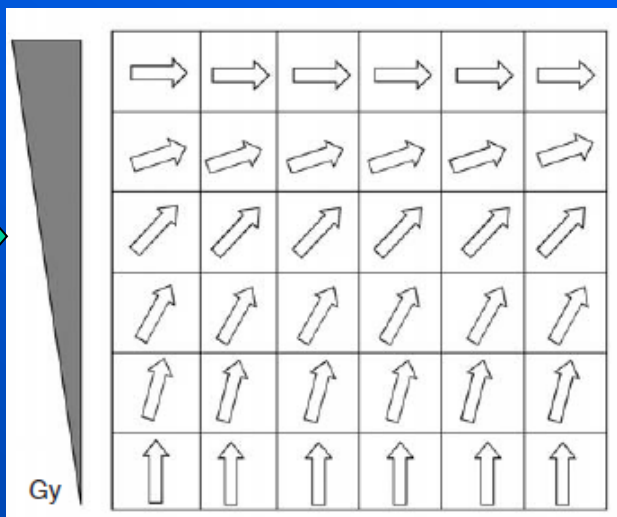
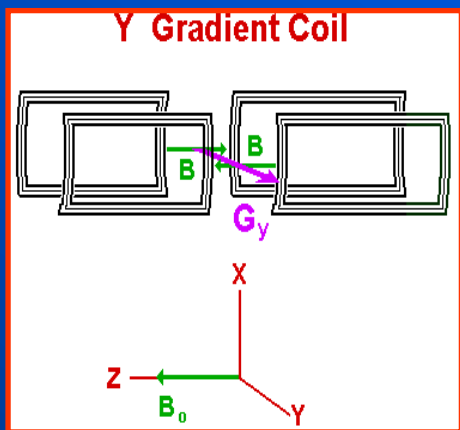


$$\omega = \gamma B$$





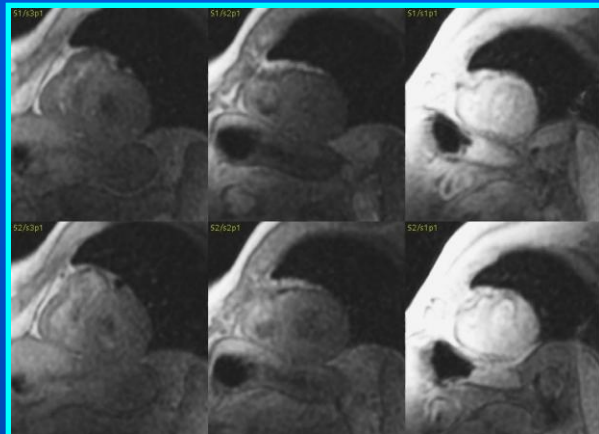
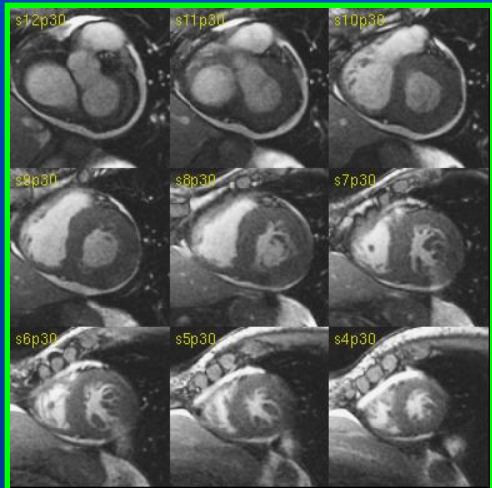
# THE GRADIENT COIL





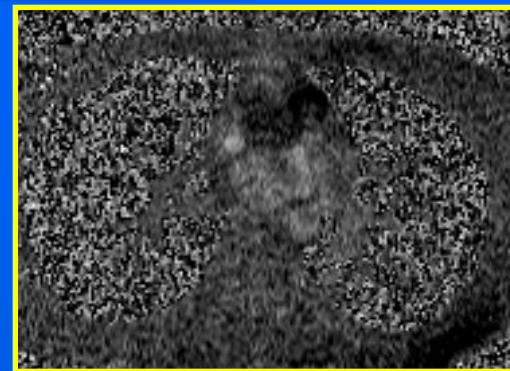
# CMR PULSE SEQUENCES

**SSFP**

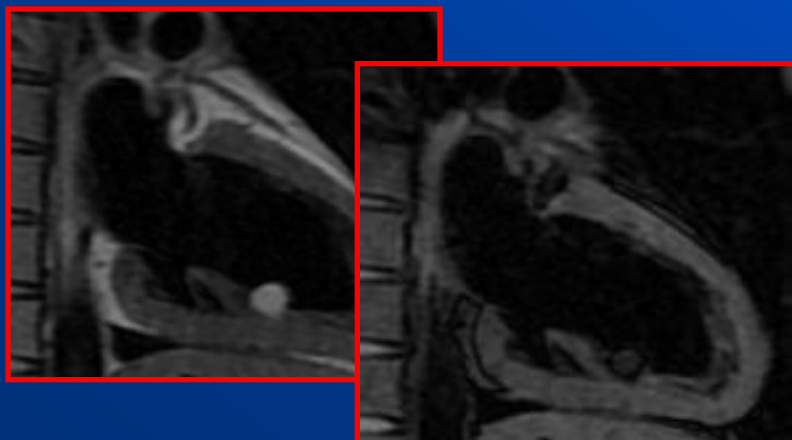


**EPI ET**

**PC**

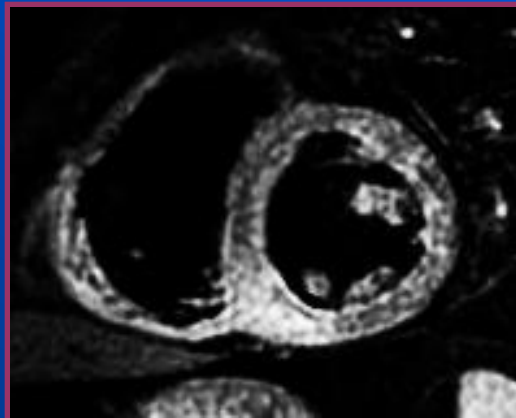


**FSE or Double IR**



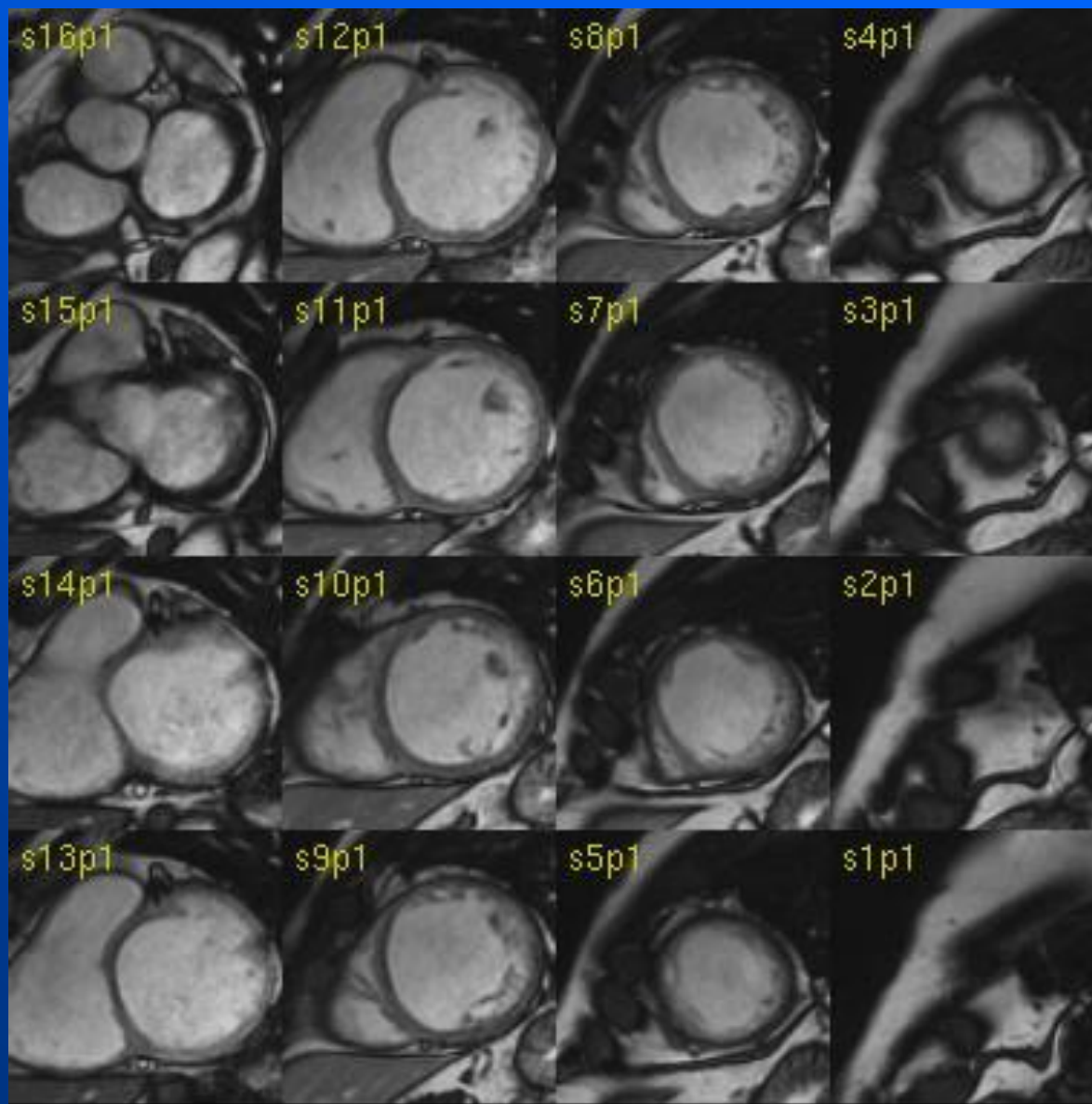
**FSE or Double IR Fat sat**

**STIR**



**IR GRE T1 post m.d.c  
Delayed Enhancement**

# Cine-CMR: SSFP & GRE



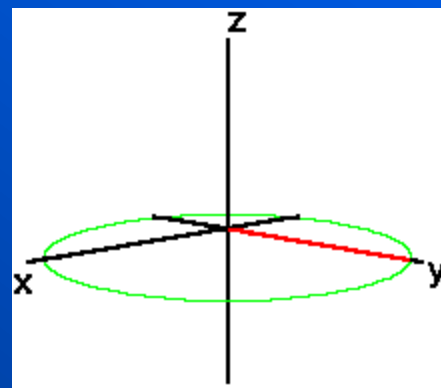
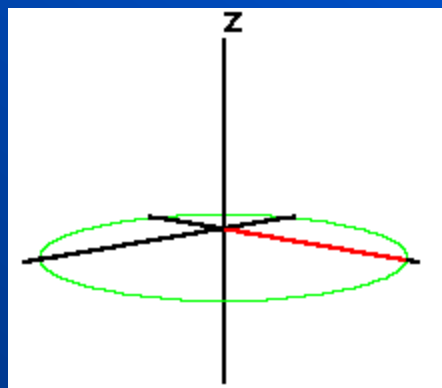
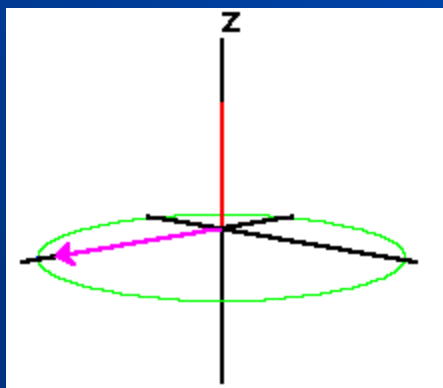


