



中華民國醫事放射學會

Taiwan Society of Radiological Technologists (TWSRT)

MRI
Lecture 1

MRS

原理與臨床應用

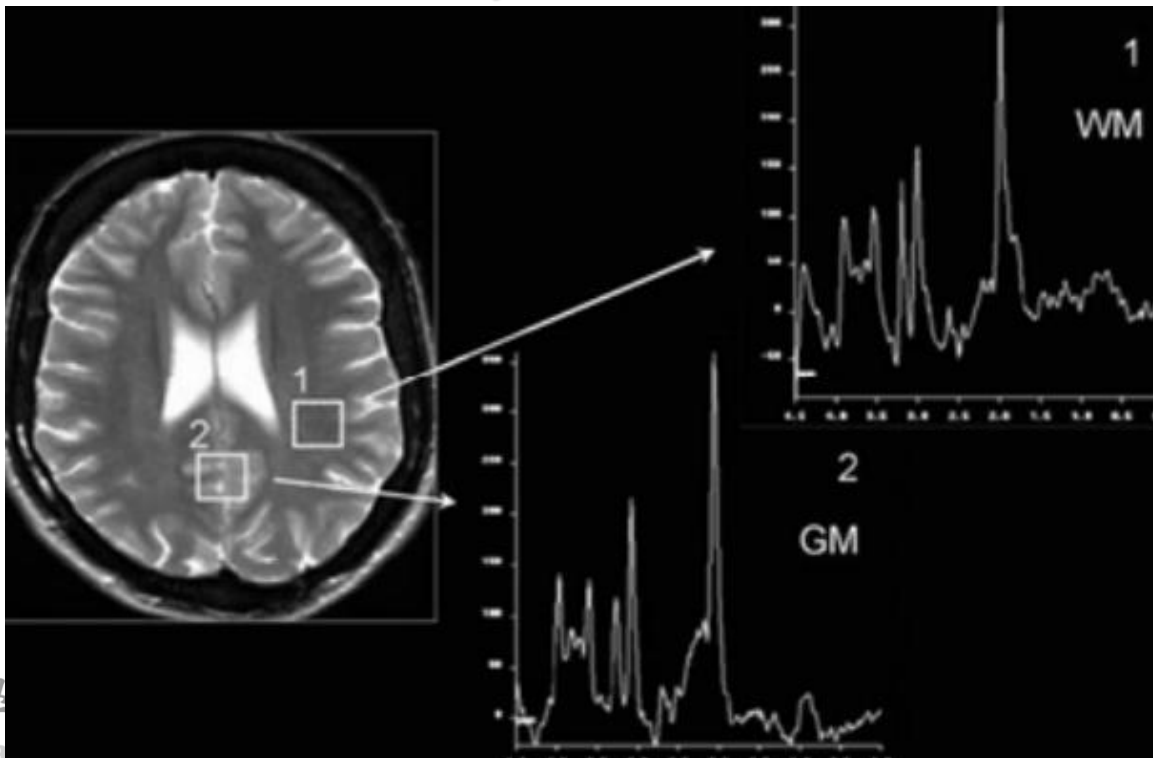
台北醫學大學 一部立雙和醫院
影像醫學部 醫學物理師 李宜恬

2017.04.16

國泰

What's MR Spectroscopy?

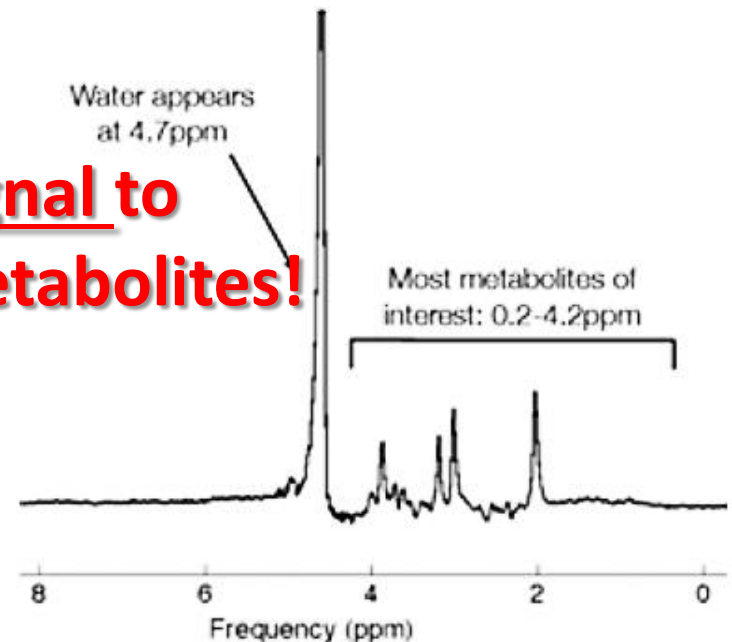
- Rather than providing images, it usually provides spectra consisting of individual peaks, **the chemical shift of metabolites.**
- Provide **bio-chemistry information.**



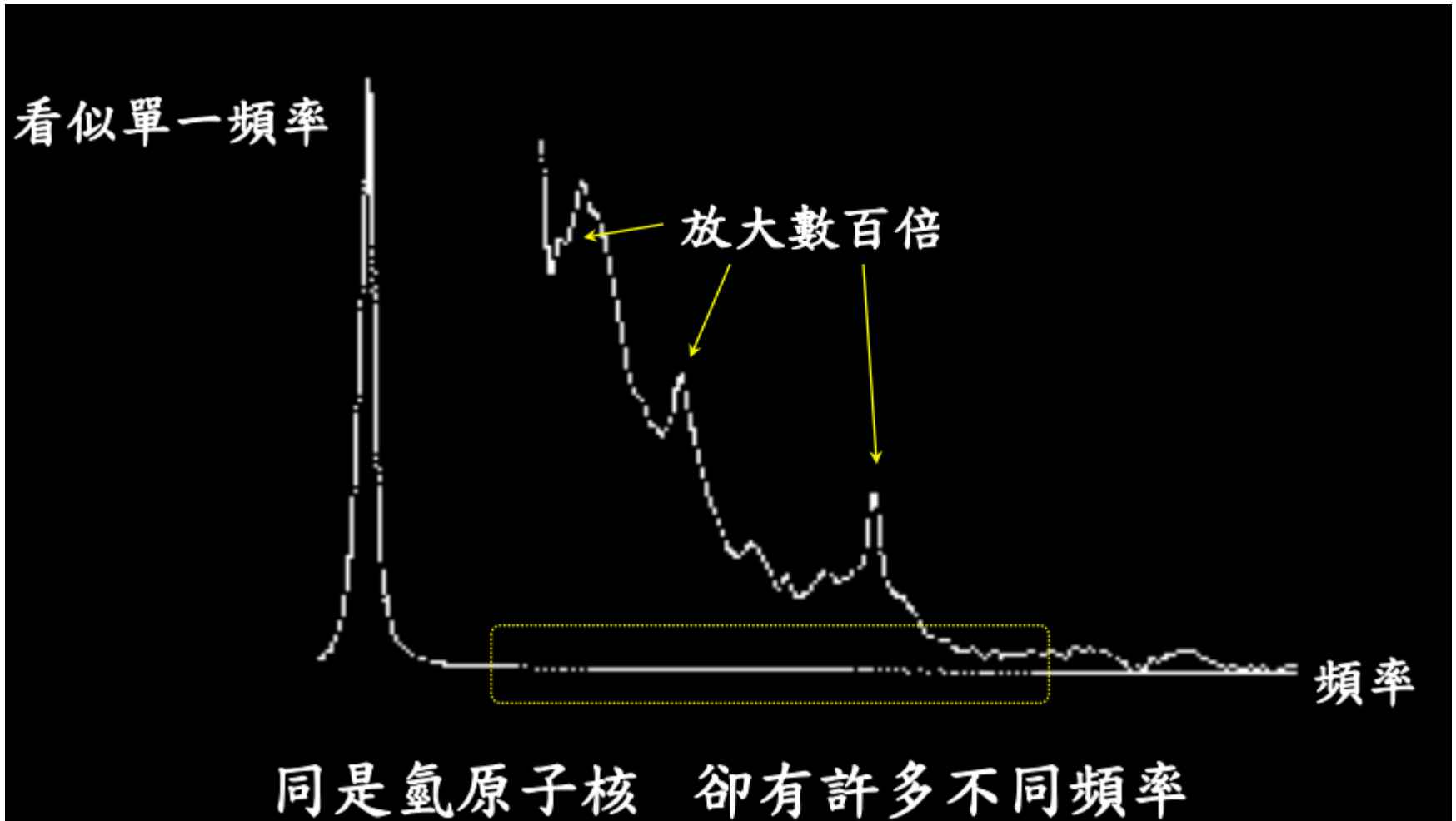
^1H (Proton) Spectroscopy

- **Proton spectroscopy** is easier to perform and provides much **higher SNR** than either sodium or phosphorus.
- Proton concentration in water $\sim 100\text{M}$
Other metabolites: $1 \sim 10\text{mM}$

✓ **Need to suppress the water signal to investigate the signals from metabolites!**

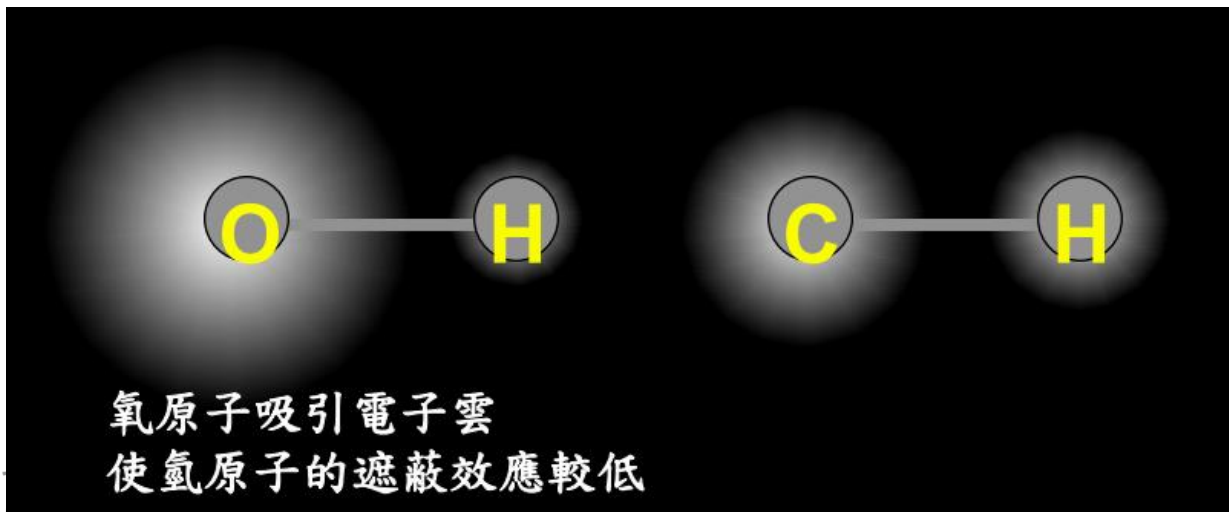


^1H (Proton) \rightarrow Single Frequency?

