

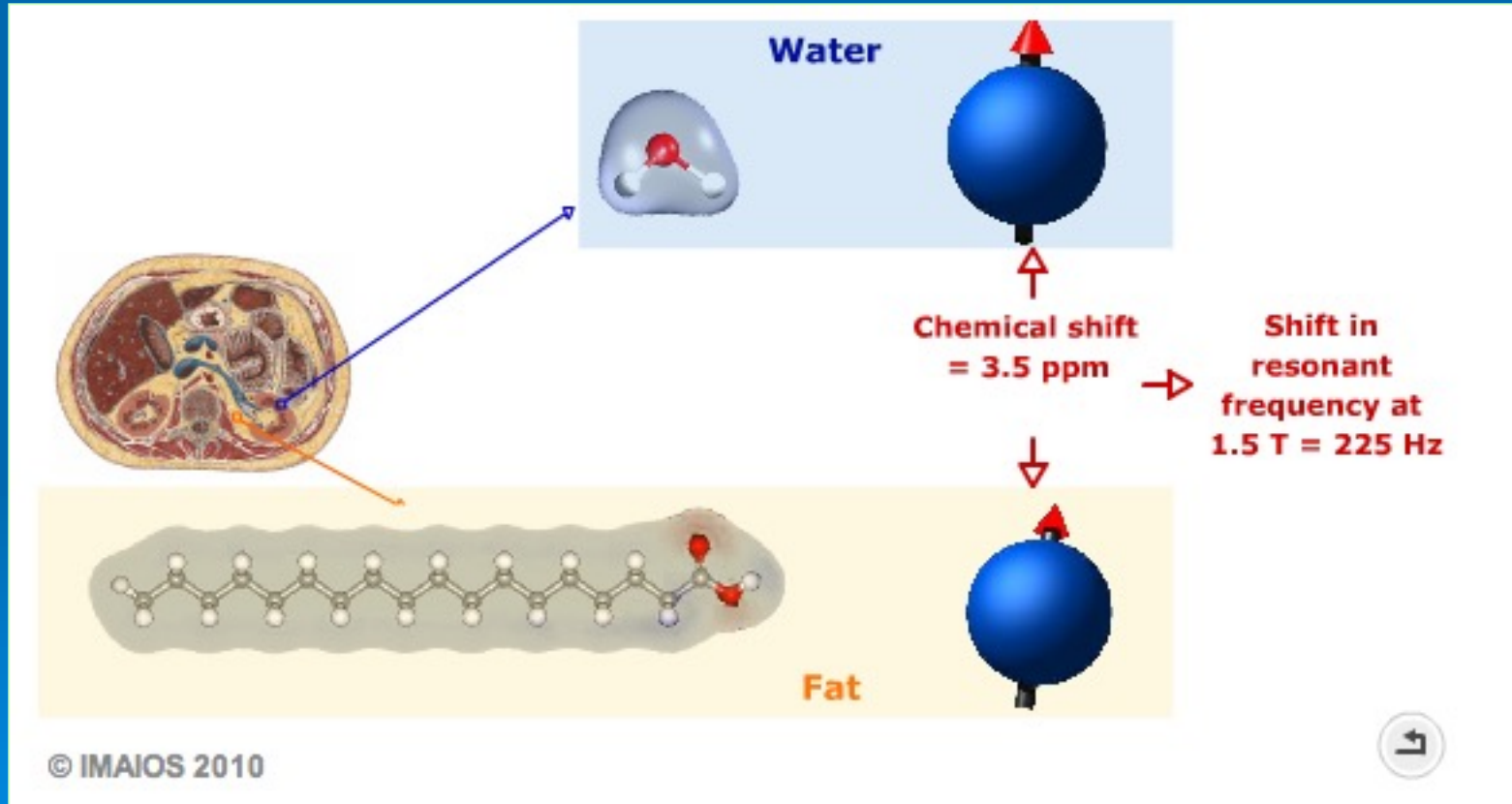
# MRS

# Magnetic Resonance Spectroscopy

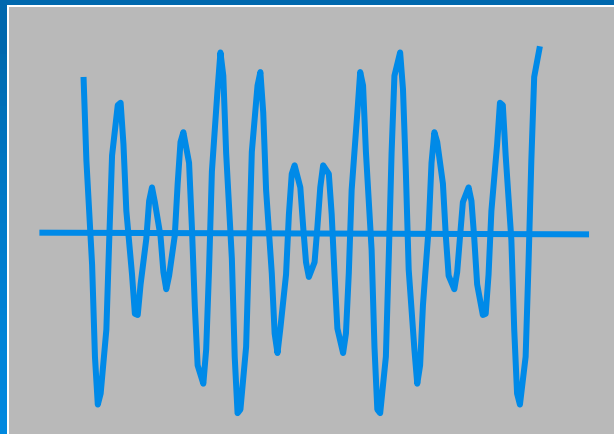
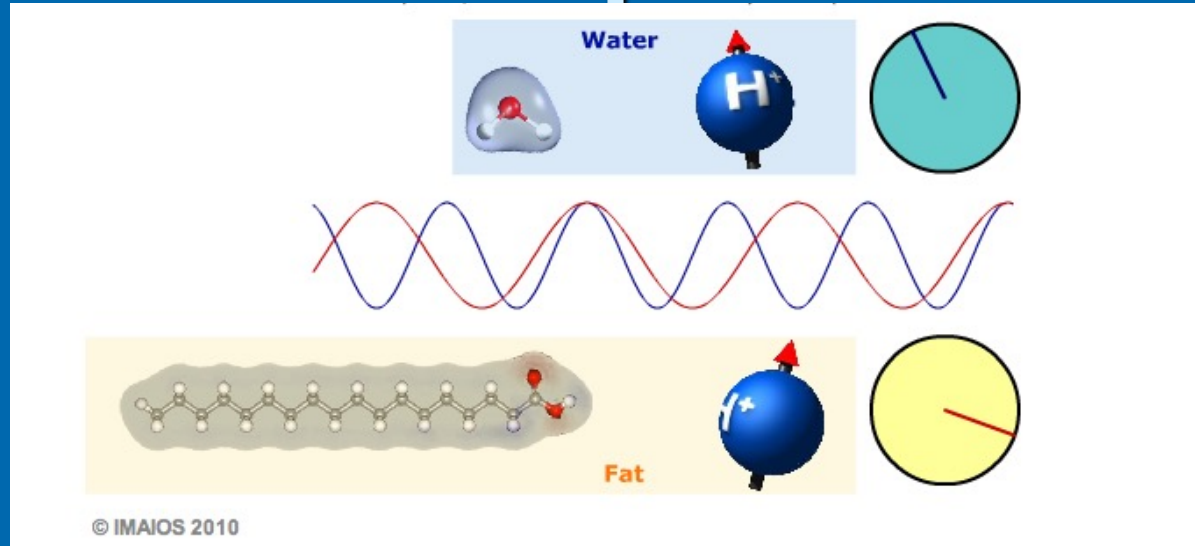
In vivo biochemistry

The background of the slide is a solid blue color. In the lower right quadrant, there are several sets of concentric, light blue circles that