# GRADIENT-ECHO

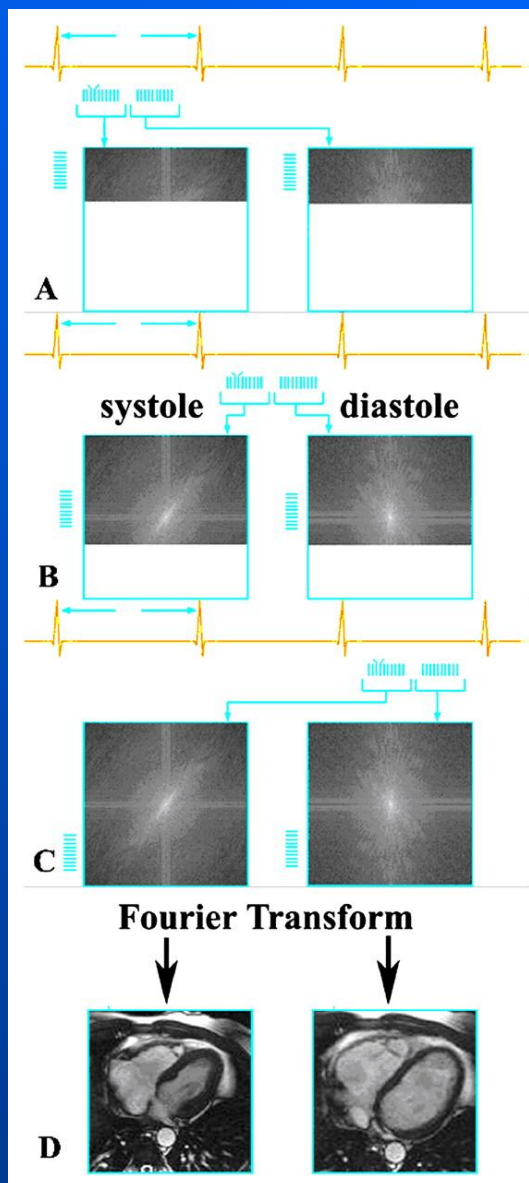
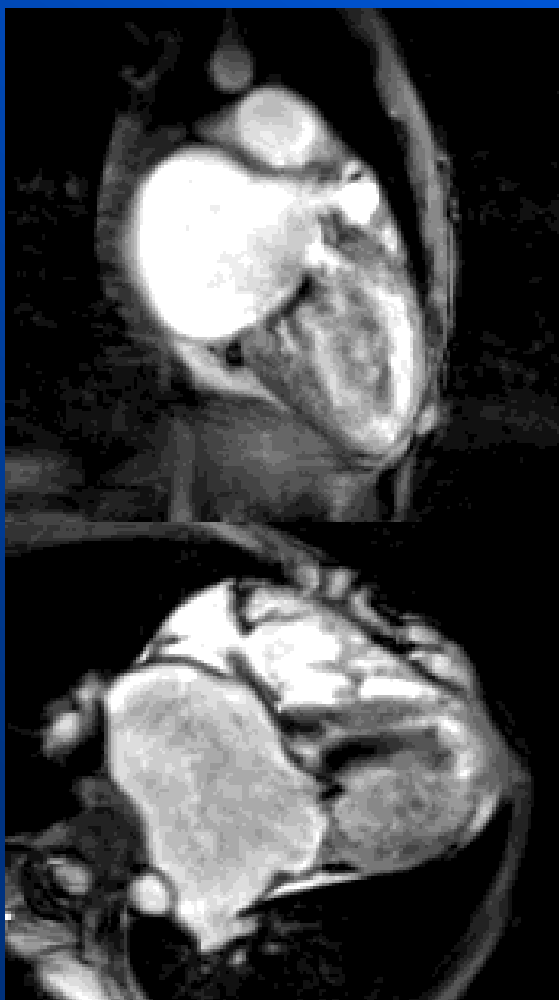
← TR →

**RF pulse**  
( $FA \leq 90^\circ$ )

Signal acquisition FID

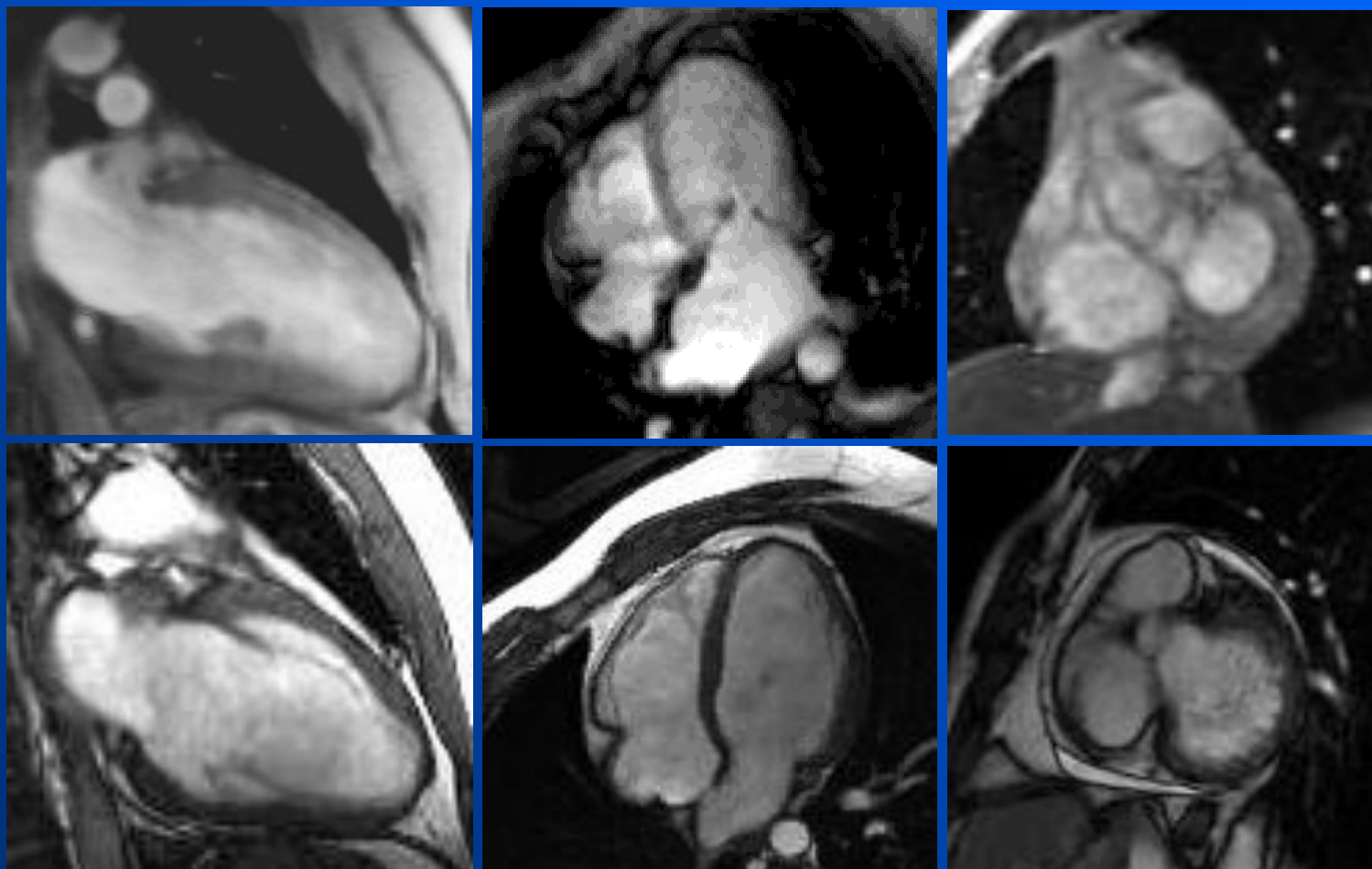


# Cine - GRE (SPGR)





# CINE-CMR



**GRE**

1994



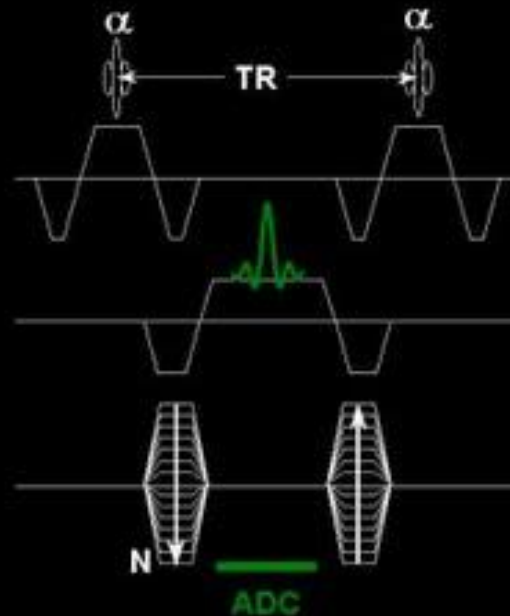
2002

**SSFP**





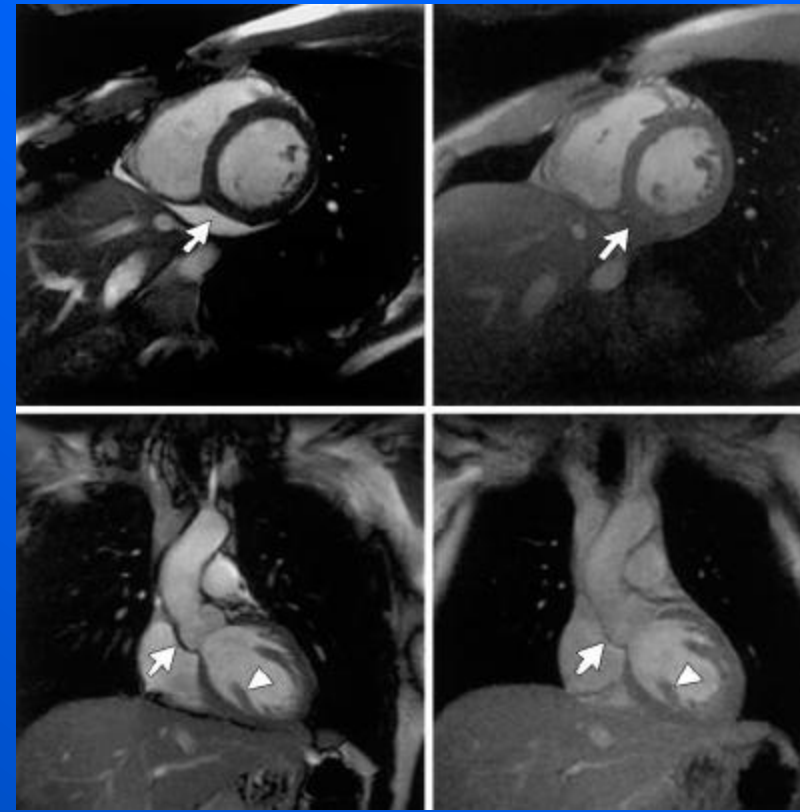
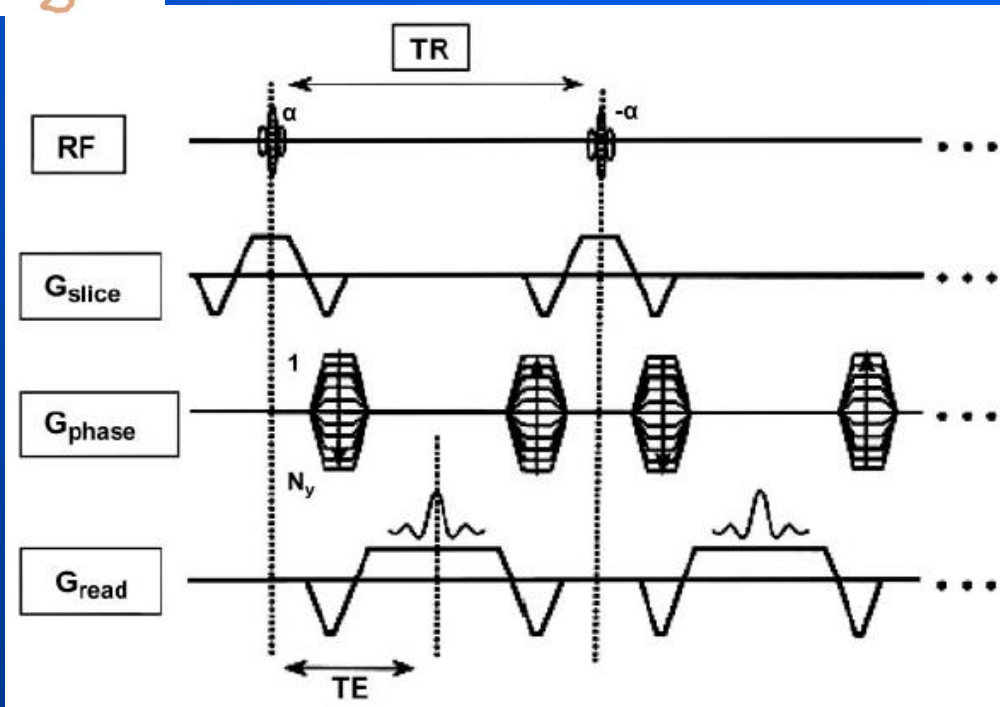
## Steady State Free Precession (TrueFisp)



Steady State Free Precession (SSFP) with Balanced Gradients  
Image contrast depends on T2/T1 relaxation rates of tissues  
and the steady state signal of spins

Steady state: short TE and  $TR < 10 \text{ msec} \rightarrow T2/T1w$

# SSFP

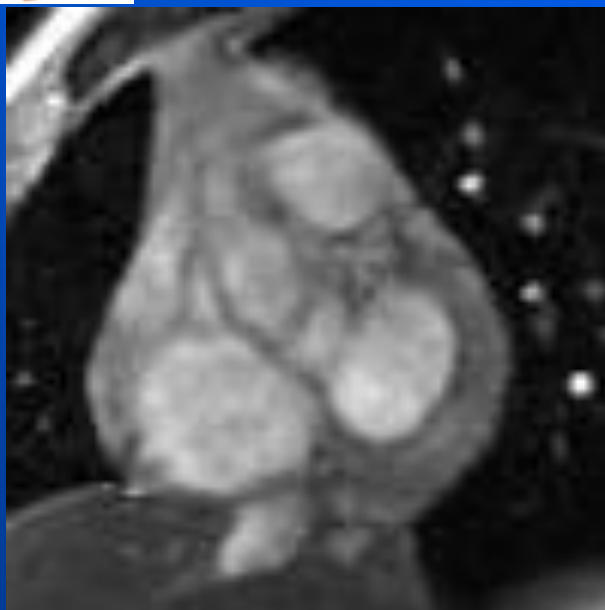


Gradients ( $G$ ) are fully balanced along all three (section-selective, phase-encoding, and readout) axes.