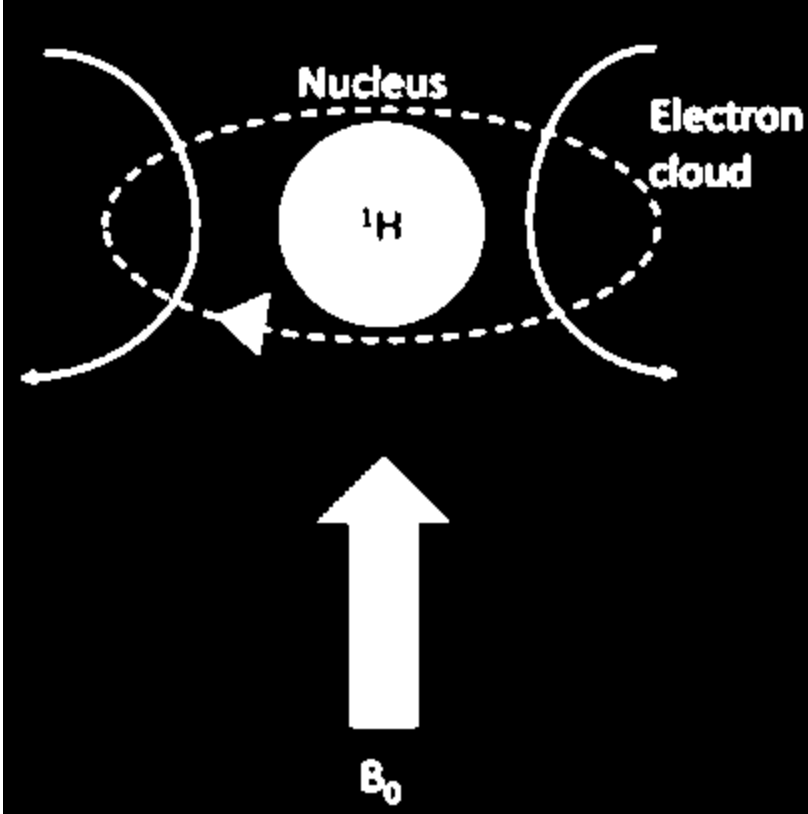


Different frequency?

- Different chemical environment.
- Different atom bonding.
- Electron screening effect (**Shielding effect**, 電子雲遮蔽效應)
 - Magnetic field weakened \rightarrow frequency \downarrow
 - **Chemical Shift**



Chemical Shift & Shielding Effect



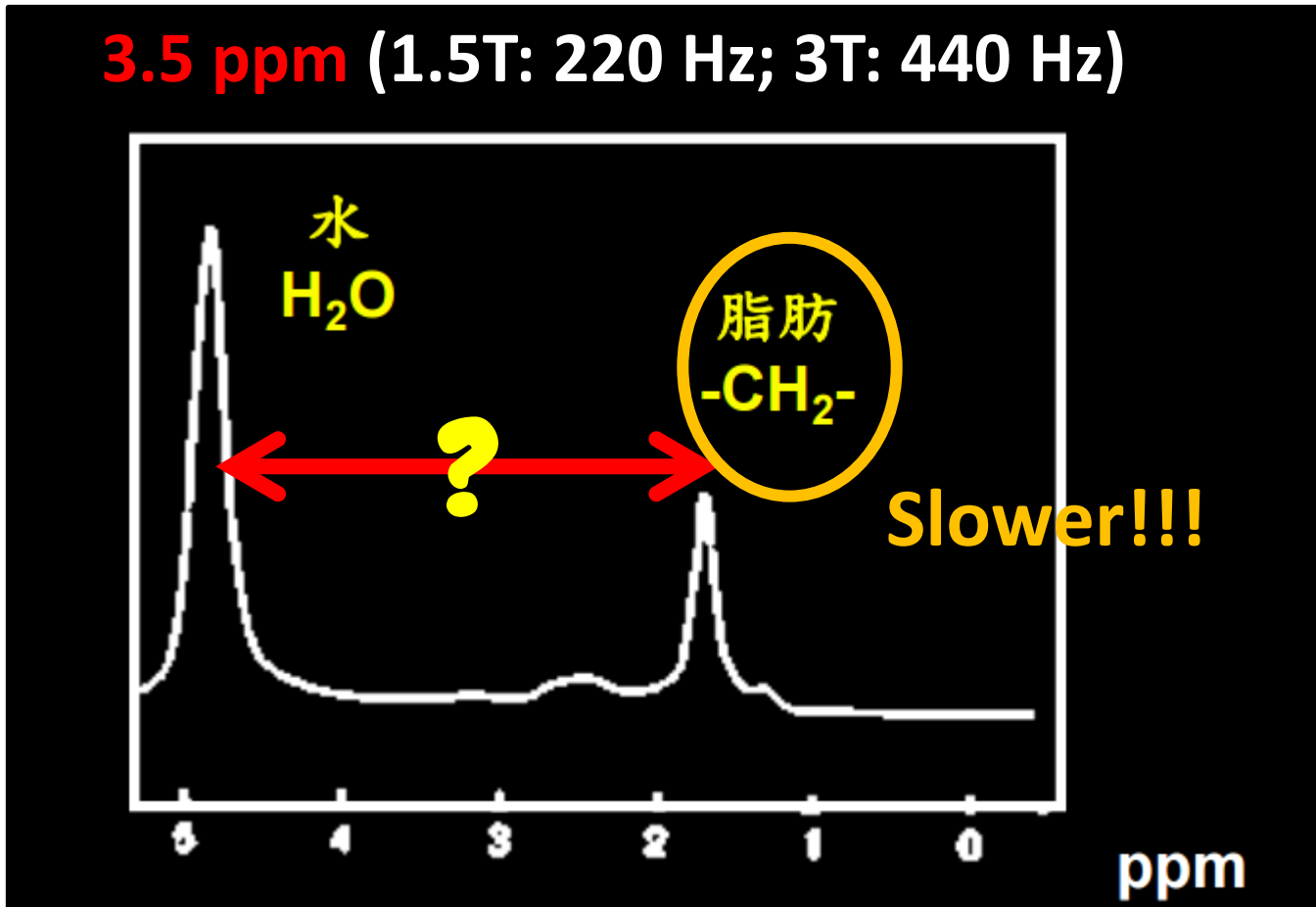
$B_{\text{local}} = -\sigma B_0$

$\omega = \gamma B_0(1-\sigma)$

$\sigma = \text{chemical shift shielding constant}$

For example: Chemical Shift (Review)

3.5 ppm (1.5T: 220 Hz; 3T: 440 Hz)

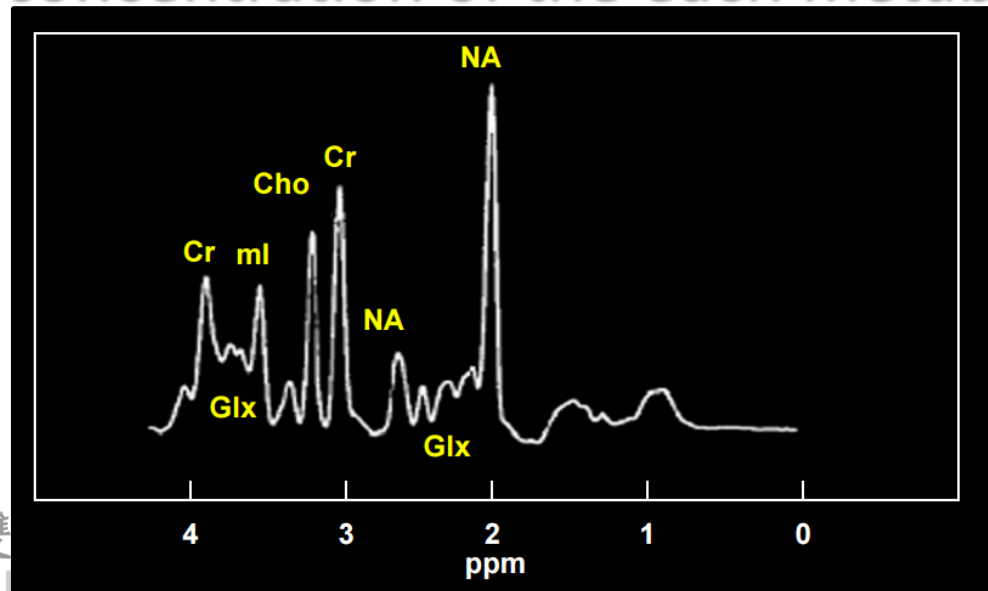


Upper field

Lower field

MR Spectroscopy

- The area under a given peak is proportional to the number of protons (**concentration**) contributing to the peak.
- MRS requires a species to be present in at least **1 mM** concentration to be seen.
- **Quantified** the concentration of the each metabolites!

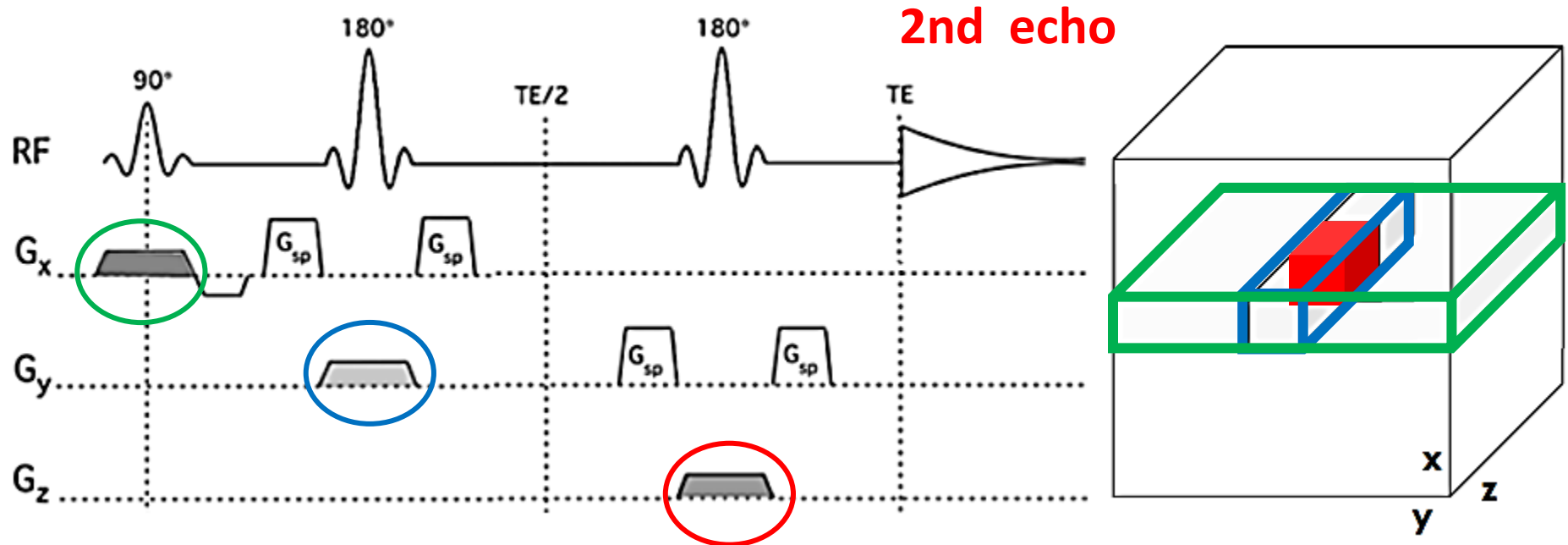


MRS pulse sequence

- **Localization**: Covering lesion and normal sites for comparison
- Two major sequences
- Point-Resolved Spectroscopy, PRESS
- Stimulated Echo Acquisition Mode, STEAM