resemble ripples on water. These circles are of varying sizes and are arranged in a way that suggests movement or a field of activity.

# Chemical shift

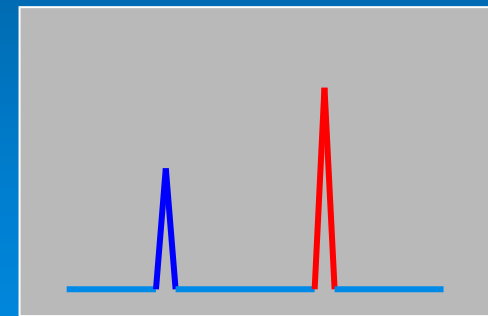


# Chemical shift and MR spectrum



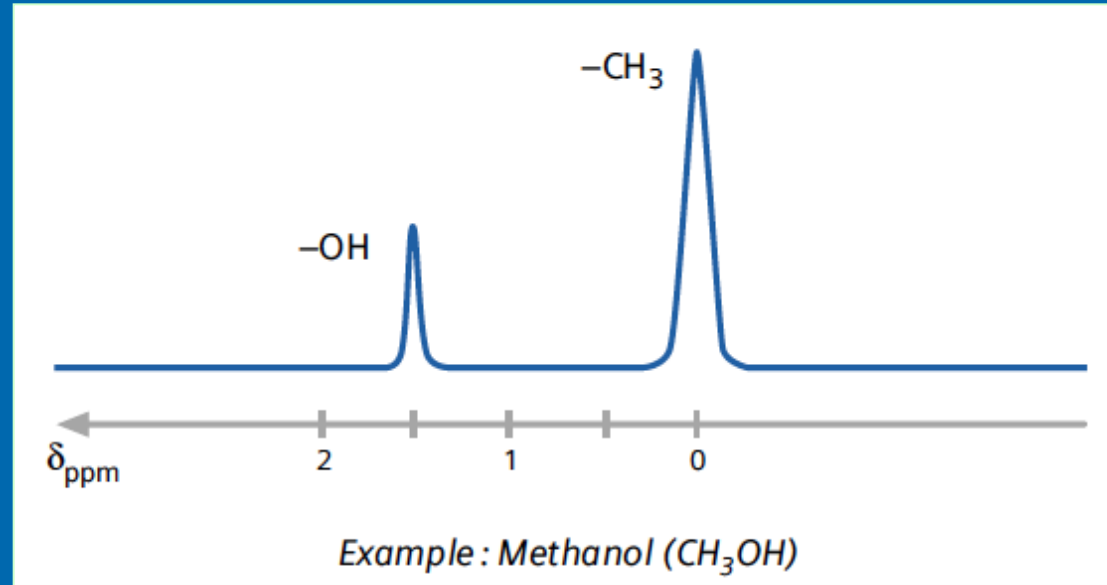
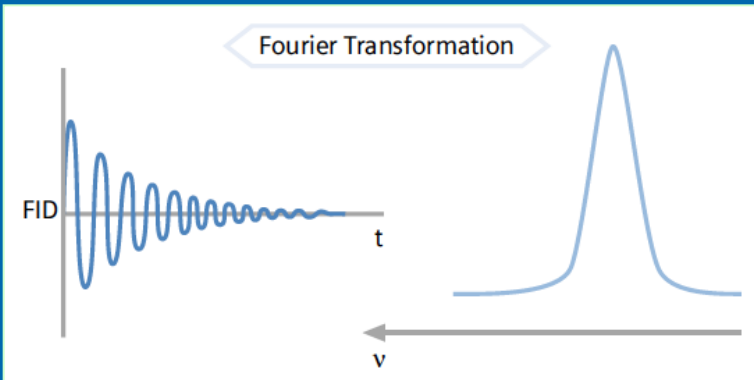
Received Signal