The sum of positive gradient areas is exactly balanced by the sum of negative gradient areas. In this case, echo and readout occur midway between RF pulse.



# MRI: cine-pulse sequences



**GRE**

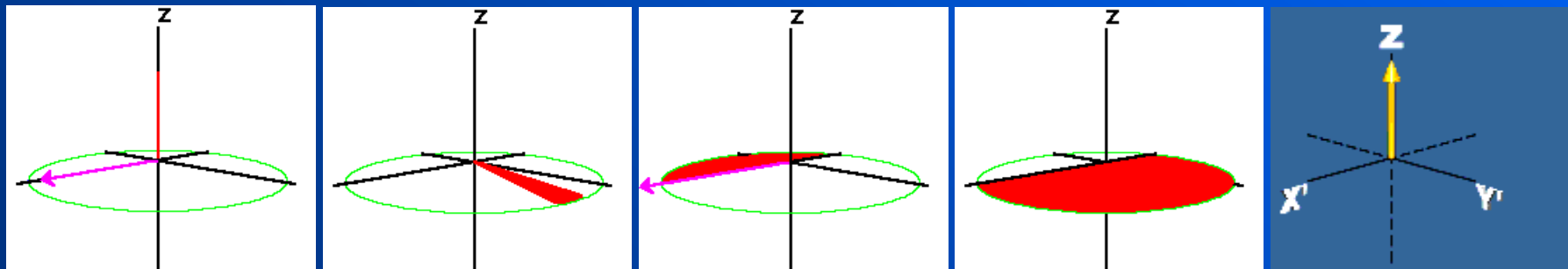
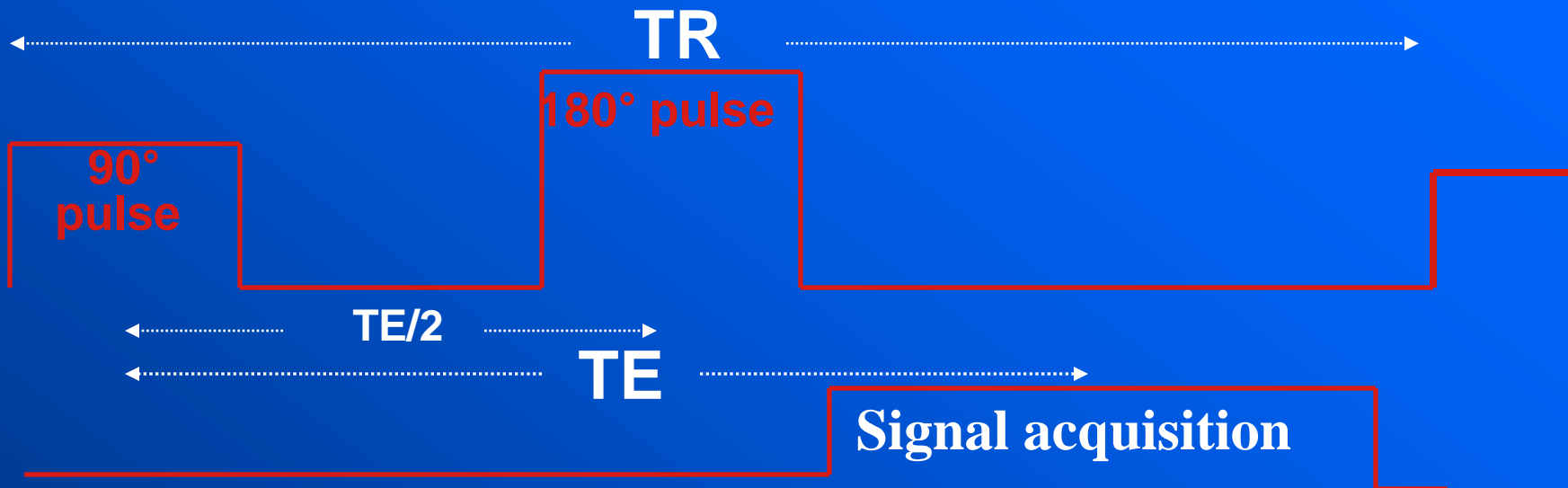


**SSFP**

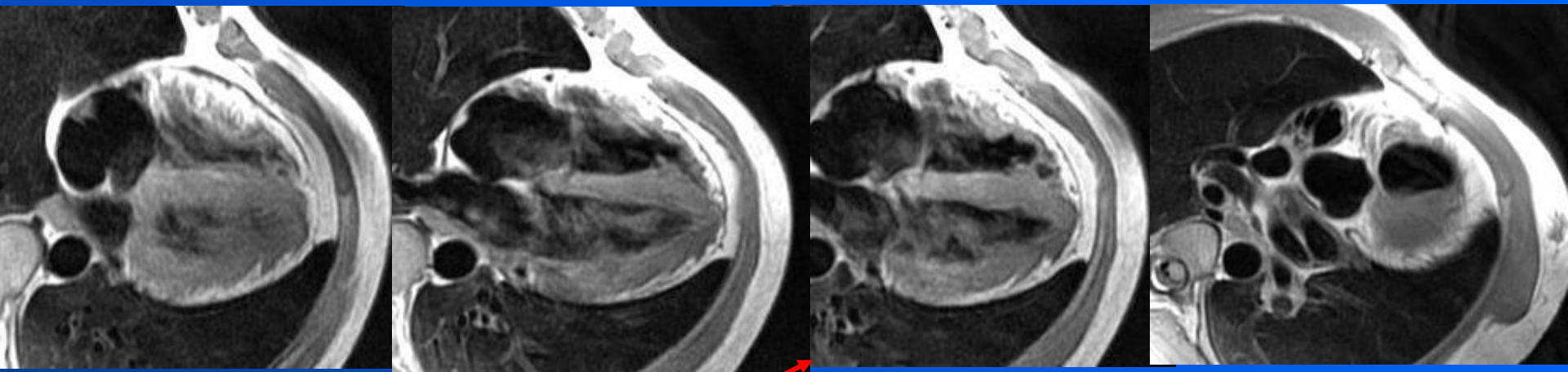
	<b>SSFP</b>
<b>TE</b>	<b>Minimum</b>
<b>TR</b>	<b>HR depending</b>
<b>FLIP ANGLE</b>	<b>45° to 60°</b>
<b>RBw</b>	<b>125 kHz</b>
<b>FOV</b>	<b>20-40</b>
<b>THICKNESS</b>	<b>5-8 mm</b>
<b>SPACING</b>	<b>0.0</b>
<b>NEX</b>	<b>1+ (BH or FB)</b>
<b>N° Phase</b>	<b>30</b>
<b>View per segm.</b>	<b>8-16 to 30-32</b>
<b>R-R</b>	<b>1</b>



# SPIN-ECHO

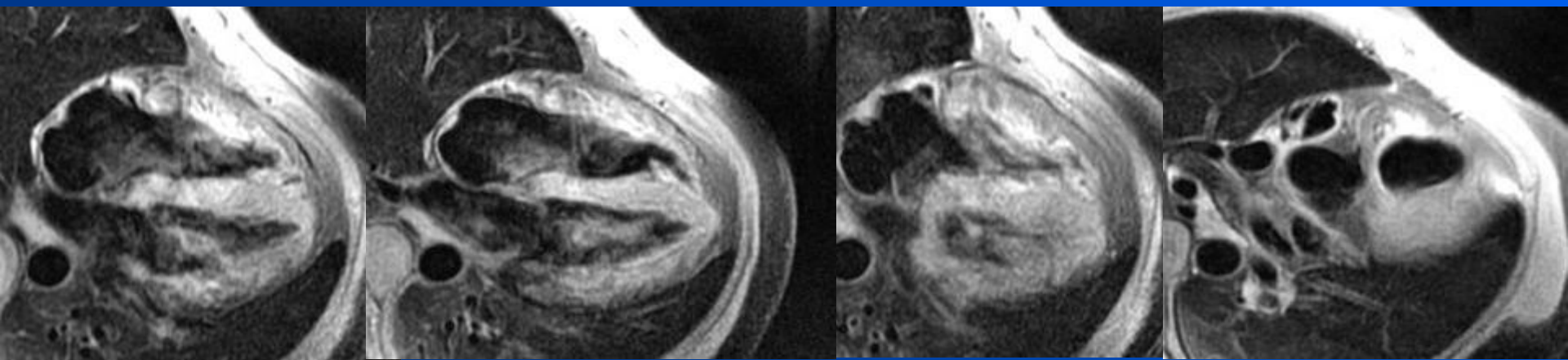


# FSE BLACK BLOOD



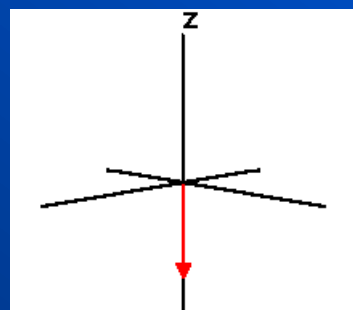
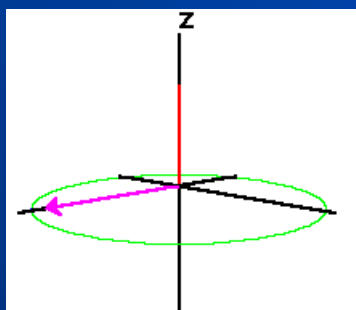
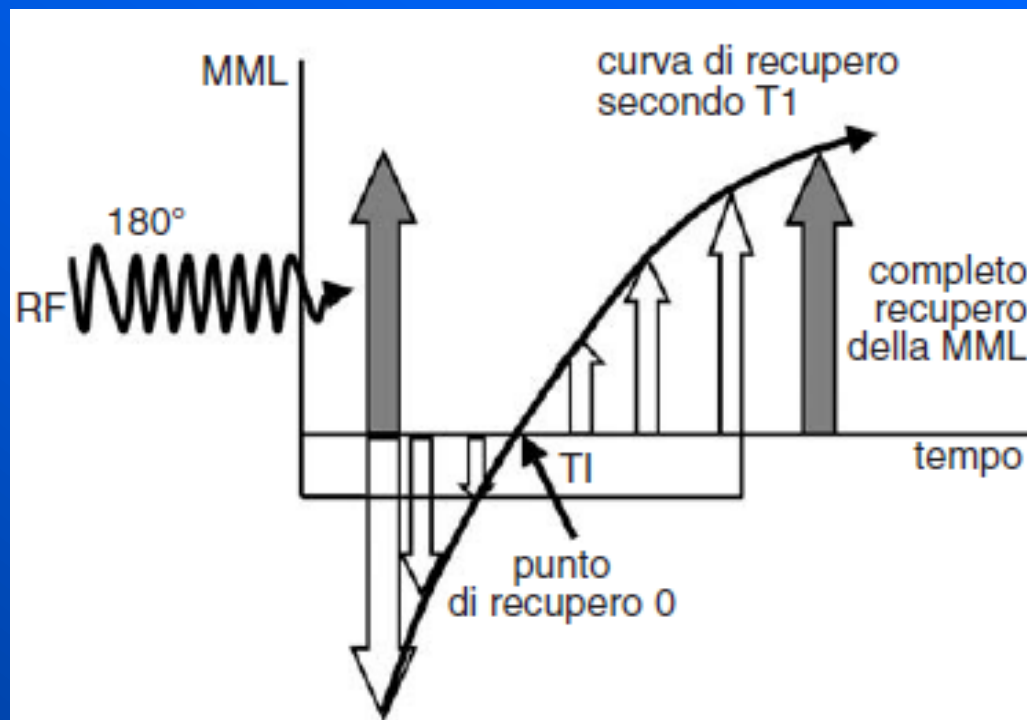
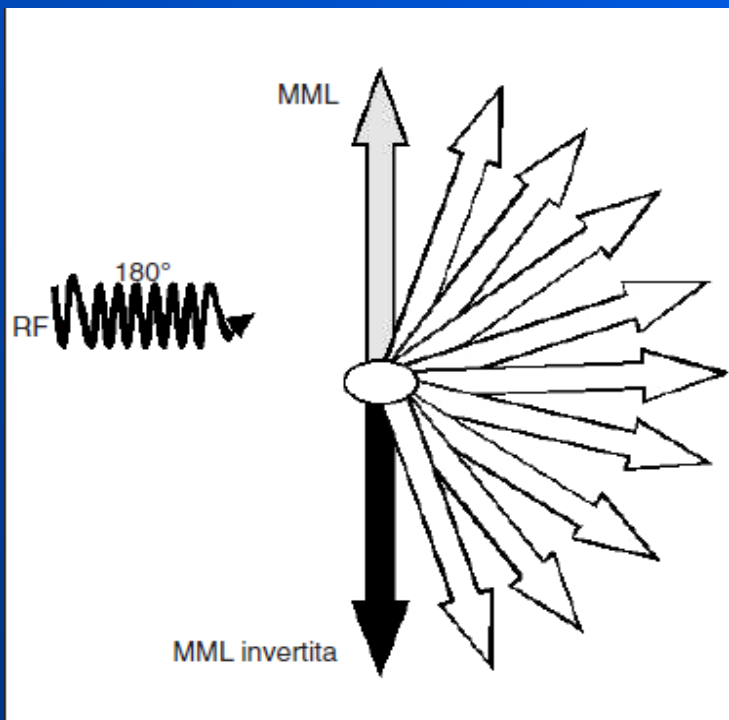
T1 weighted SpinEcho