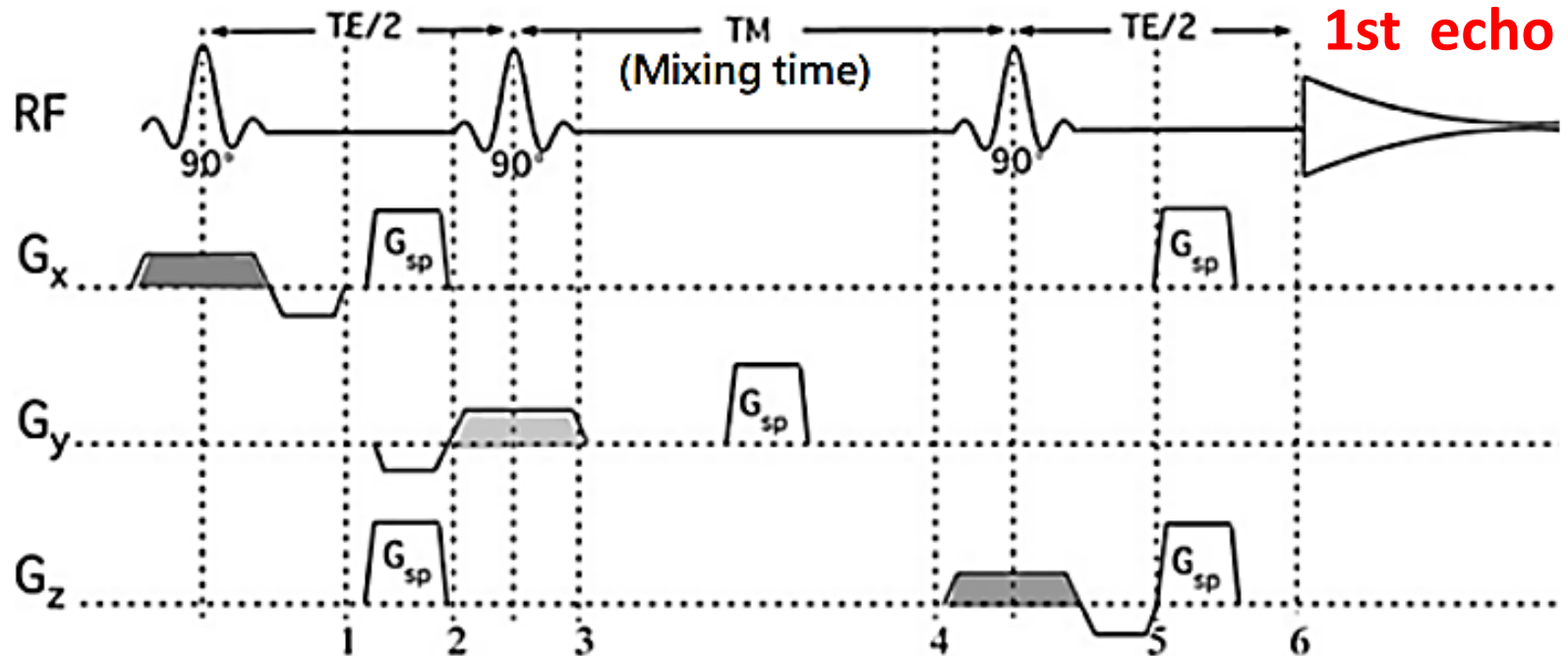


Point-Resolved Spectroscopy (PRESS)



- **Dual spin-echo sequence** consisting of 3 slice selective pulses in orthogonal planes (**90-180-180**)
- Signal comes from the **intersection** of the 3 planes!
- But....**TE too long!!** (Can't see short T_2 metabolites)

Stimulated Echo Acquisition Mode (STEAM)

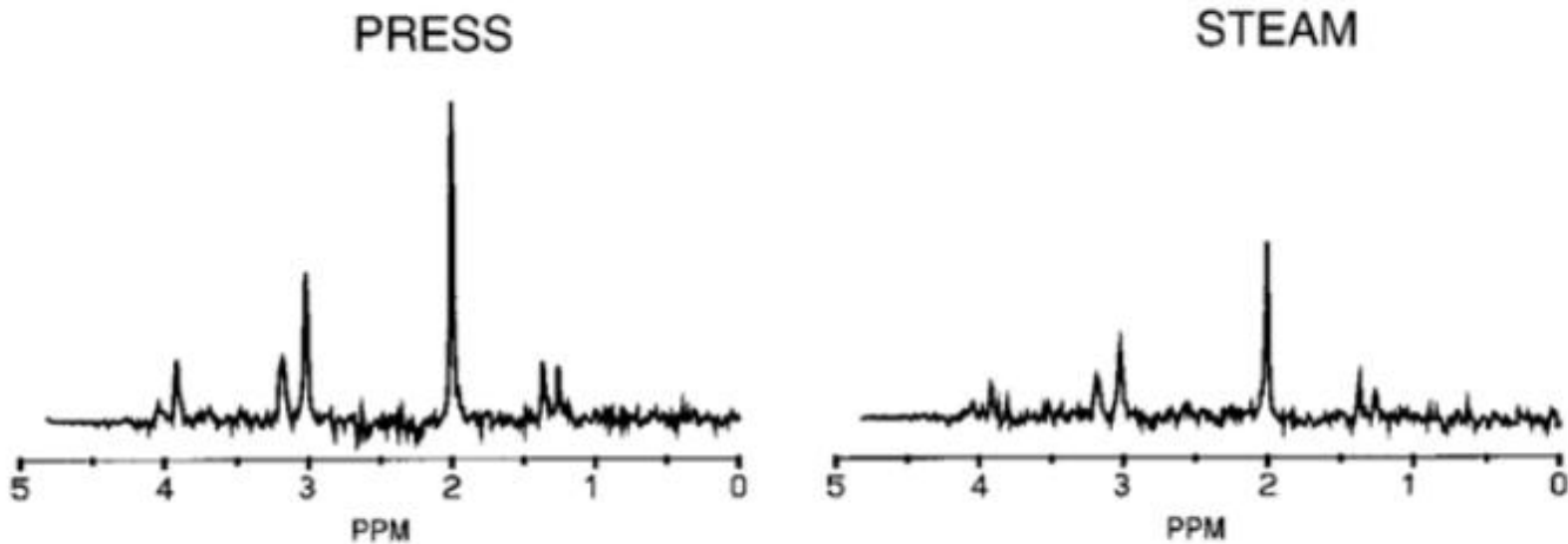


- Consists of three orthogonally slice selective 90 pulses
- Separate 180 to 90 + 90 (90 -90-90)
- T₂ decay does not occur during TM

But SNR decreased due to incompletely refocusing!!

PRESS vs. STEAM

- Stimulated echo amplitude is only half the size of a PRESS spin echo.



PRESS vs. STEAM

- Stimulated echo amplitude is only half the size of a PRESS spin echo.

	PRESS	STEAM	Note
SNR	S	S/2	PRESS SNR 2x STEAM SNR
TE	Short TE difficult	Short TE possible	STEAM: Better for metabolites with short T2
SAR	High	Low	90 transmit lower power than 180
Location	Sharp	Sharper	90 pulses have sharper profiles than 180s

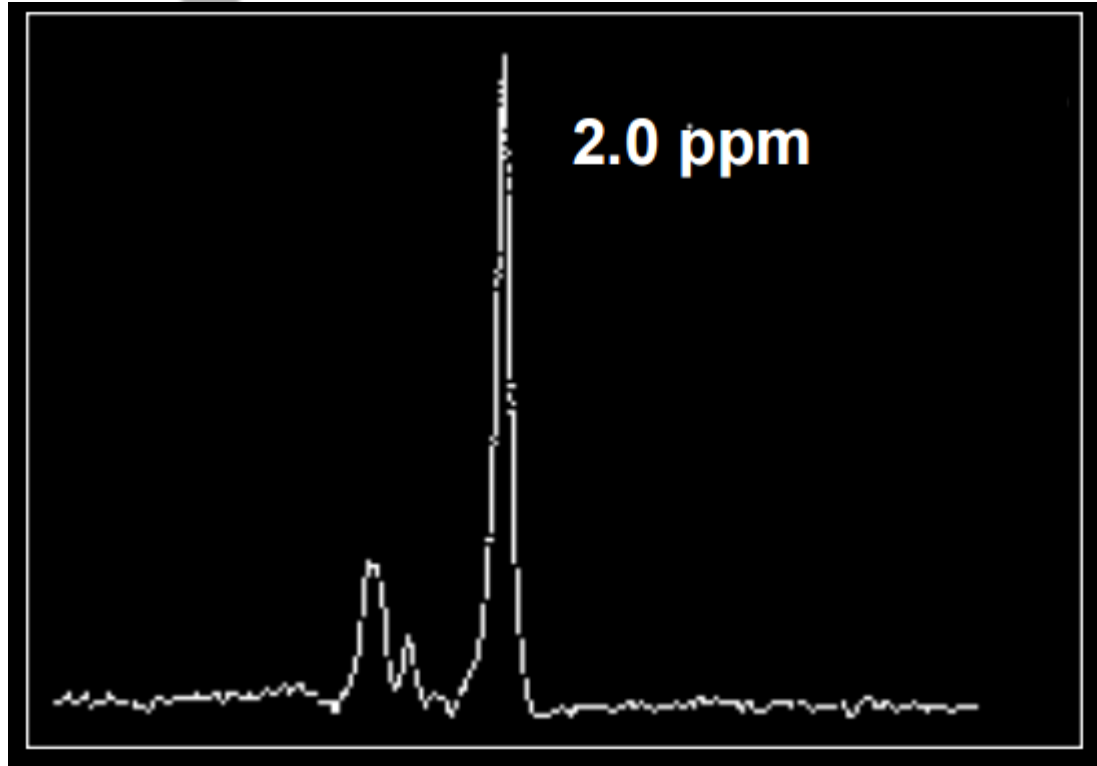


Short T₂ Metabolites

- Short T₂ → Wide Spectrum
- Wide spectrum → Wide bandwidth → dephasing faster → Short T₂