Fourier Transform



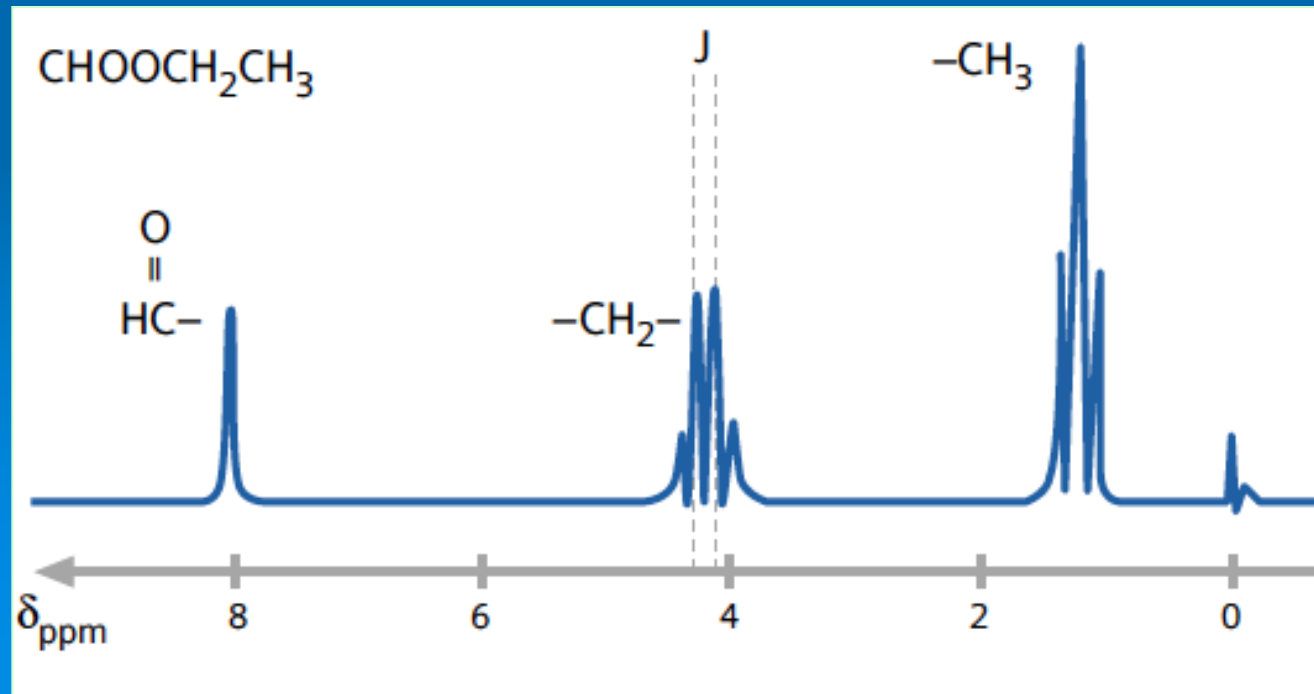
Image

# $^1\text{H}$ NMR spectra

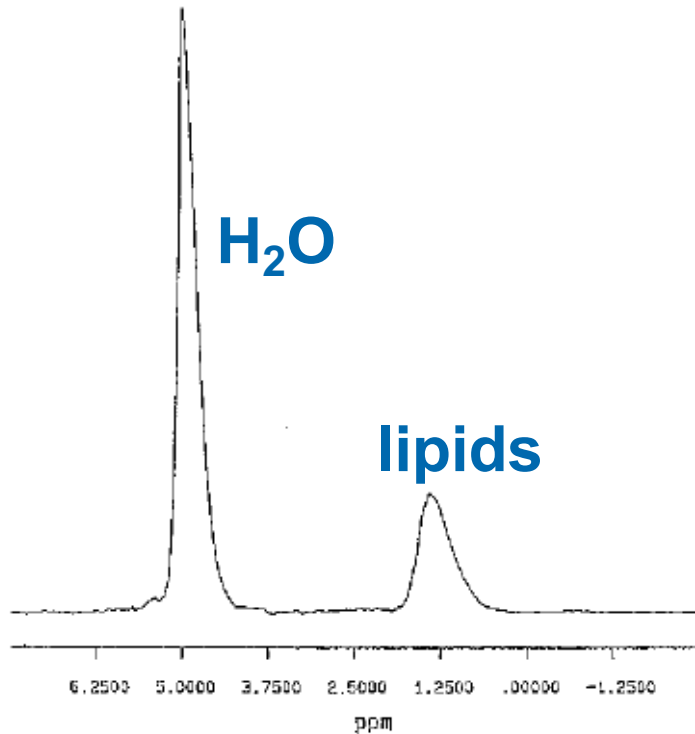


Example: Methanol ( $\text{CH}_3\text{OH}$ )

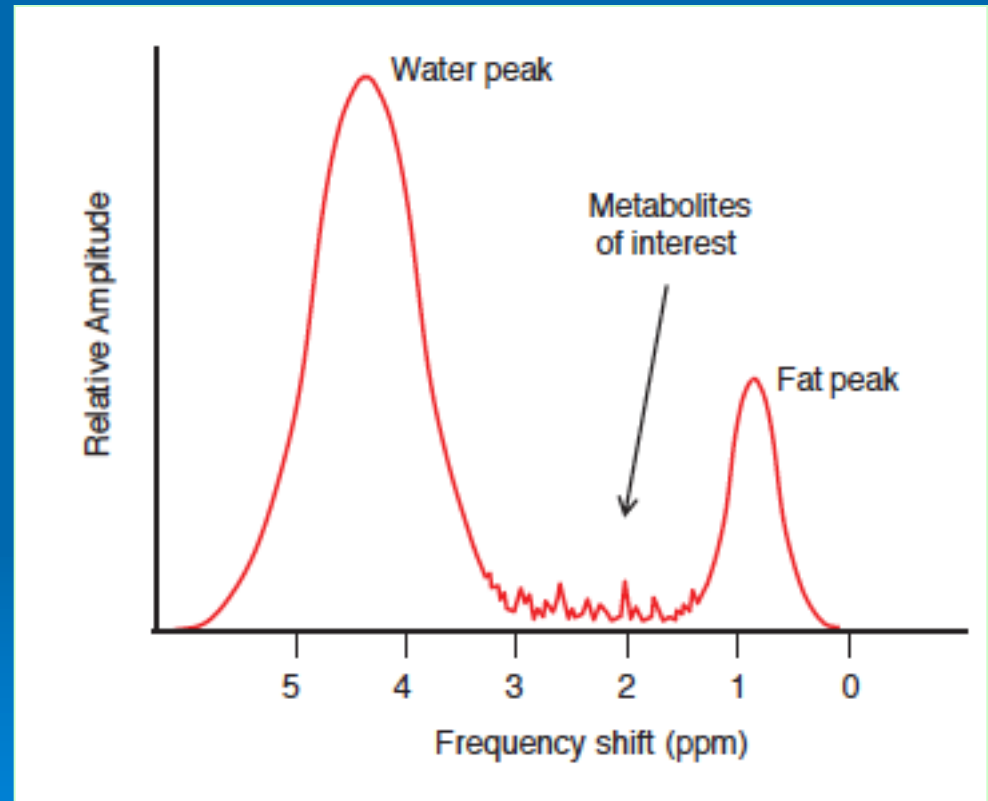
- ✓ Peaks pattern is footprint of molecules
- ✓ Peak area is proportional to the number of  $^1\text{H}$  nuclei