T1 weigthed Spinecho with fat saturation





# INVERSION RECOVERY: 180° RF

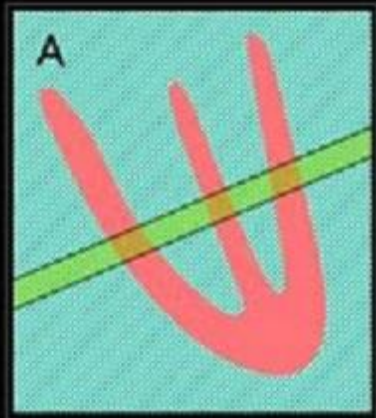


# DOUBLE-IR



## DOUBLE IR

A non-selective  $180^\circ$  inversion pulse excites all the tissues and blood within the entire heart when the R-wave trigger is detected at the beginning of the cardiac cycle



R-wave Trigger

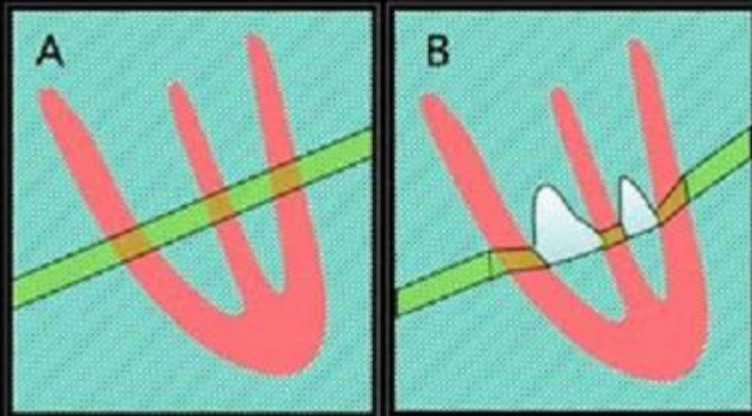


non-selective  
inversion

# DOUBLE-IR



## Dark Blood Pulse



R-wave Trigger

Systole



non-selective  
inversion



slice-selective  
reversion

## DOUBLE IR

Immediately following, a slice-selective re-inversion pulse excites only the tissues and blood within the image slice.

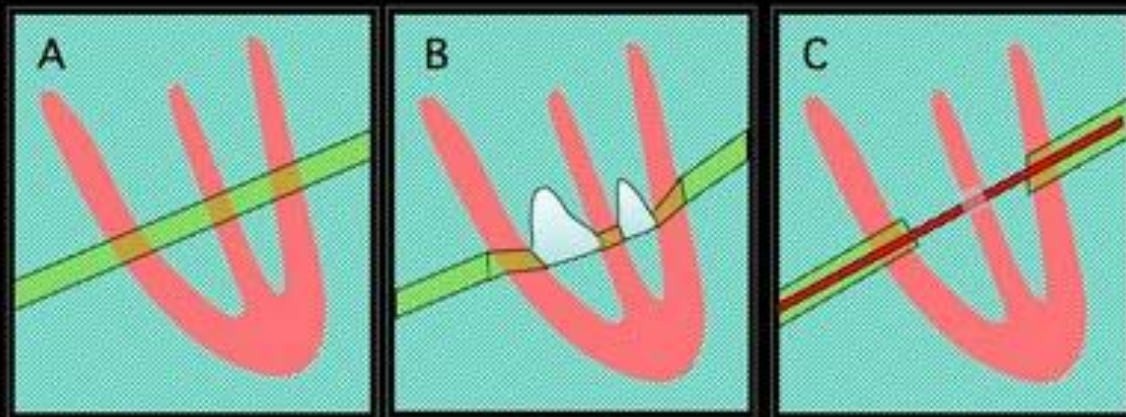
The net result is that everything within the slice is flipped back to normal because it experienced both the inversion and re-inversion pulses, whereas everything outside the slice remains inverted.



# DOUBLE-IR



## Dark Blood Pulse



R-wave Trigger

Systole

Late diastole

non-selective  
inversion

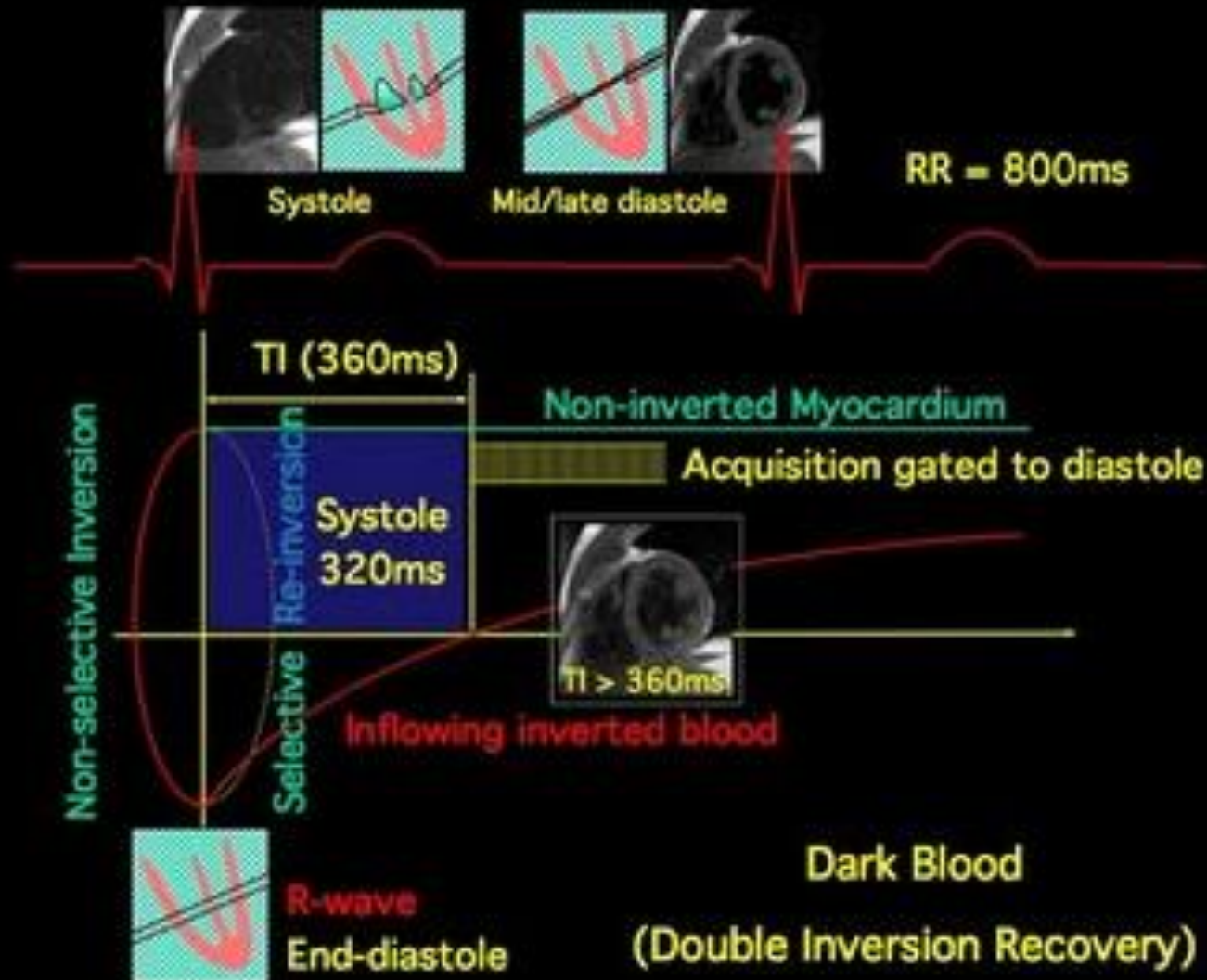
slice-selective  
reversion

imaged slice

systolic contraction forces  
the blood within the slice to  
be replaced by blood from  
outside the slice.

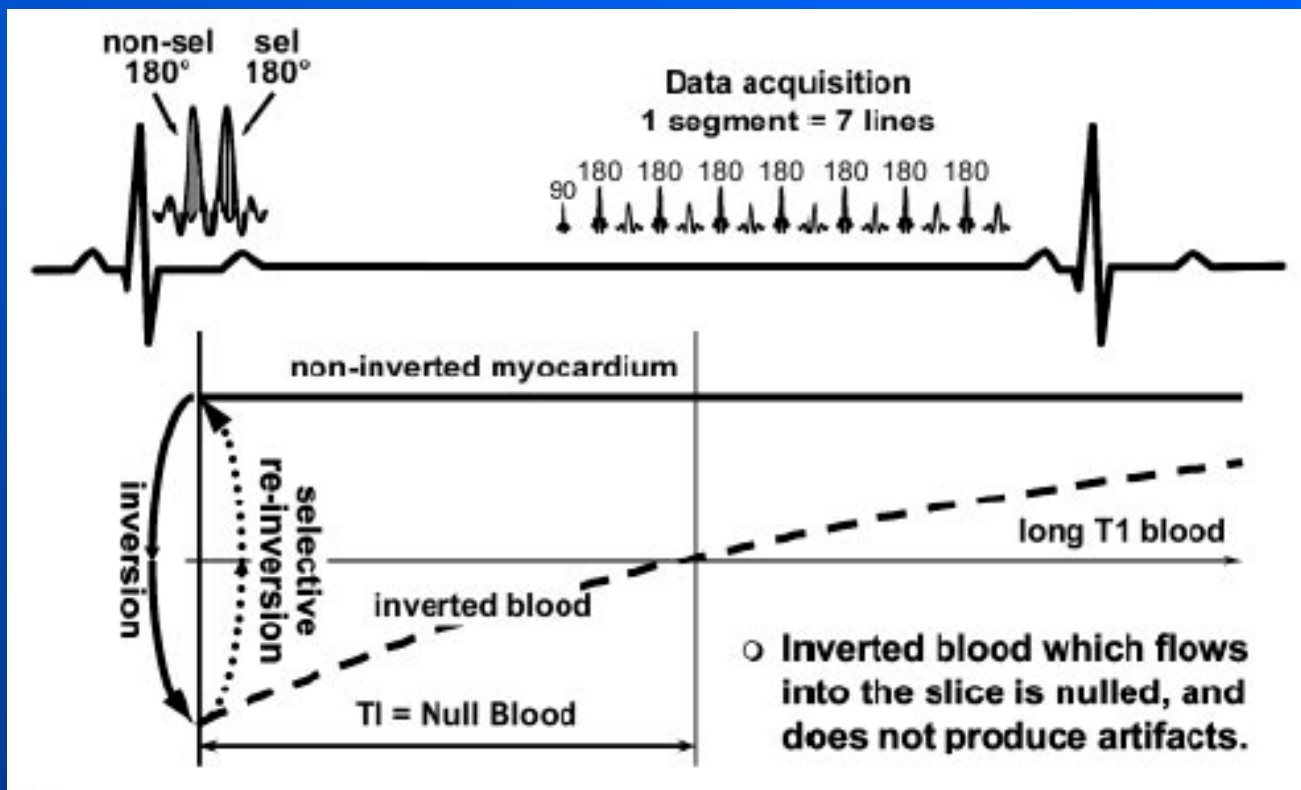
After enough time delay has occurred to allow the blood to be fully replaced within the slice, the image data is collected during mid to late diastole of the cardiac cycle.