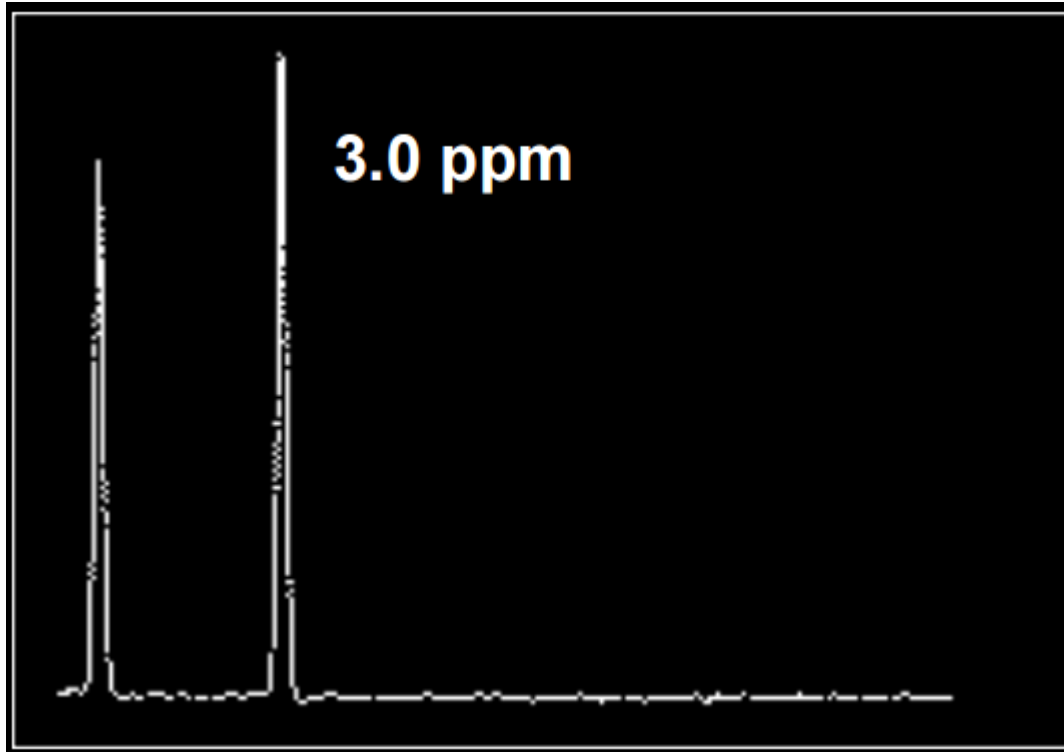


N-acetylaspartate (NAA) : Long T₂



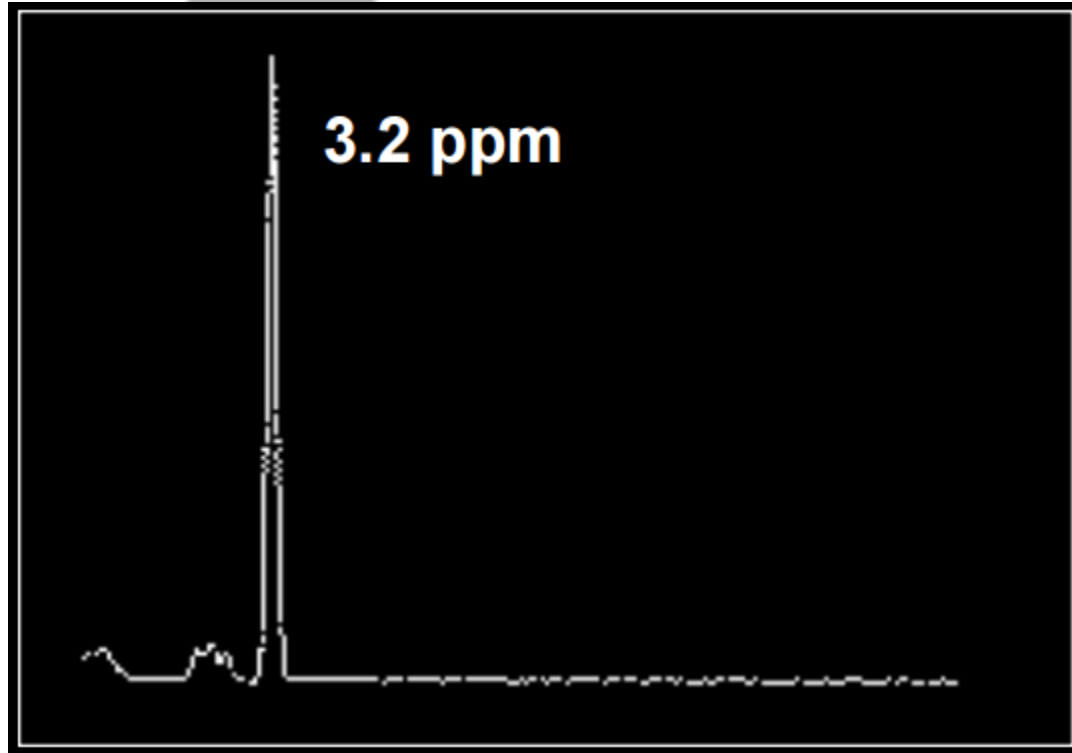
- A neuronal marker
- Register marker: Adjust the frequency shift across subject and session.

Cr/ PCr: Long T₂



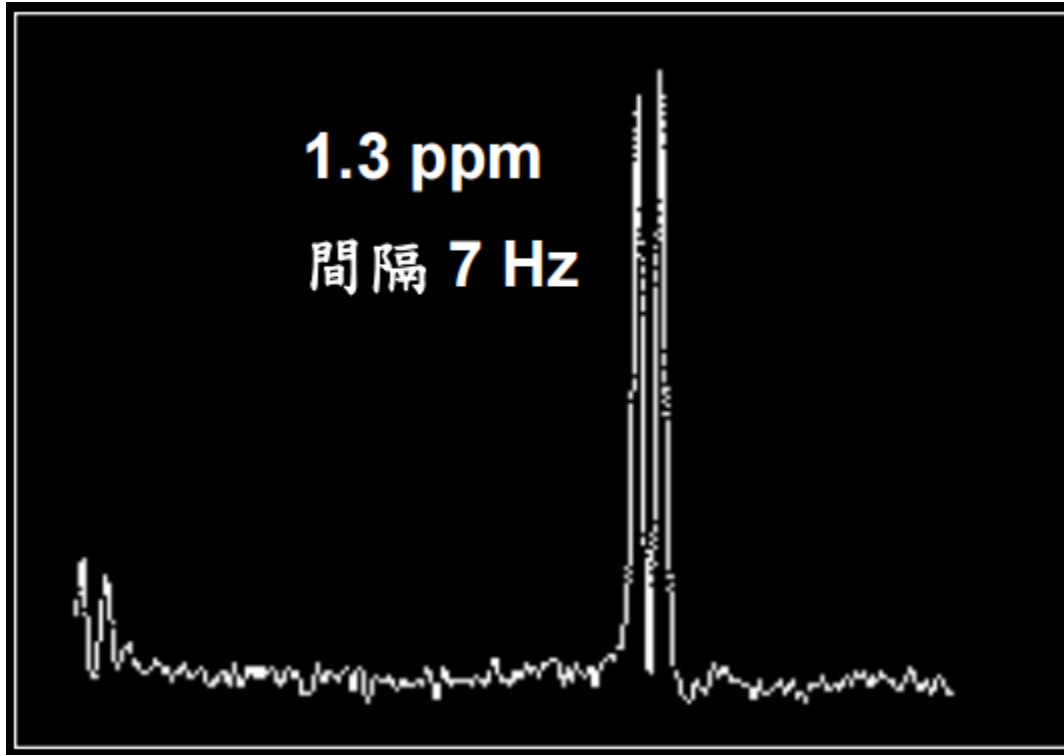
- Provide Phosphate

Choline: Long T₂



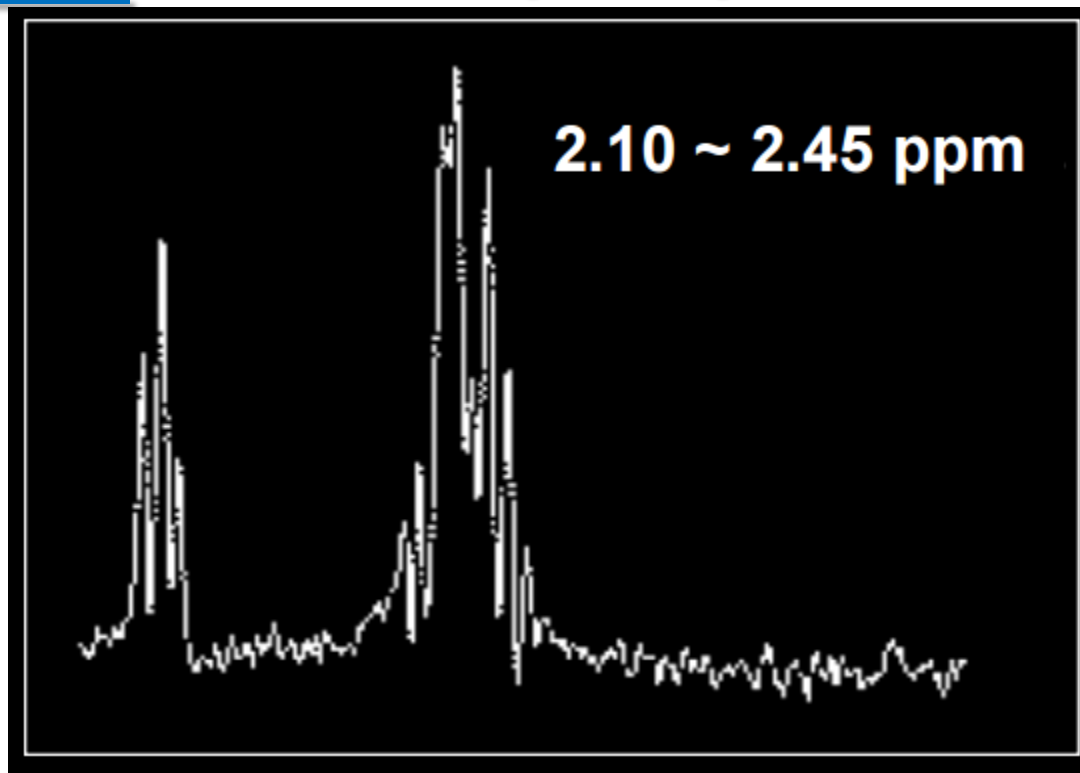
- Neurotransmitter (**Acetylcholine**) and others.