# Proton spectroscopy

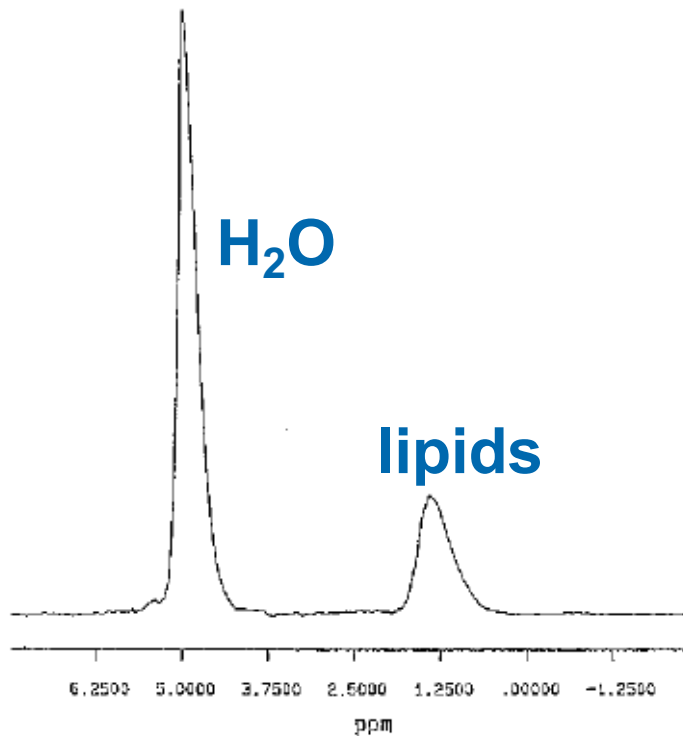


**Figure 1.** Typical in vivo H-1 spectrum from a steatotic liver (TR = 3 seconds, TE = 24 and 50 msec, four signals acquired, 262 cm<sup>3</sup> volume of interest). No signal filtering before Fourier transformation and no baseline correction were applied.