# DOUBLE-IR





# DOUBLE-IR

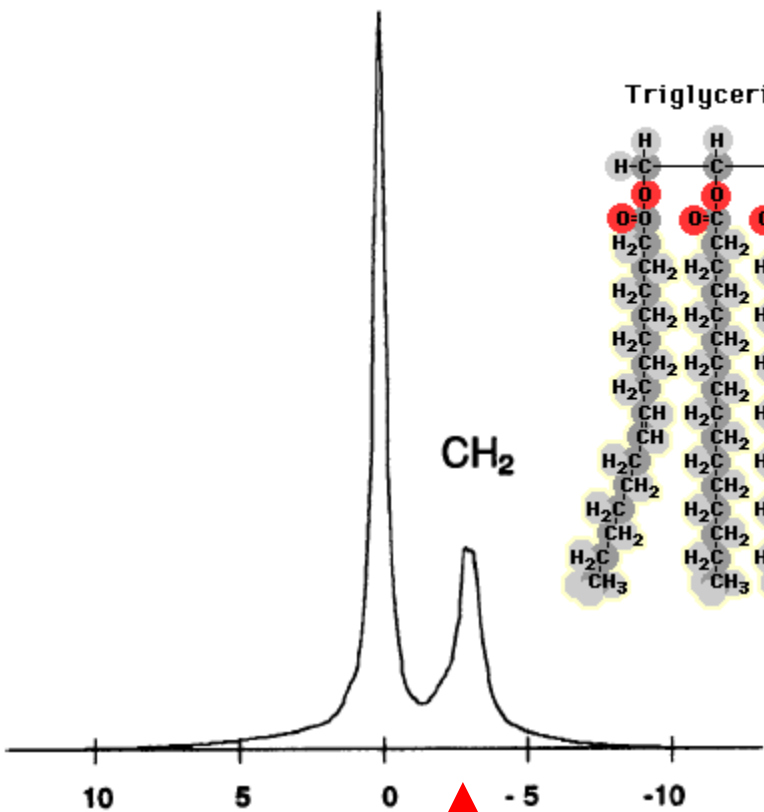


the readout module  
 (a 90° pulse followed by a train of refocusing 180° pulses)  
 is applied when blood is relaxing to zero, in-  
 flowing blood produces no signal.

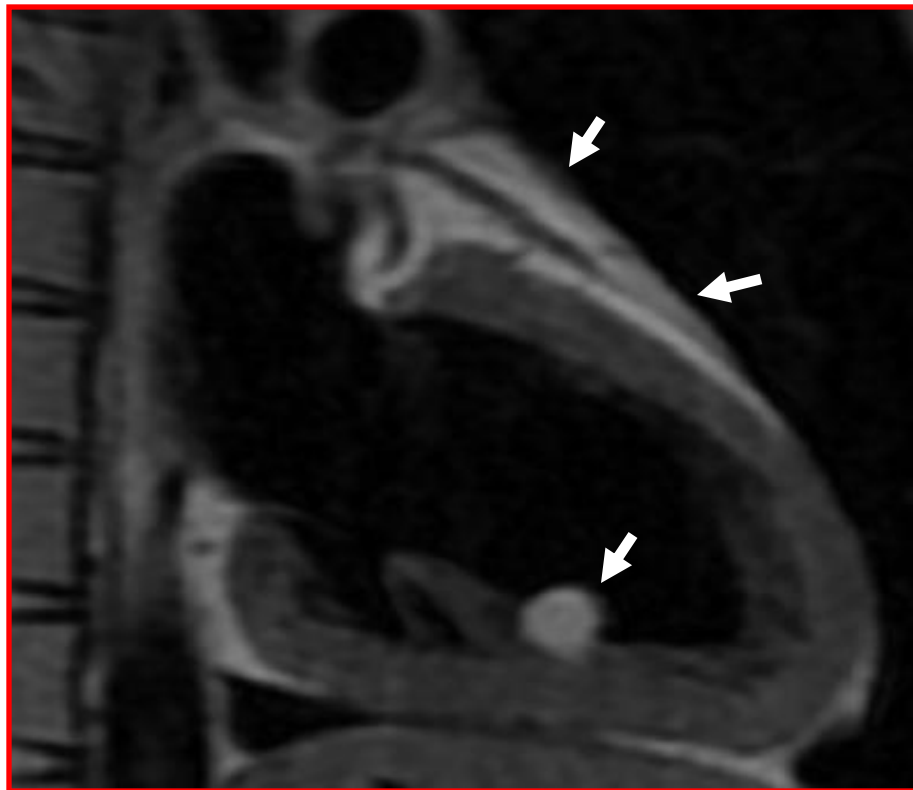
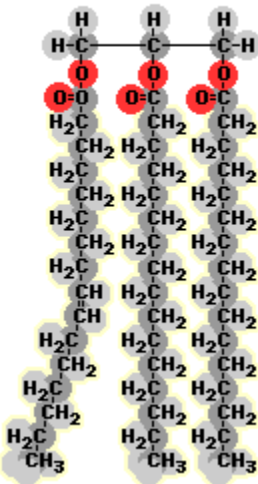
# FSE and Double IR



H<sub>2</sub>O



Triglyceride



←  
δ [ppm]

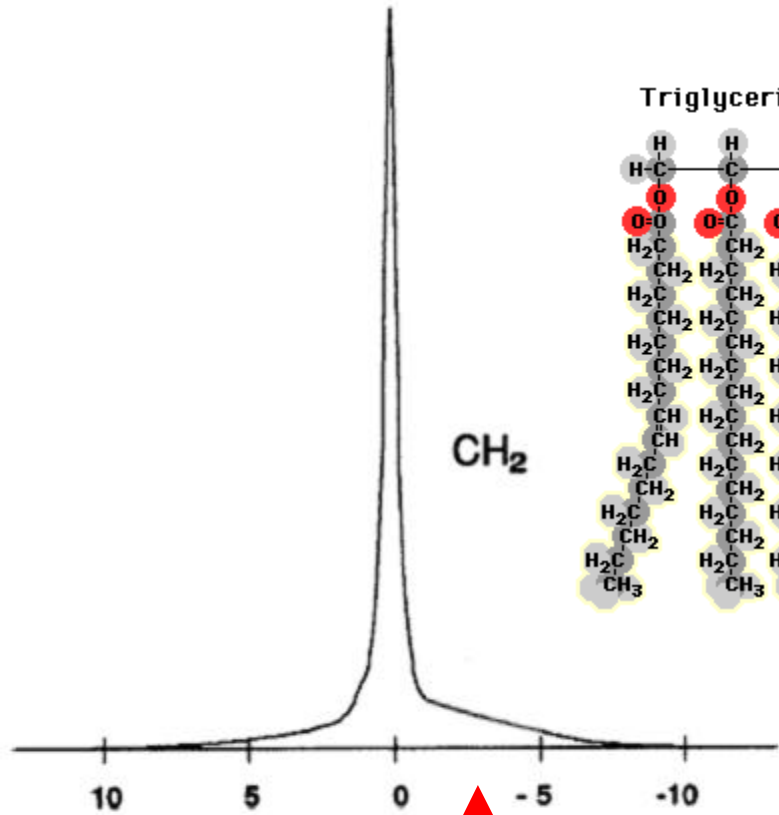


Chemical-Shift

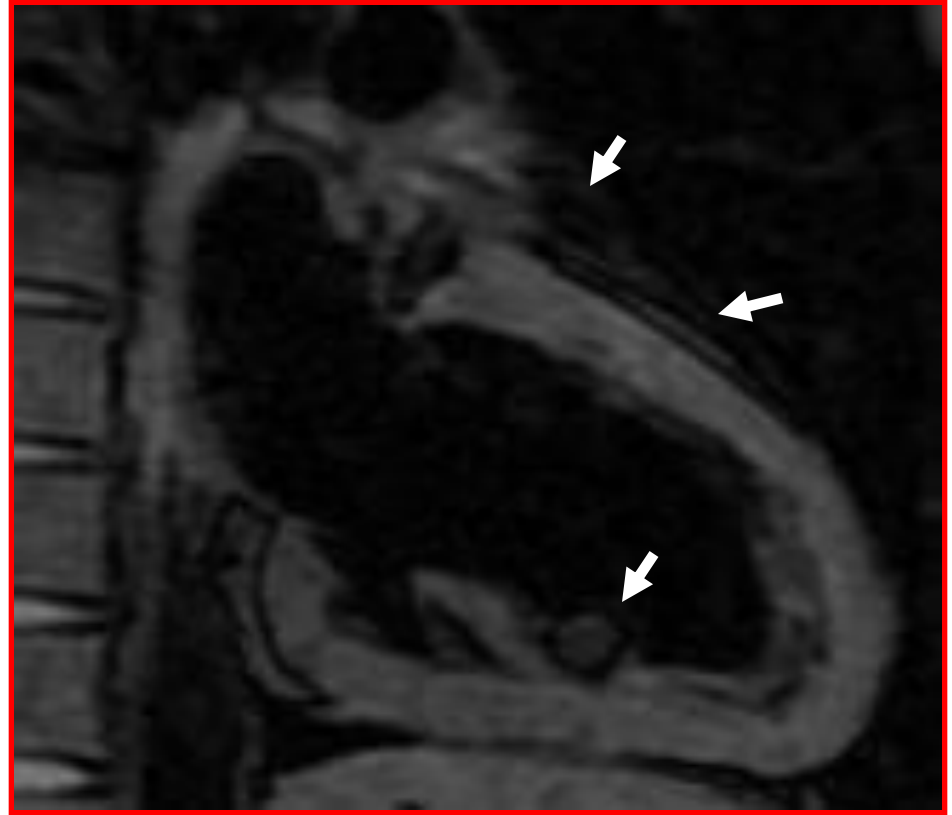
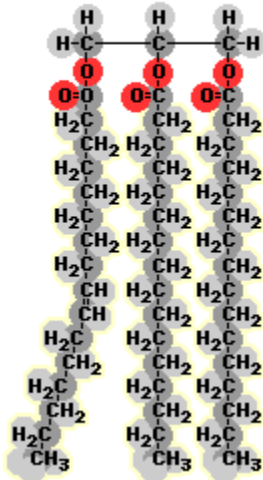
# FSE and Double IR



H<sub>2</sub>O

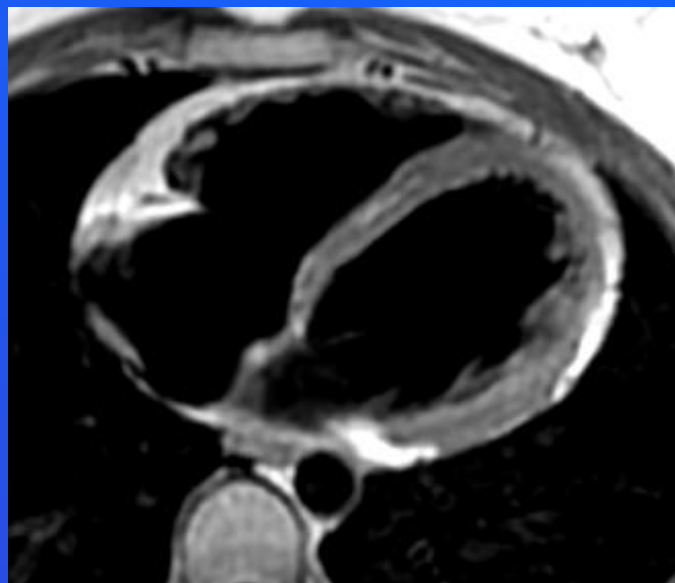
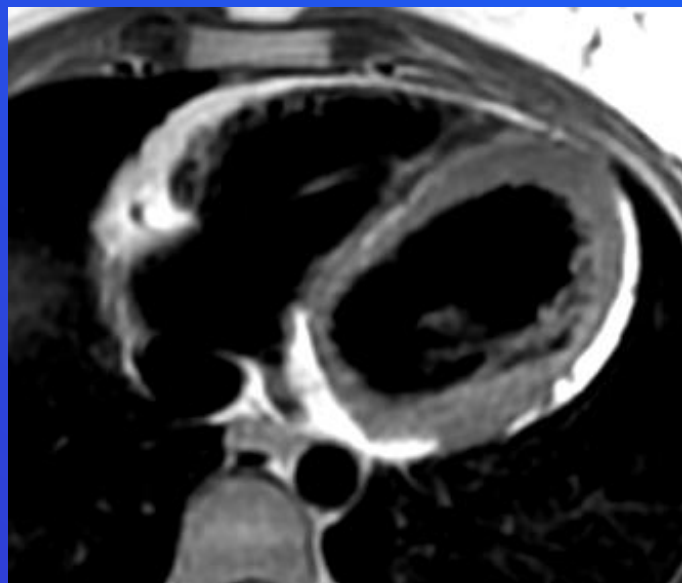