Lactate: Long T₂



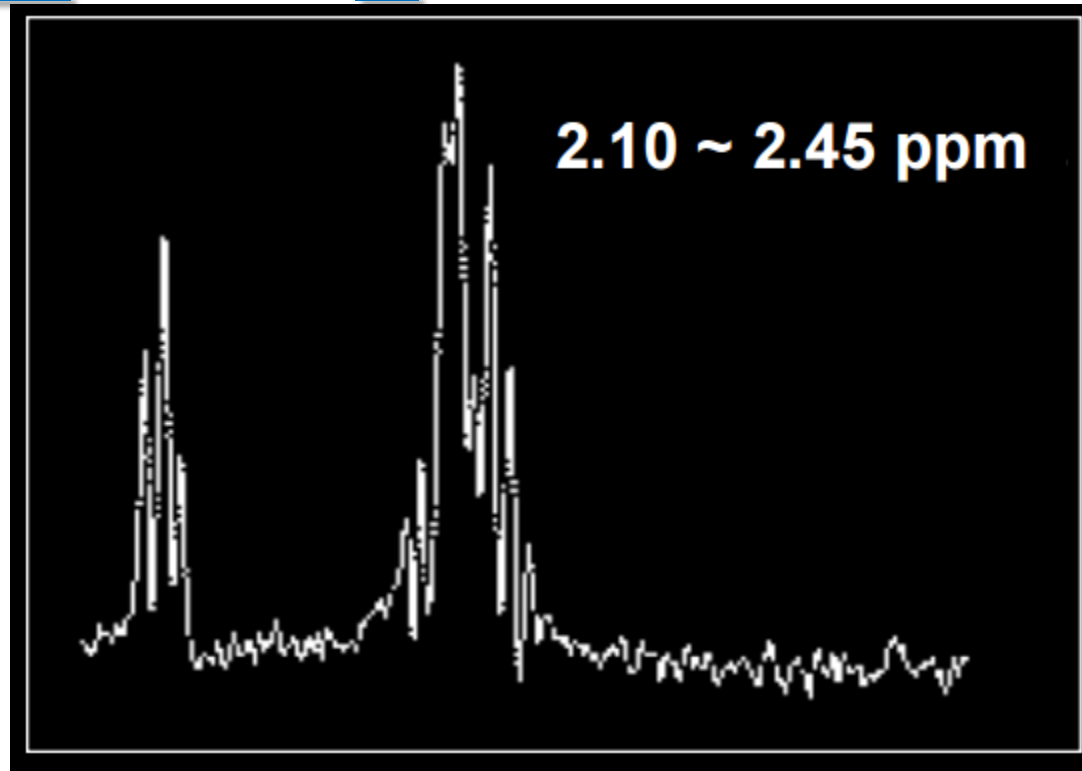
- Product of **anaerobic metabolism**
- **J-coupling**: methyl and methine share a bond → peak splitting (**increase with TE**)

Glutamate (Glu): Short T2



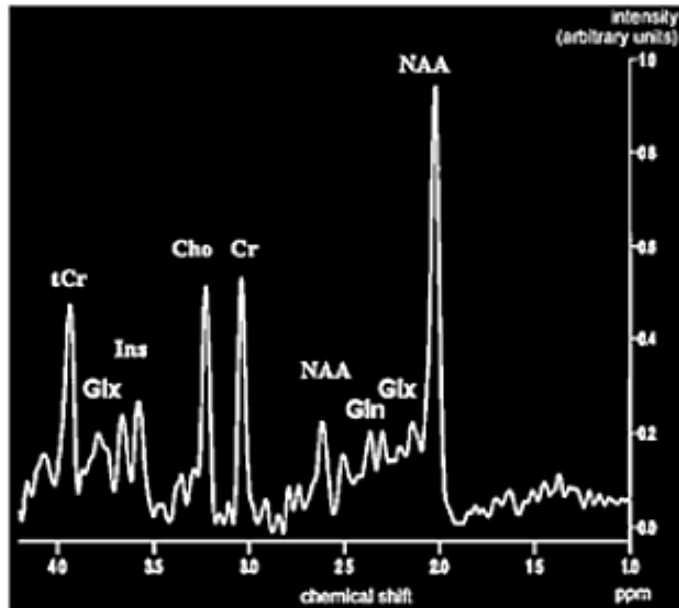
- **Excitatory** neurotransmitter

Glutamine (Gln): Short T₂



- Product of glutamate

MRS peaks in Brain



4.0 ppm ← 1.0 ppm

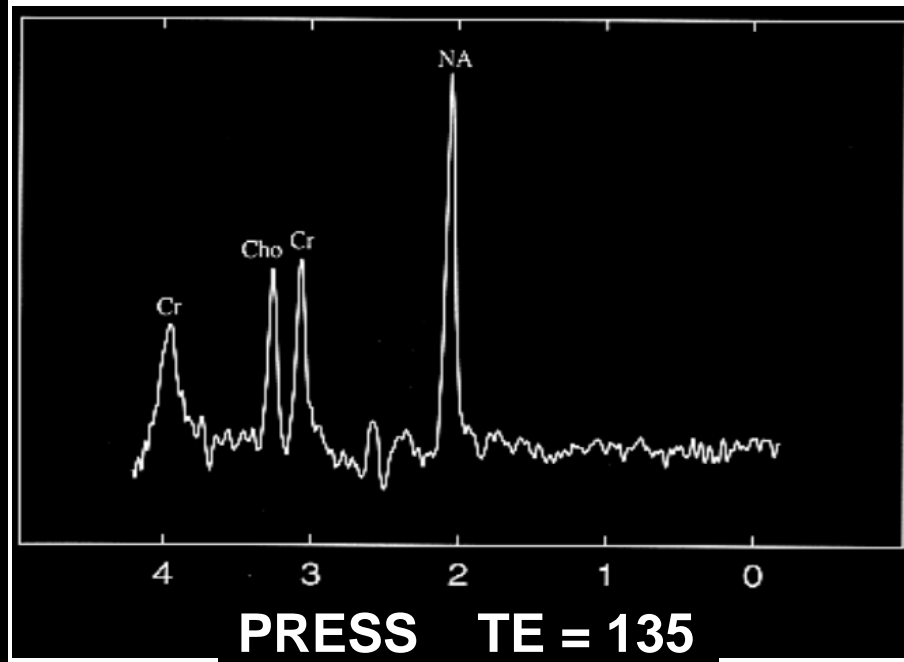
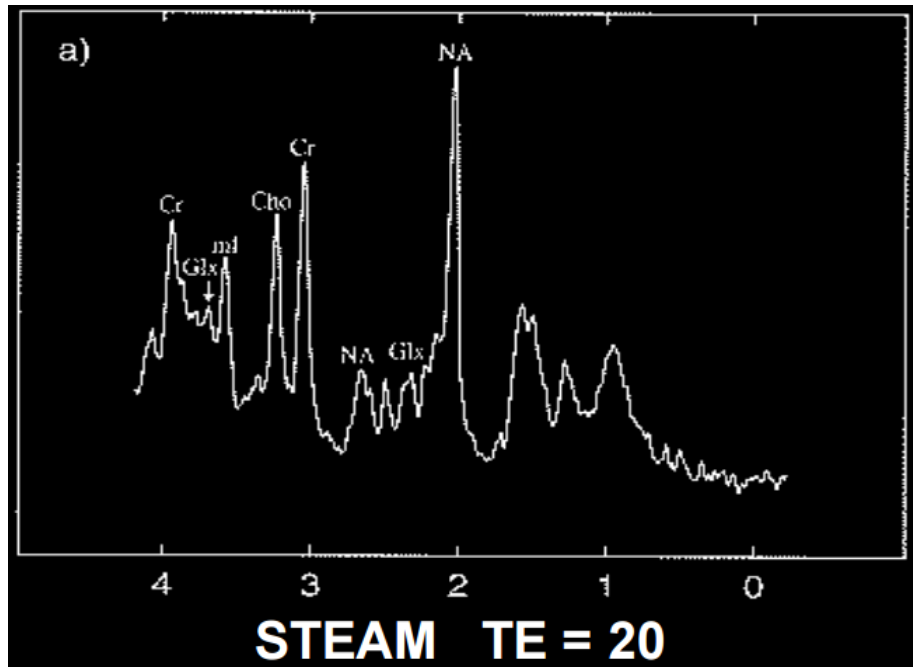
Metabolite	Major Resonance (ppm)	Effect	Visible only at short TE
Lipids (Lip)	0.9, 1.3	Breakdown of tissue	Y
Lactate (Lac)	1.3	Marker of anaerobic glycolysis	N
N-acetyl aspartate (NAA)	2.0	Marker of neuronal health	N
Glutamate/Glutamine (Glx)	2.1, 3.8	Excitatory neurotransmitter	Y
Choline (Cho)	3.2	Marker of membrane metabolism, cell proliferation	N
Creatine (Cr)	3.0	Marker of cellular energetics	N
Myo-inositol (MI or Ins)	3.5, 3.6	glial cell marker	Y

Short T₂ Metabolites

- Short T₂ → Wide Spectrum
- Wide spectrum → Peak Broadening → dephasing faster → Short T₂
- Looks like baseline drift??



Effect of TE



More information!

Specific!