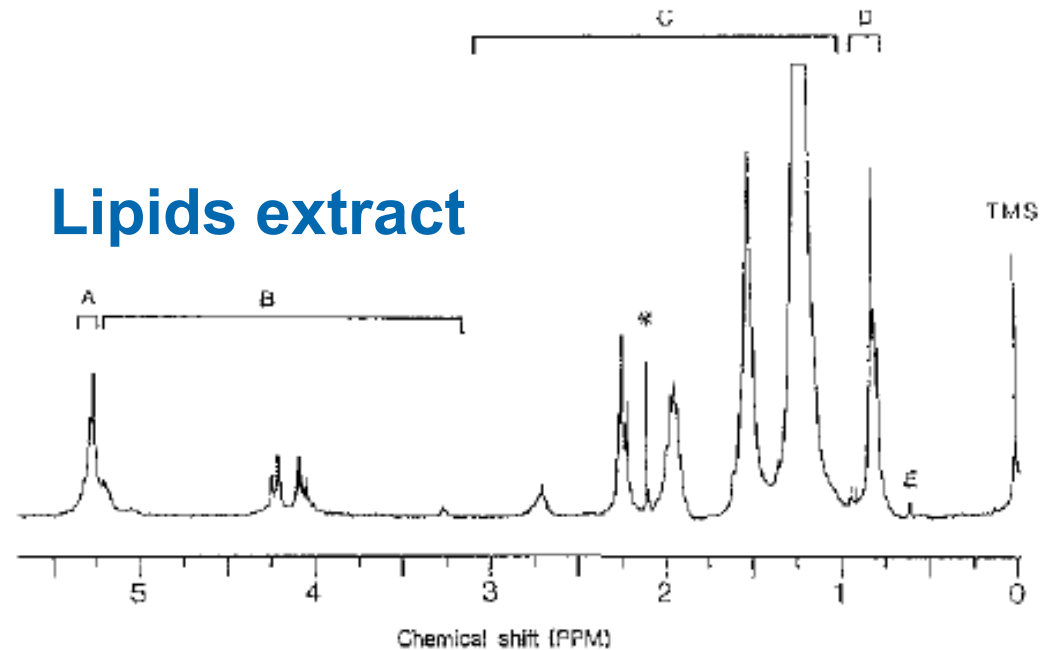


➤ The water signal has to be suppressed

# Proton spectroscopy



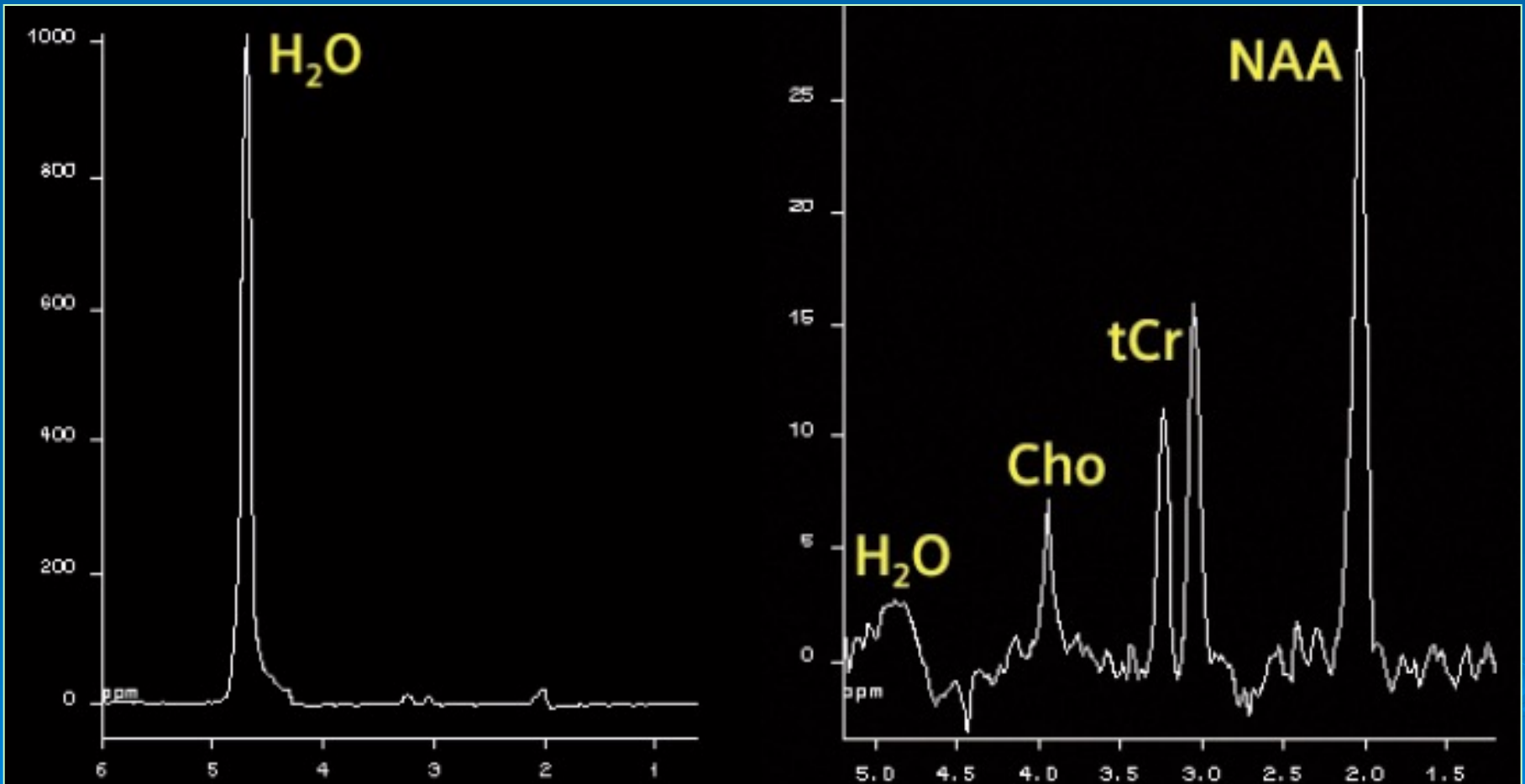
**Figure 1.** Typical in vivo H-1 spectrum from a steatotic liver (TR = 3 seconds, TE = 24 and 50 msec, four signals acquired, 262 cm<sup>3</sup> volume of interest). No signal filtering before Fourier transformation and no baseline correction were applied.



**Figure 2.** H-1 300-MHz spectrum of lipid extract obtained from a steatotic liver specimen. The major peaks assignable to protons in different positions on lipid molecules are (A) double bonds, (B) protons belonging to di- or triacylated glycerol and to the phosphocholine and phosphoethanolamine components of phospholipids, (C) methylene