Triglyceride

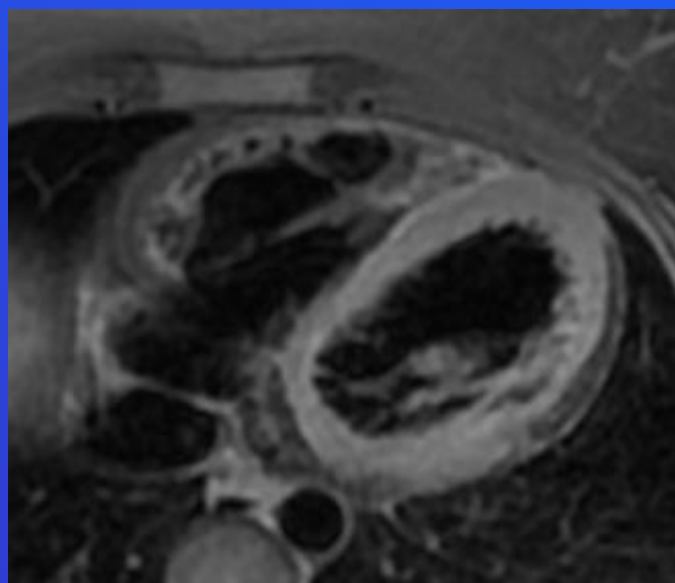
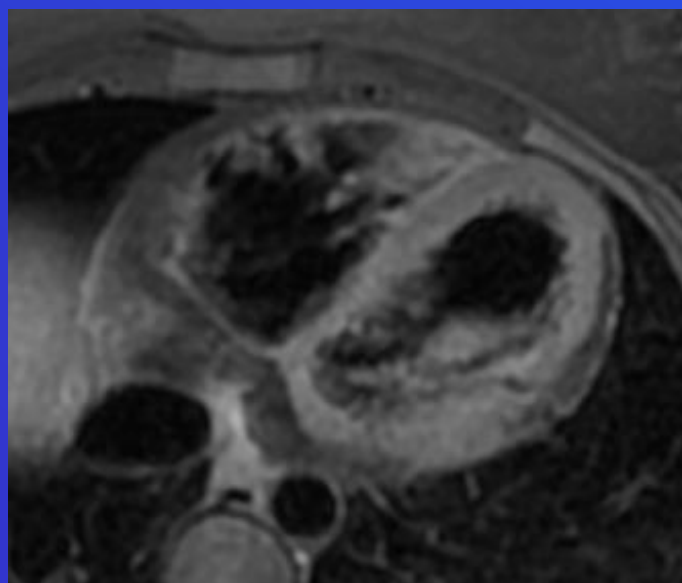


Chemical-Shift

# FSE and Double IR

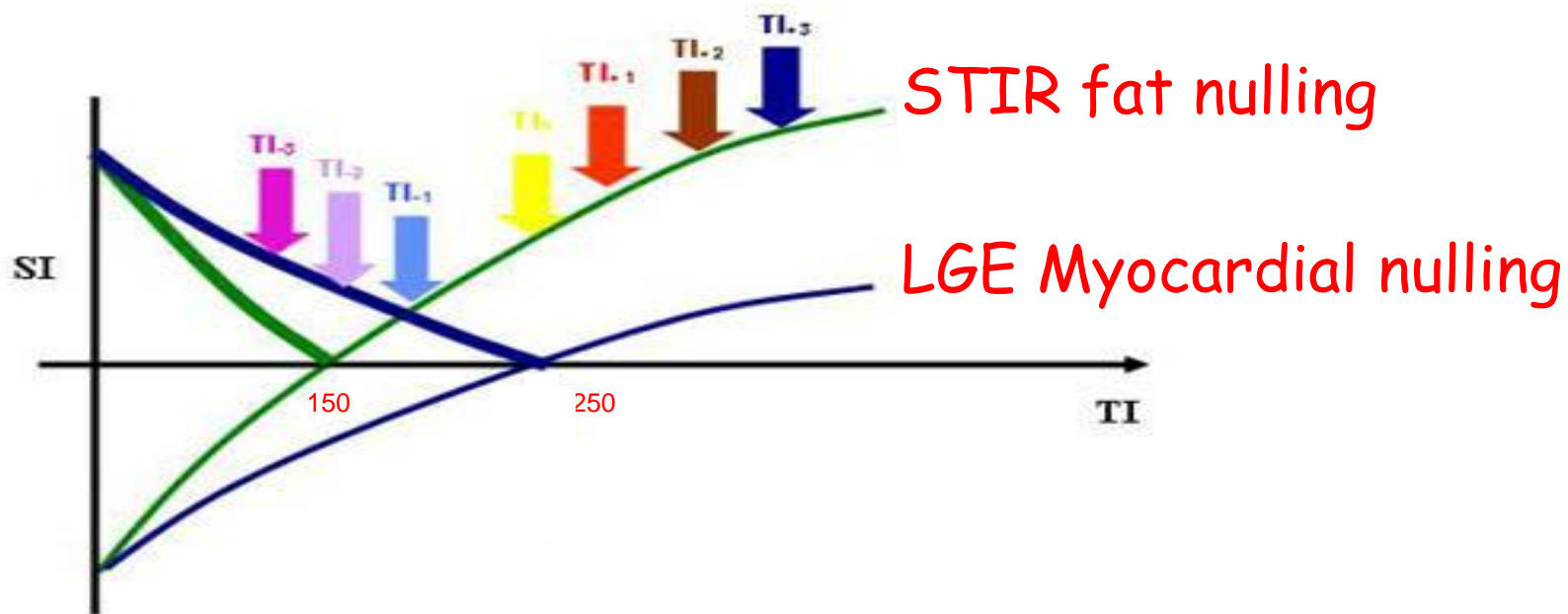
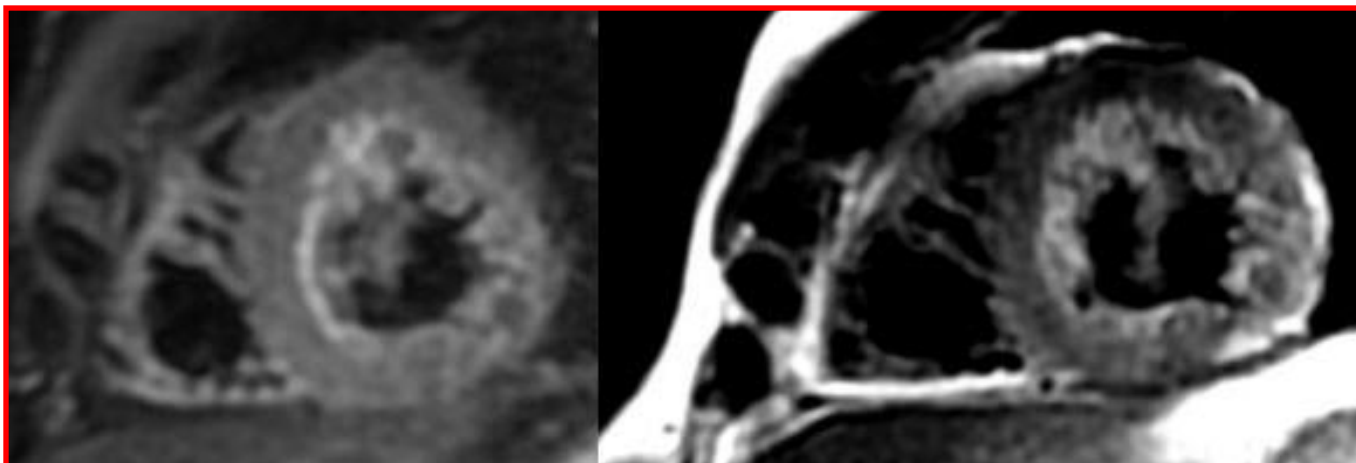


FSE



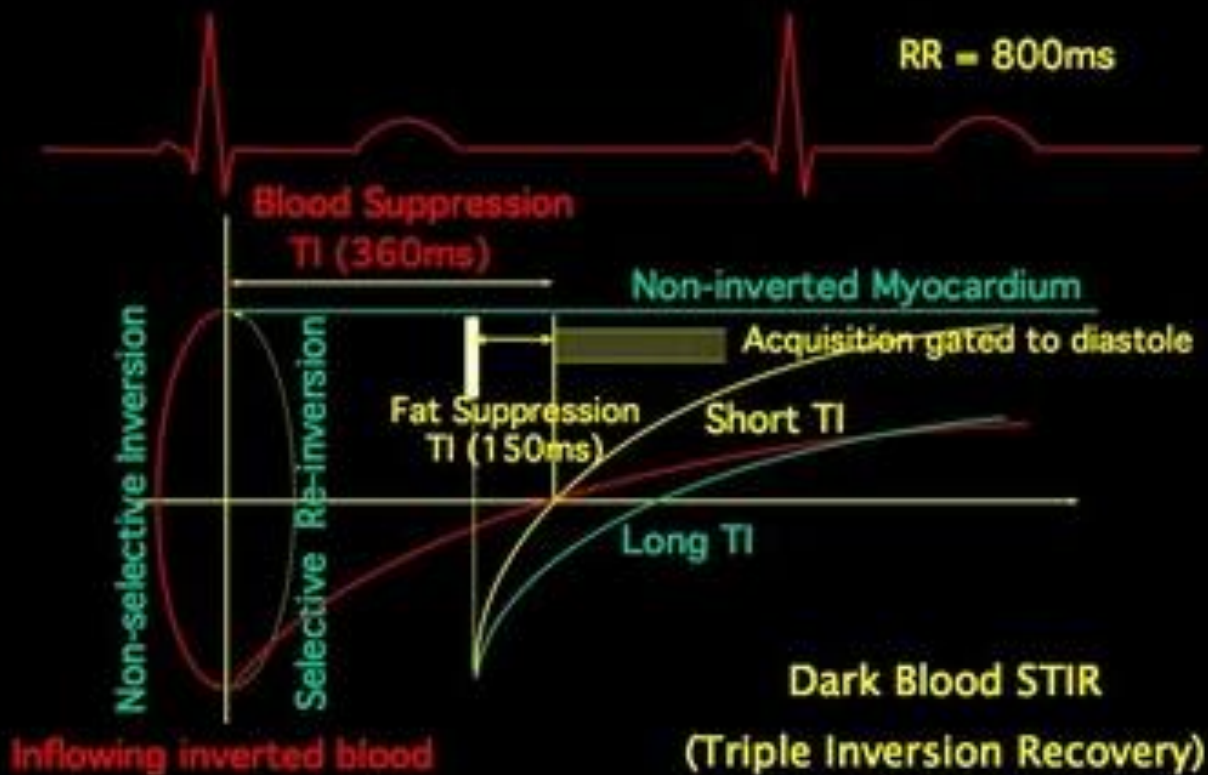
FSE FatSat

# FSE and Double IR





# STIR: TRIPLE IR



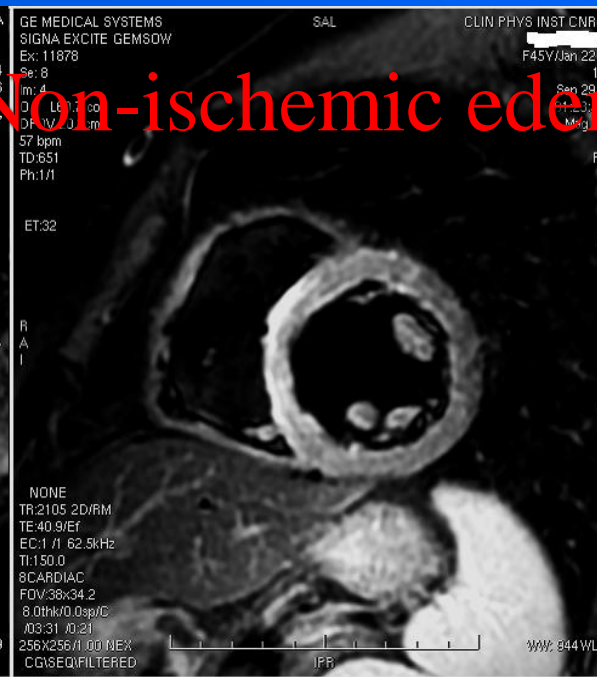
The third IR pulse and the fat signal are shown in yellow.

The delay time for fat to cross its null point is much shorter.