Use **PRESS** at **long TE** metabolites could achieve **higher SNR!!**

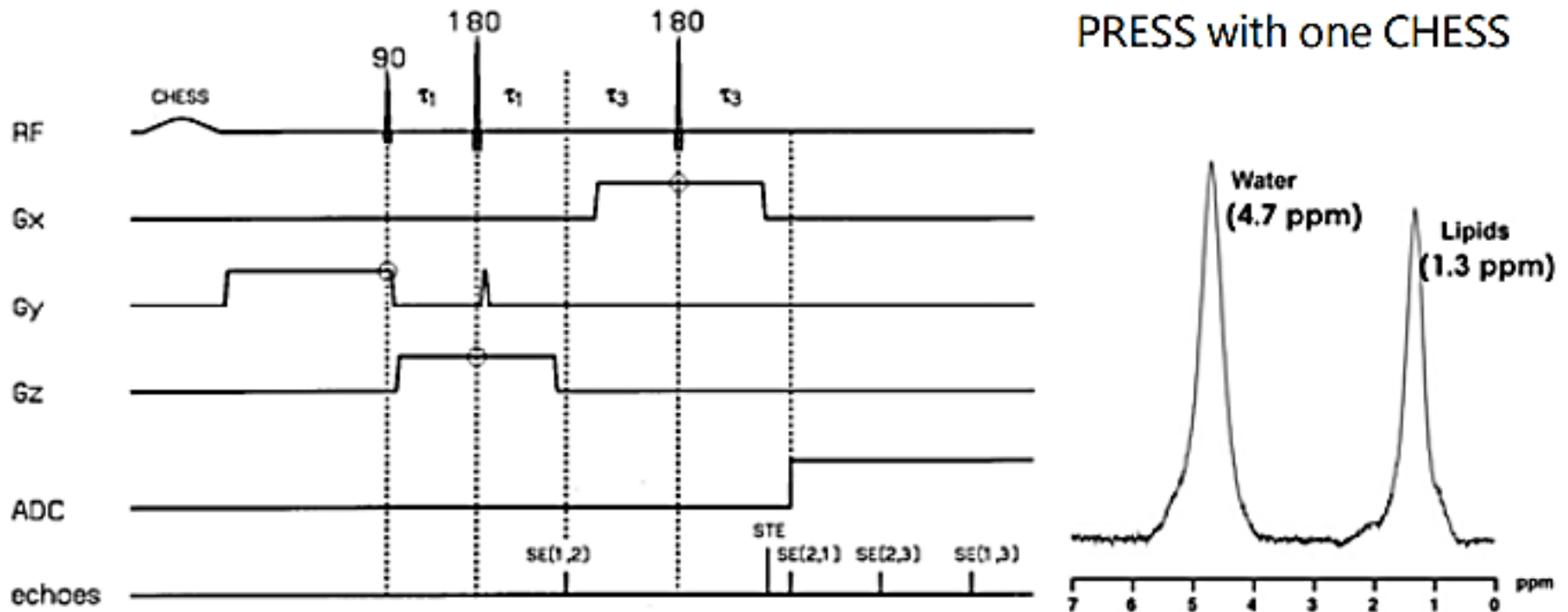
TR & TE in MRS

- Most institutions use a **TR of 1500 msec** and the shortest possible **TE of 30 or 35 ms** to **maximize the SNR**.
- This also allows the detection of **short T₂** species (like myoinositol and lipid), which would otherwise have already decayed at longer TE.
- **Peak width** is proportional to **1/T₂**, thus short T₂ species will lead to peak broadening.



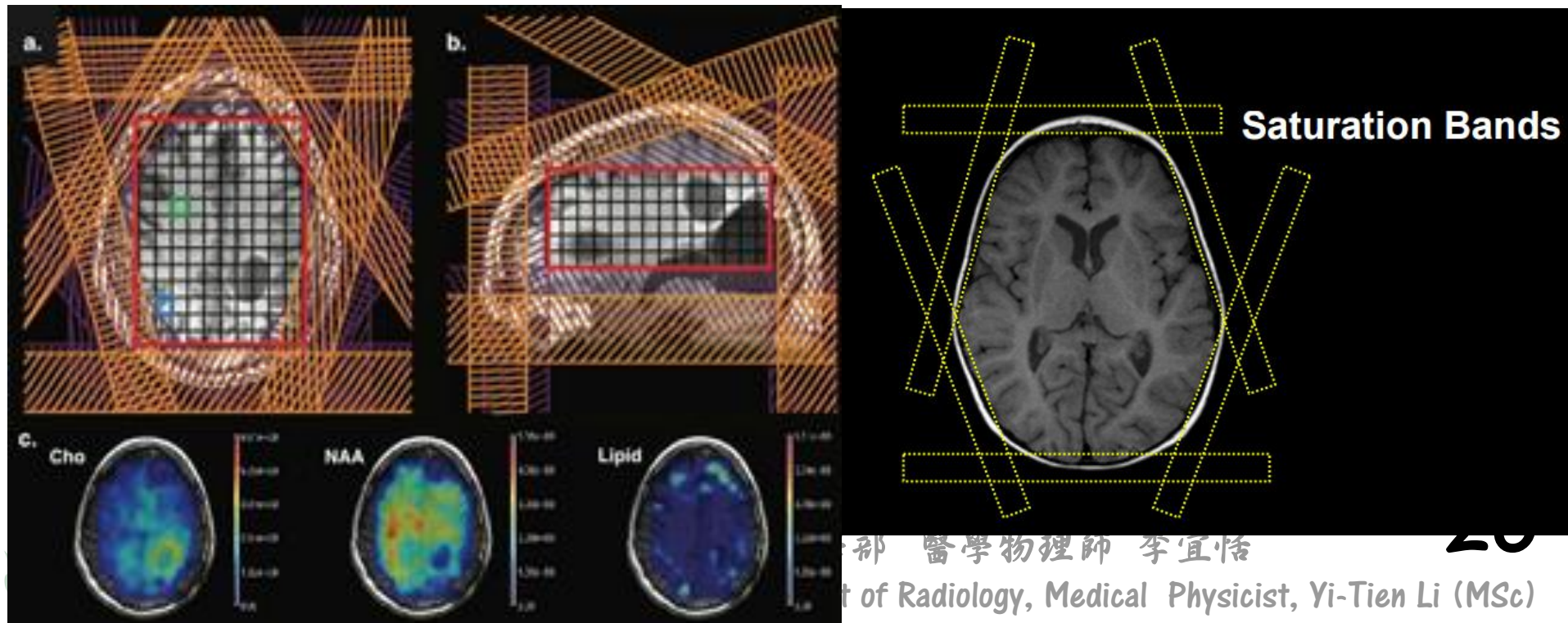
Water Suppression

- Chemical Shift Selection, CHESS
 - Frequency-selective presaturation pulse



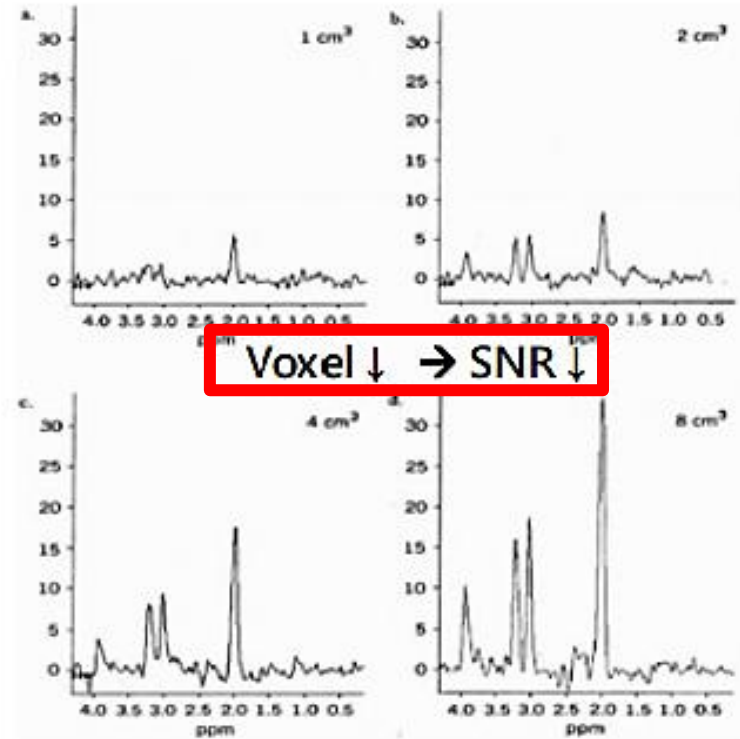
Fat Suppression

- Add **spatial saturation** bands
 - Outer Volume Suppression, OVS
- Why we didn't use frequency-selective saturation?



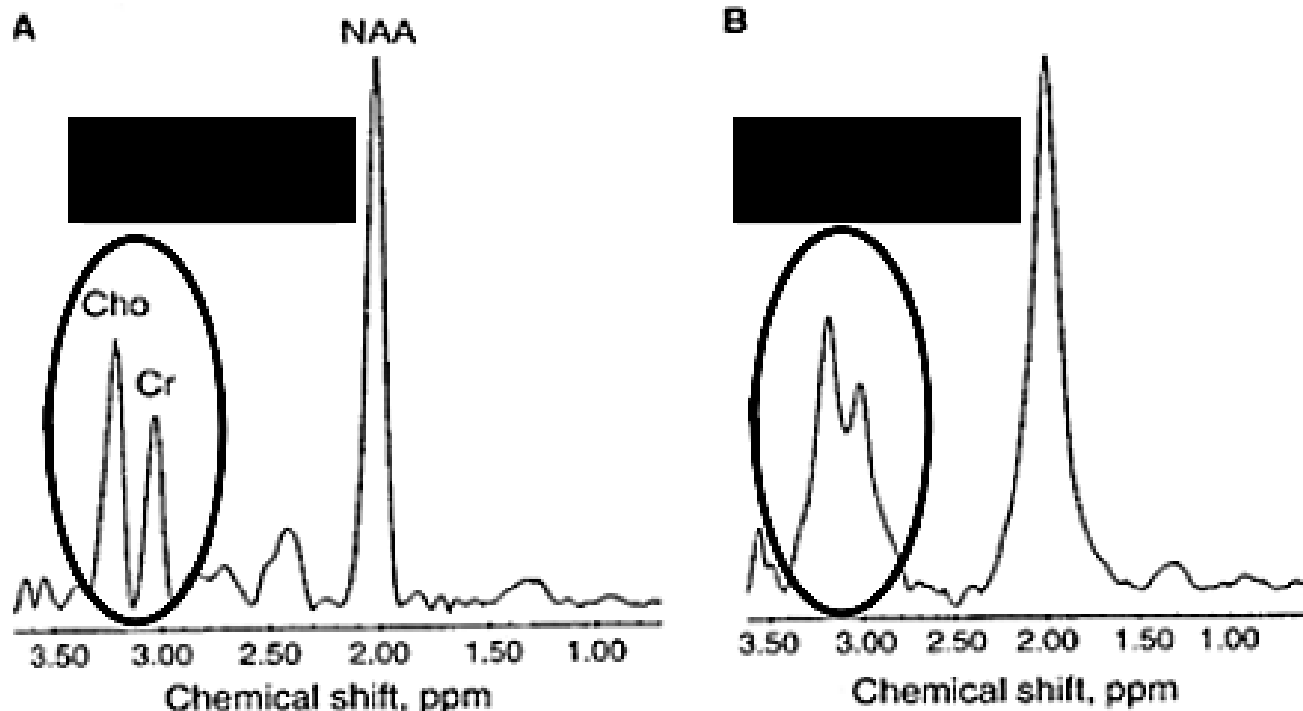
Voxel size of MRS

- MRS (6~15 min) in the brain is generally performed in conjunction with MRI.
- For single voxel techniques, a volume of 8 cc (2x2x2 cm³) is generally recommended at 1.5T.
- **Peak height is generally proportional to field strength**
 - a smaller voxel can be used at 3 T, reducing partial volume averaging.