groups, (D) methyl groups, and (E) methyl signal assigned to carbon-18 of cholesterol. Acetone (\*) and tetramethylsilane (TMS) (internal standard) are also shown.

➤ The water signal has to be suppressed

# Water suppression



**Spectrum without and with water suppression**  
*Different scaling*

# $^1\text{H}$ MRS metabolites

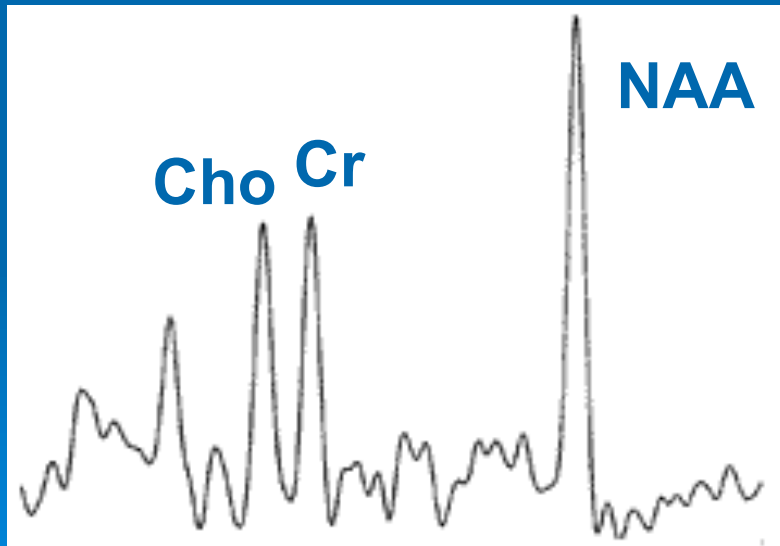
ABBREVIATION	METABOLITE	SHIFT (PPM)	PROPERTIES/SIGNIFICANCE IN THE BRAIN
Cho	Phosphocholine	3.22	Membrane turnover, cell proliferation
Cr	Creatine	3.02 and 3.93	Temporary store for energy-rich phosphates
NAA	<i>N</i> -acetyl-L-aspartate	2.01	Presence of intact glioneuronal structures
Lactate		1.33 (inverted)	Anaerobic glycolysis
Lipids	Free fatty acids	1.2–1.4	Necrosis