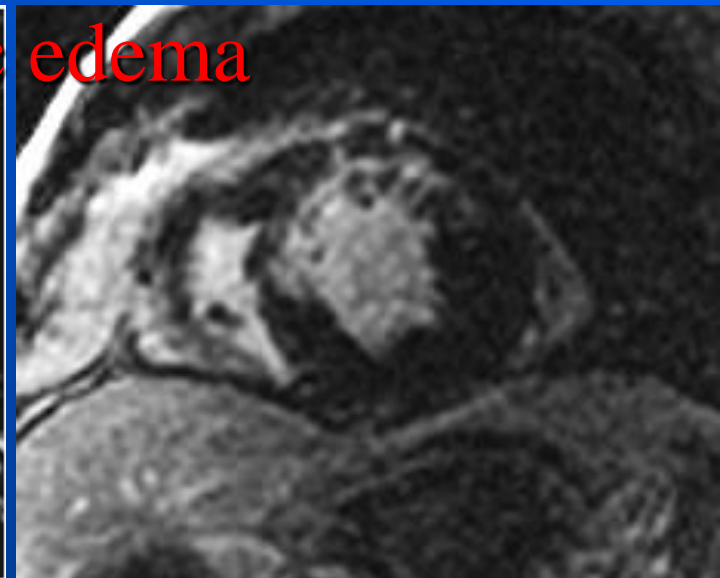
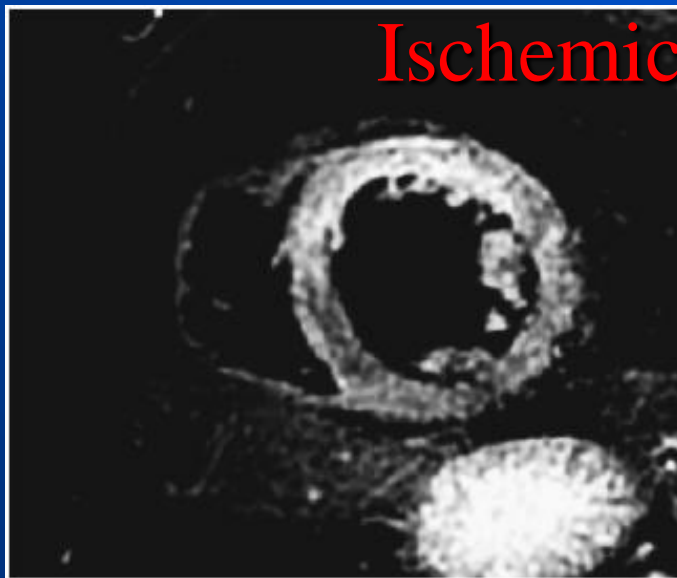
Both the blood and fat cross their null points at the same time that data is collected,  
both blood and fat will be dark in the image.



# T2-STIR MYOCARDIAL EDEMA

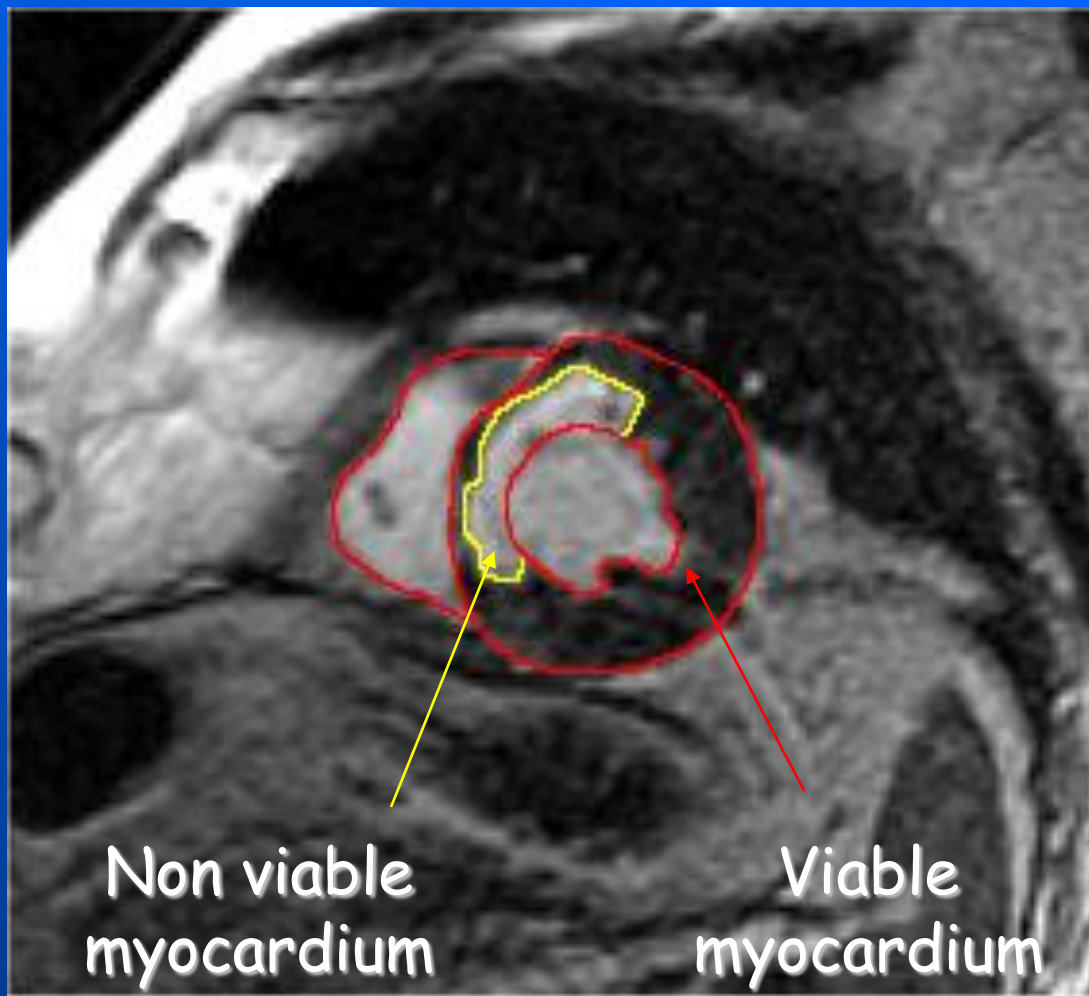
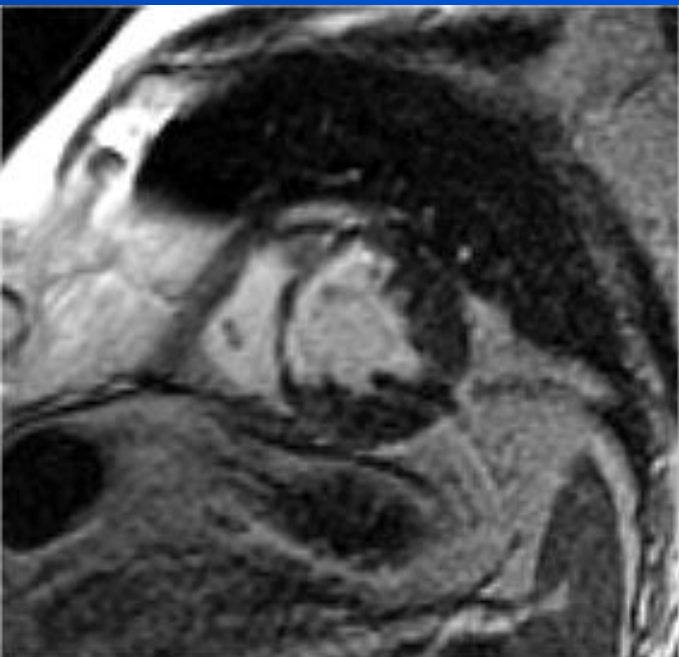


Non-ischemic edema





# LATE GADOLINIUM ENHANCEMENT



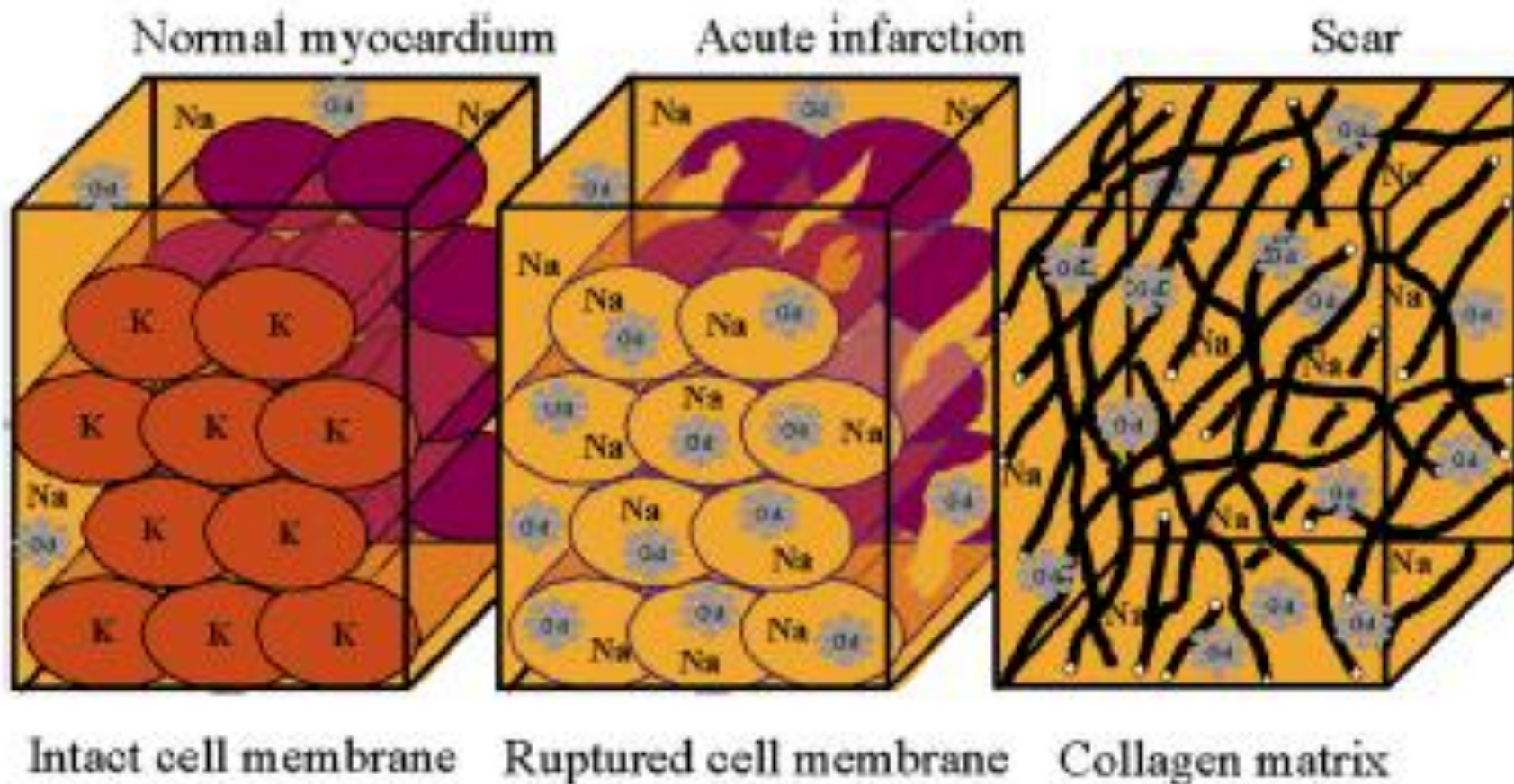
Inversion Recovery GRE T1w post mdc  
TI to null myocardium

# LATE ENHANCEMENT



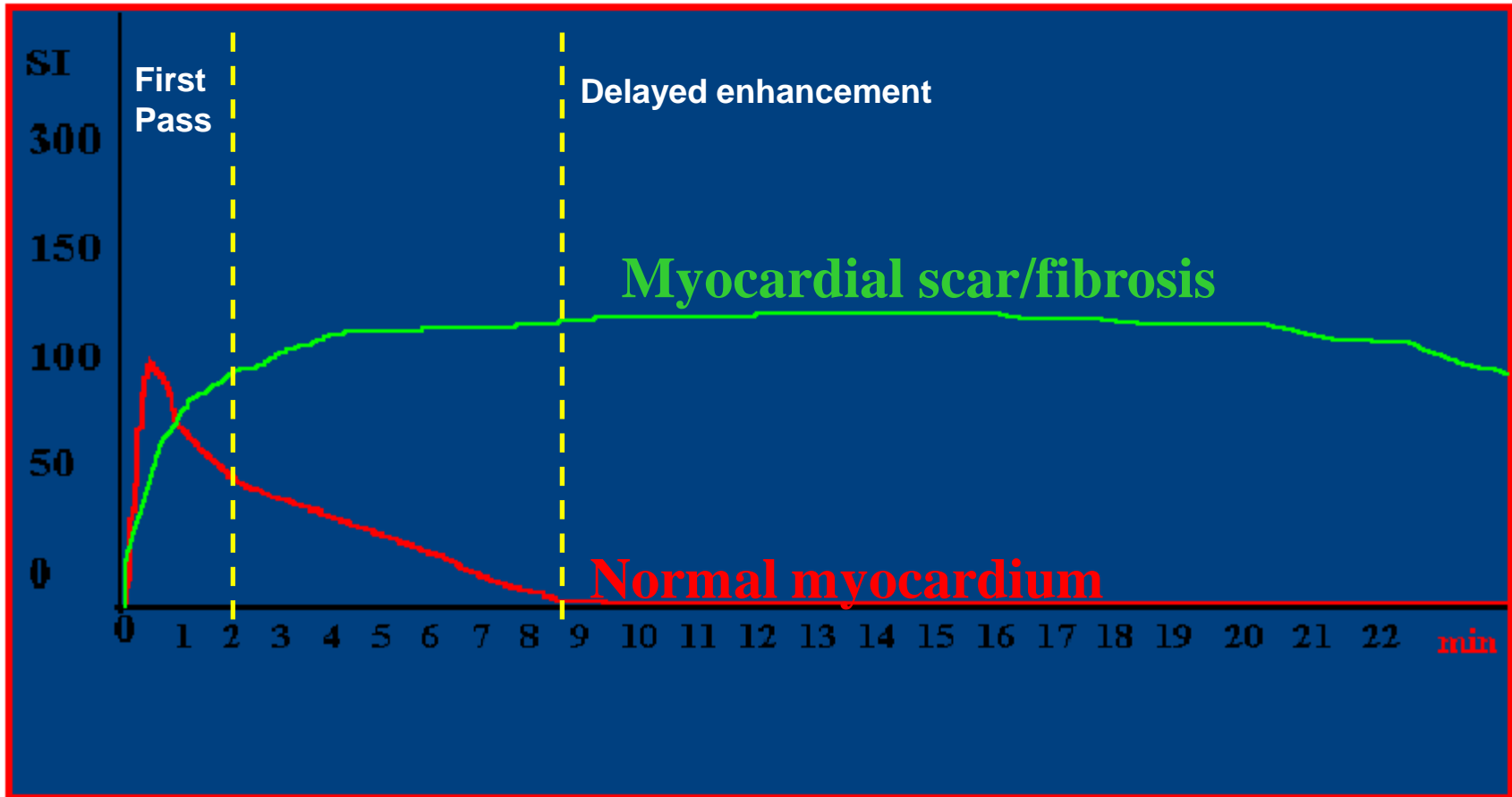
The mechanism is a combination of:

- delayed wash-in and wash-out kinetics of nonviable tissue
- acute and chronic disarrangement of interstitium
- different volumes of distribution of Gd in viable and nonviable tissue





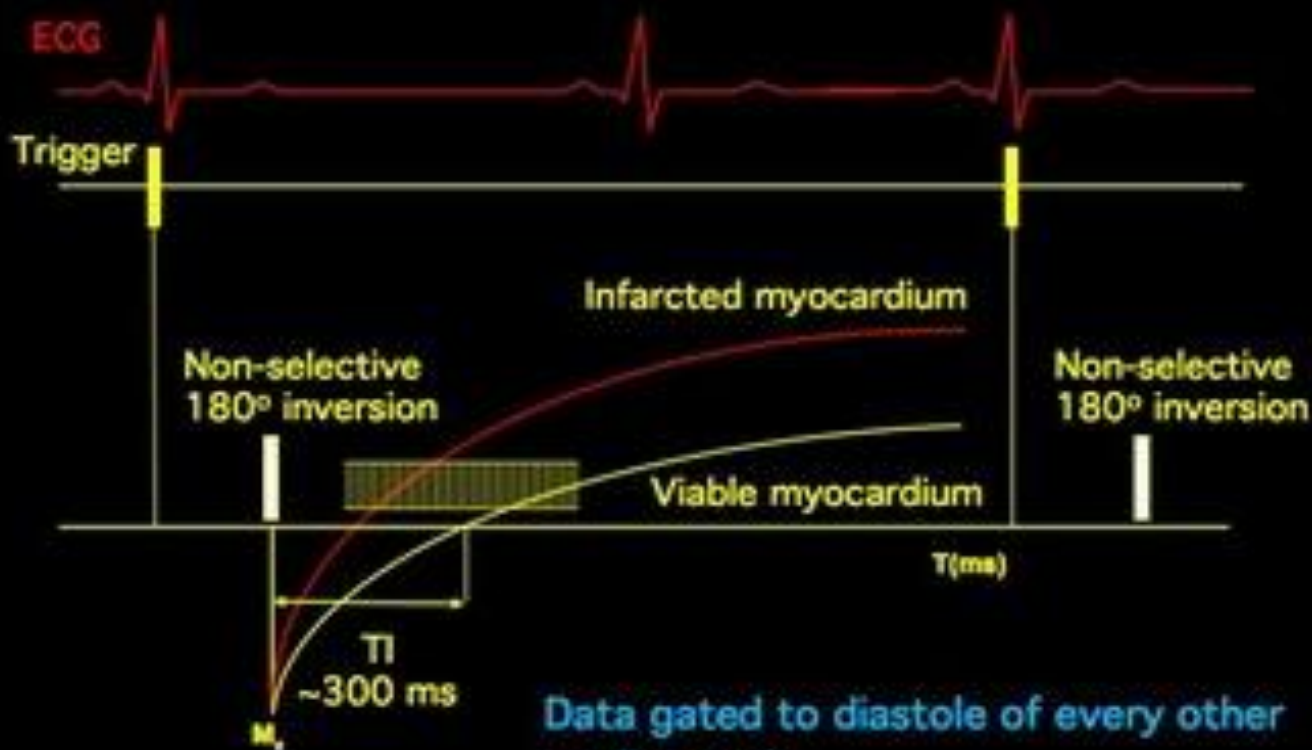
# LATE GADOLINIUM ENHANCEMENT





# LATE GADOLINIUM ENHANCEMENT

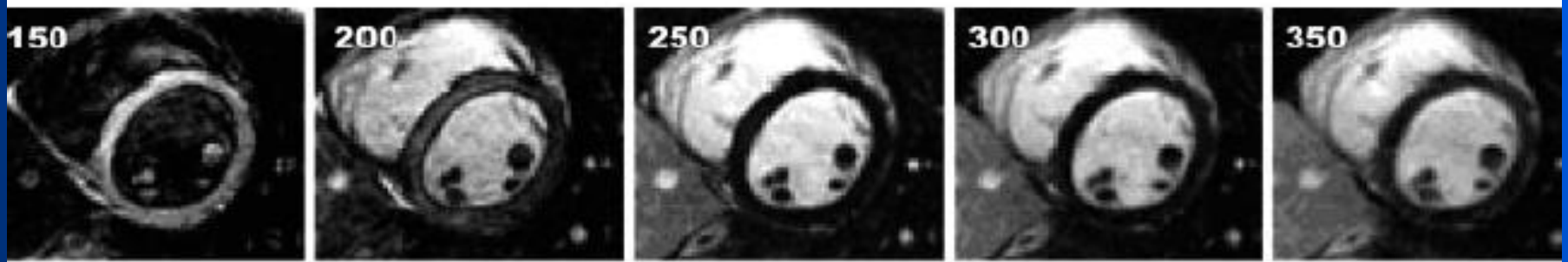
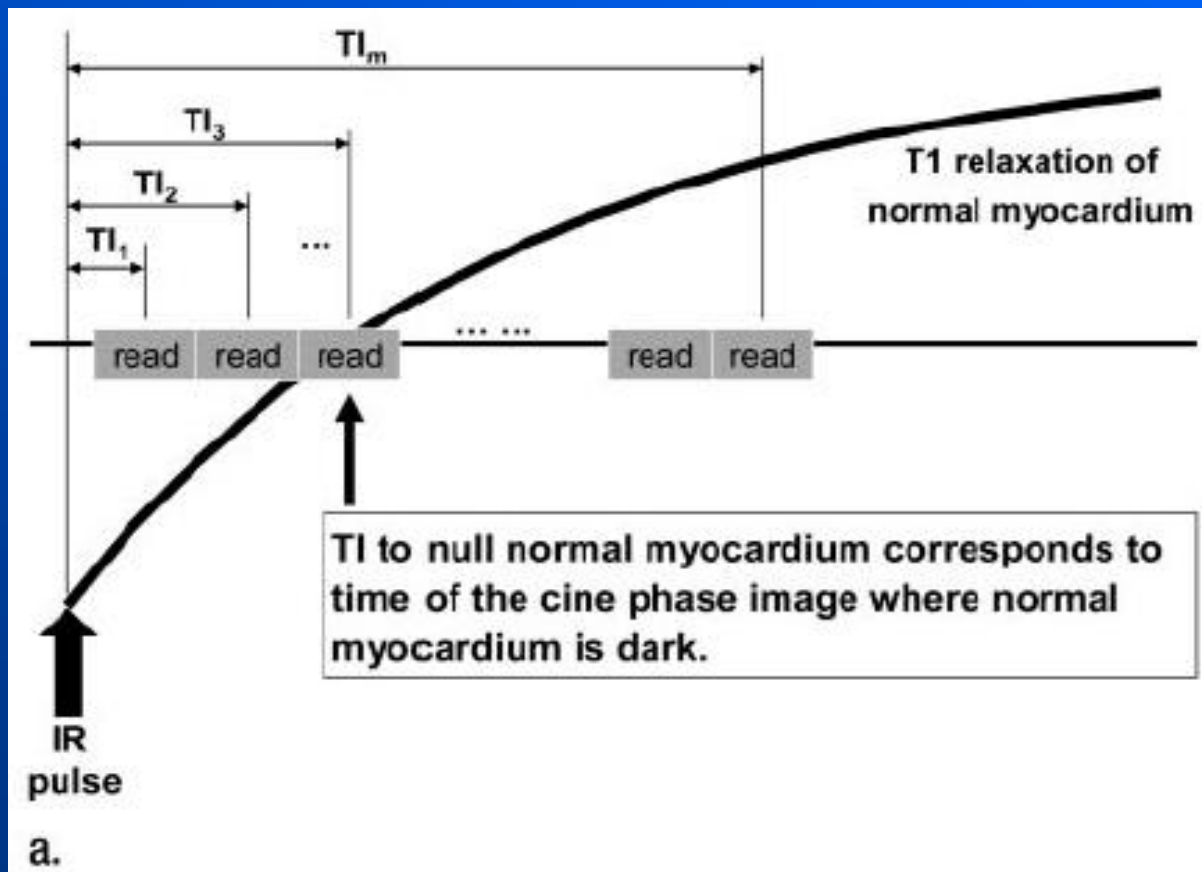
## Inversion Recovery – TurboFLASH



Data gated to diastole of every other cardiac cycle  
TI adjusted to null viable myocardium  
Infarct is brighter than viable myocardium



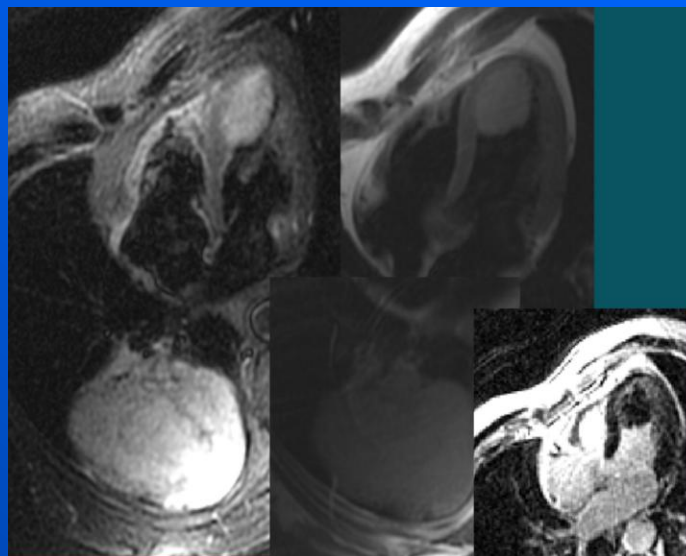
# LATE GADOLINIUM ENHANCEMENT



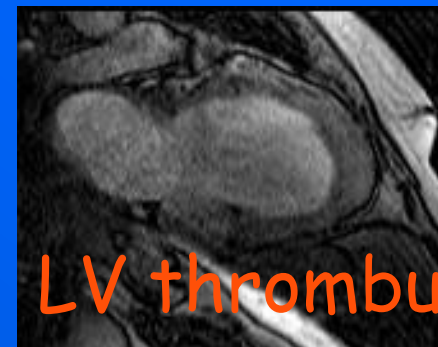
# Applications of Late Enhancement



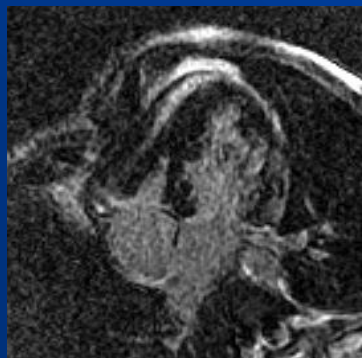
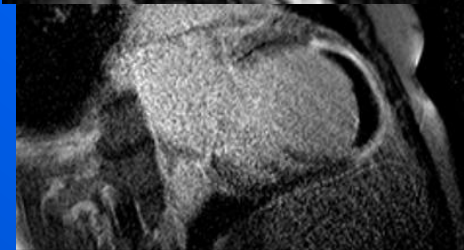
Infarction



Mass



LV thrombus



Myocarditis



Sarcoidosis



HCM

Cardiomiopathy





THANK YOU!

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[mrilab@ftgm.it](mailto:mrilab@ftgm.it)