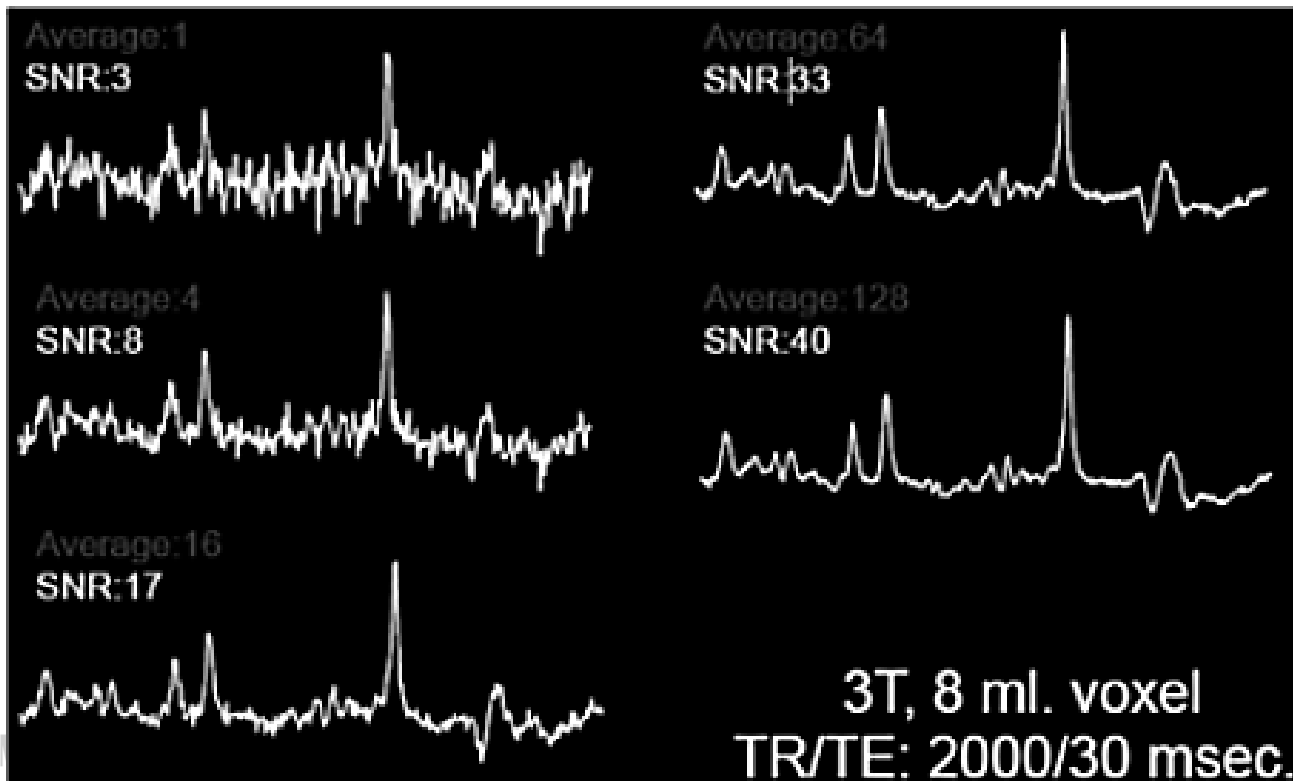
Shimming Requirement for MRS

- Shimming requirement for MRI is usually **less than 5 ppm**.
- For MRS, shimming results in improving the uniformity from 1 ppm in the main magnetic field to 0.1 ppm inside the voxel.



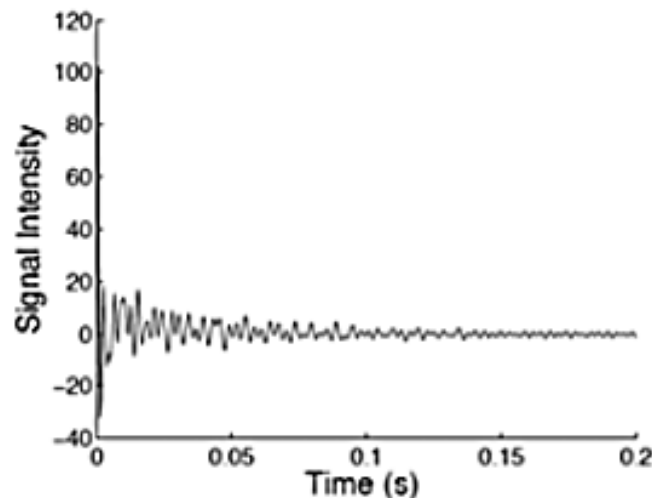
Average & SNR

- Another option to increase SNR is to increase the average (**NEX**)
- Typically, **64-128 averages** are demanded to acquire sufficient SNR for short TE.

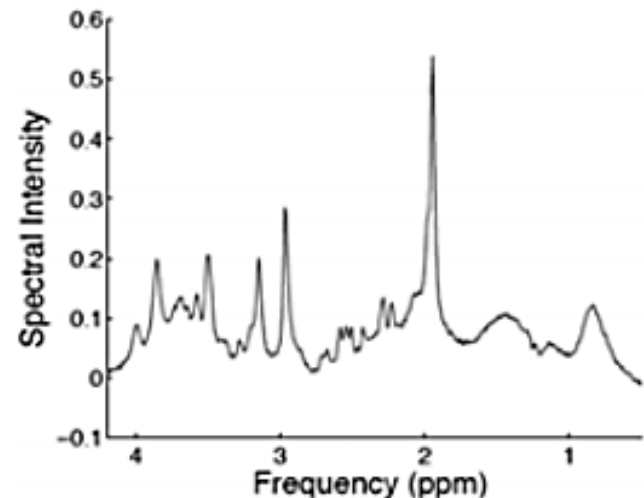


Fourier Transform

- In the simple MRS experiment, **no frequency–encoding gradients** are applied during the readout for spatial encoding.
- The signal does not contain spatial information, just information of **the different resonance frequencies** within the sample.



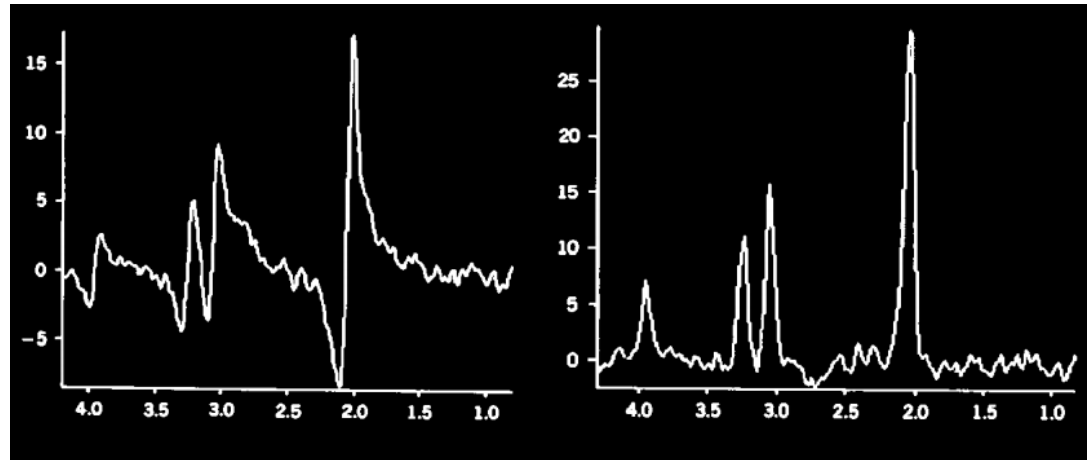
Fourier
Transform
→



30

Post Processing

- FID signal processing
 - Water suppression (removing the 4.7 ppm signal)
 - Zero filling (Increase frequency resolution)
 - Apodization (Noise filter)
- Fourier Transform
- Spectrum processing
 - Phase Correction
 - Baseline Correction

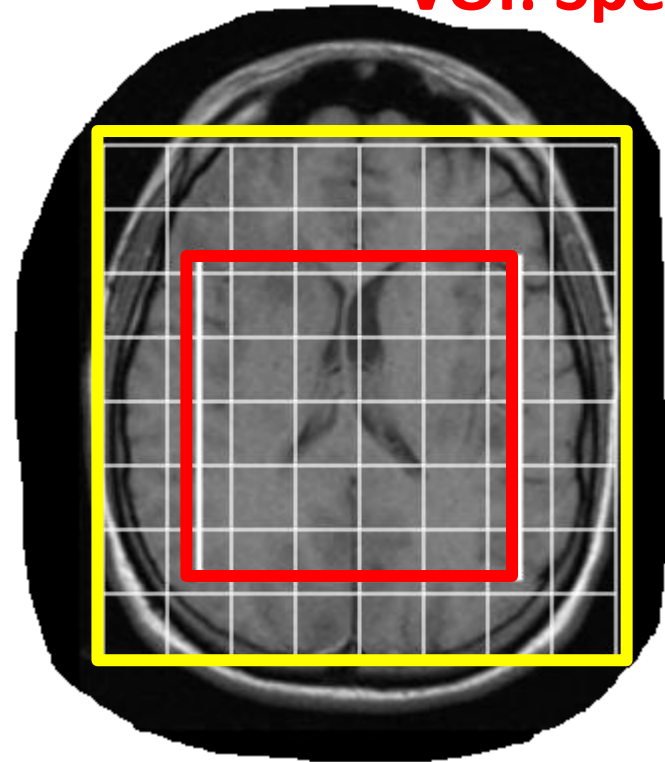
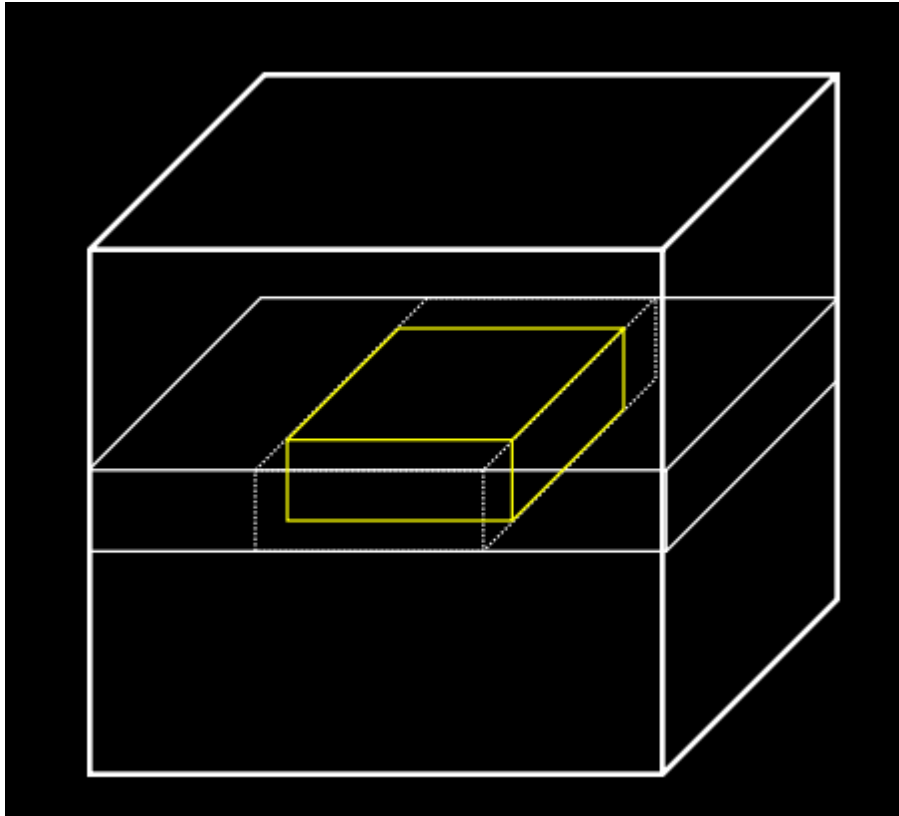