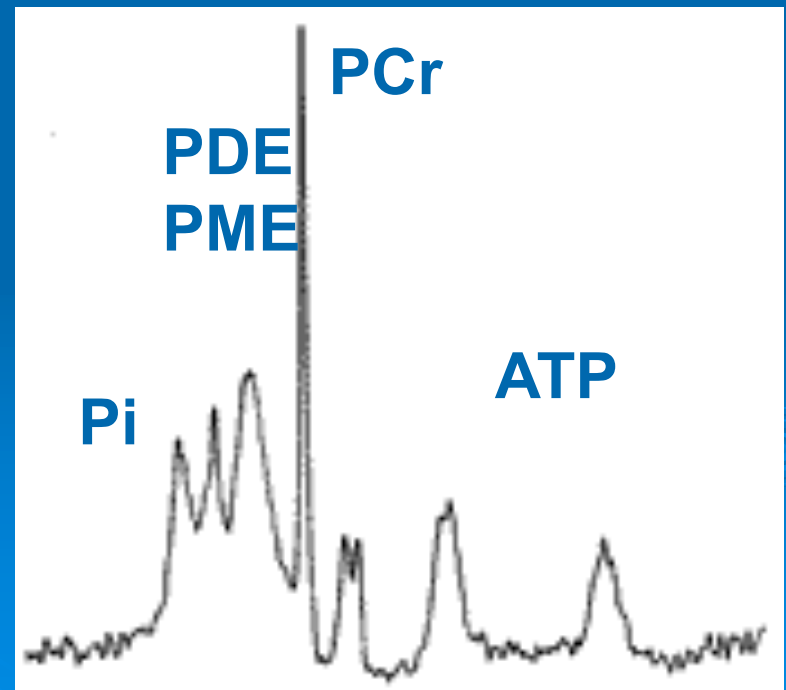
# Brain spectroscopy

- The  $^1\text{H}$  (or  $^{31}\text{P}$ ) nuclei in different molecules have slightly different resonance frequencies
- Each peak is related to a molecule (metabolite)