Spectroscopic Imaging (SI)

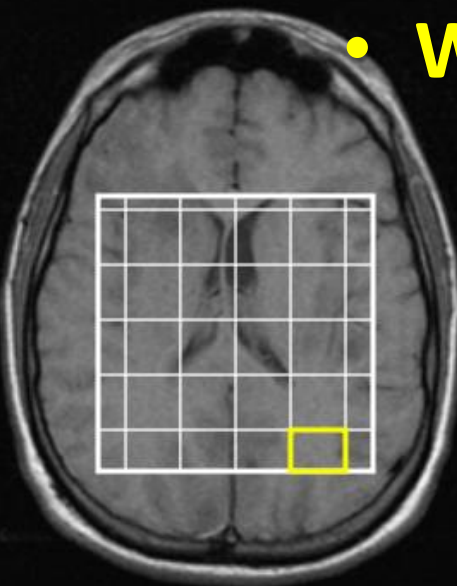
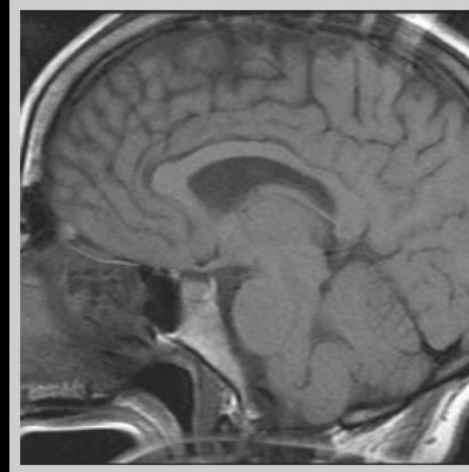
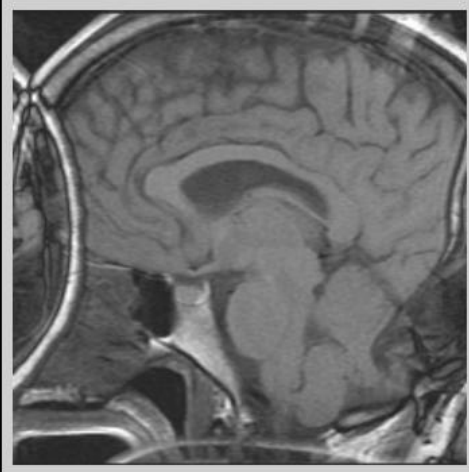
- Multi-Voxel
- Chemical Shift Imaging (CSI)

FOV: SNR \uparrow

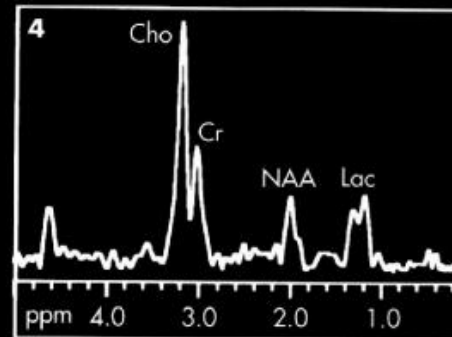
VOI: Spectrum



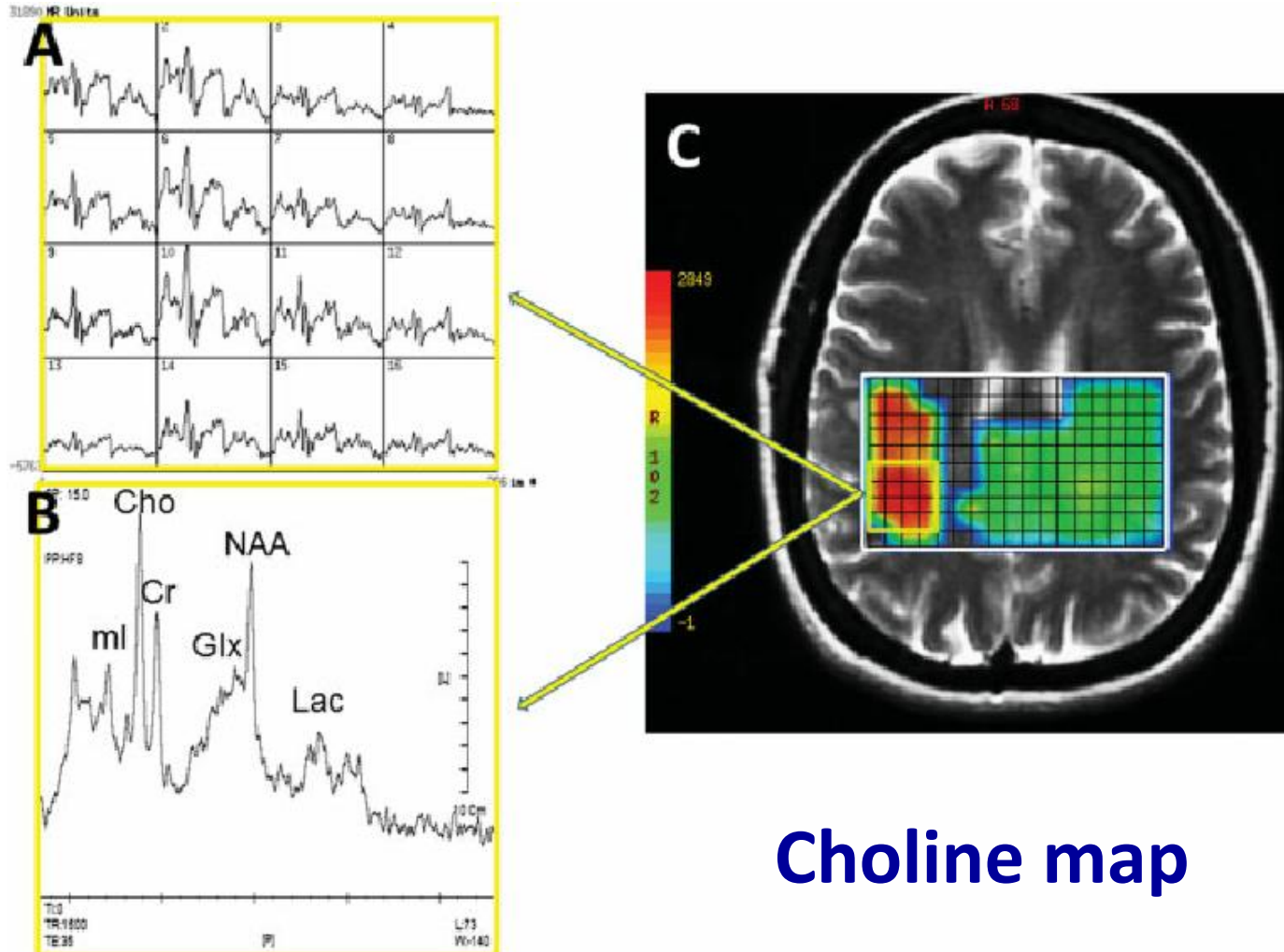
Aliasing Artifact



• Whose lactate???



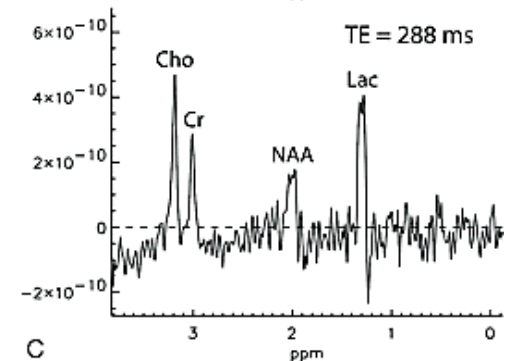
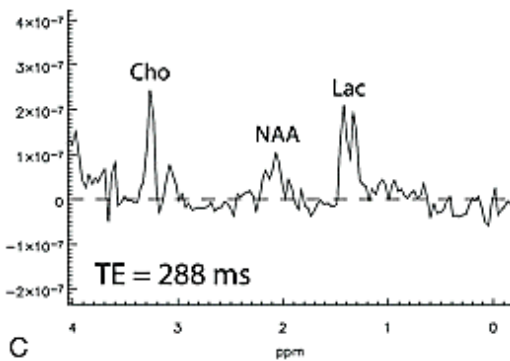
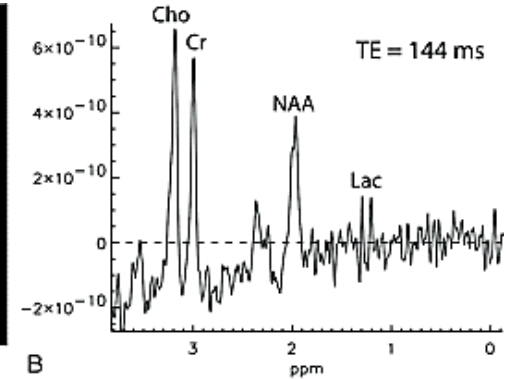
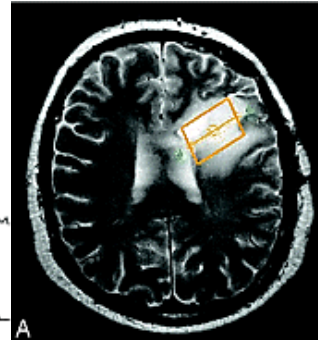
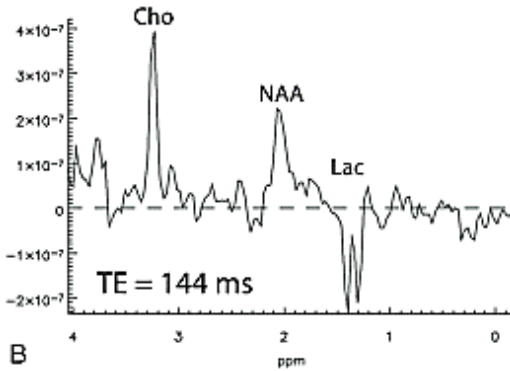
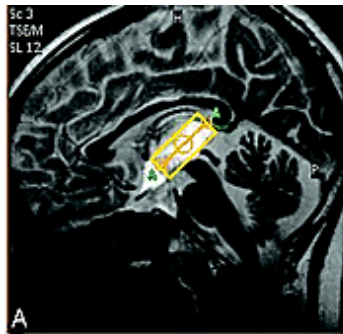
Metabolite Concentration Map



Choline map

Special Topic: Lactate

1. Inversion peak?
2. Nearly invisible at 3T short TE condition?



1.5T

3T



Special Topic: Lactate

1. Opposite phase with NAA
2. Anomalous J-modulation