*In vivo*  $^1\text{H}$  spectrum

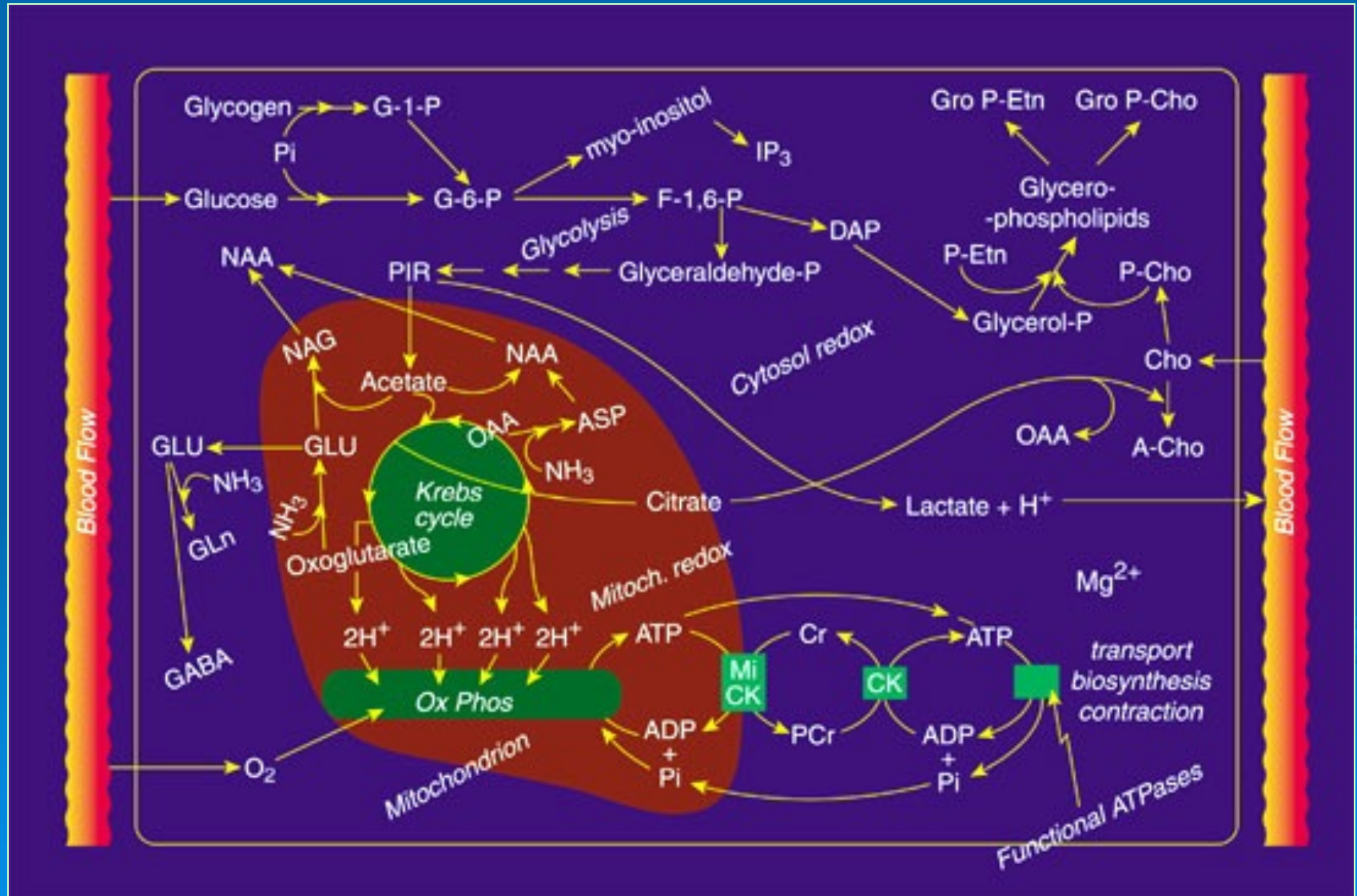


*In vivo*  $^{31}\text{P}$  spectrum

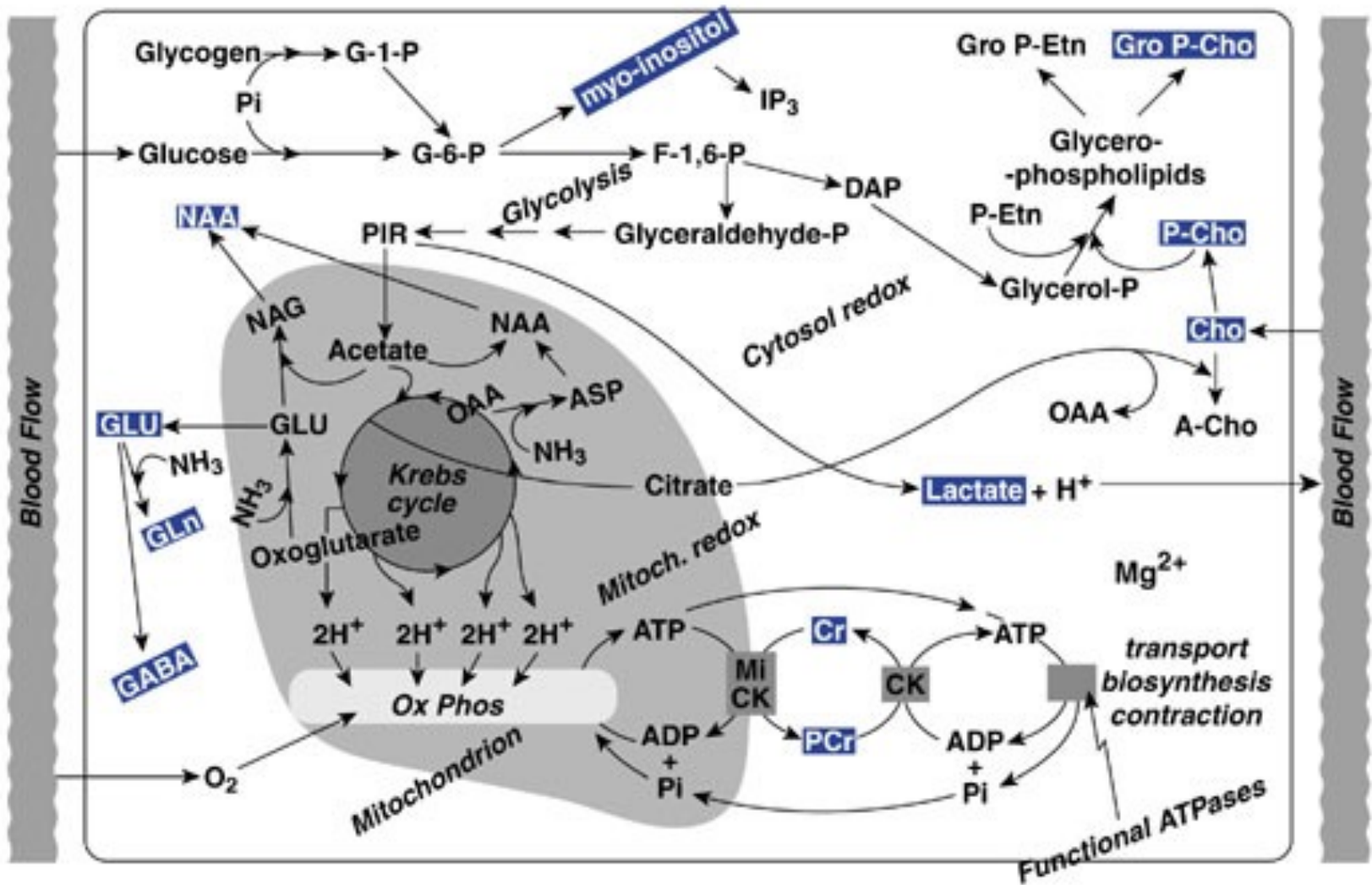


- Occipital cortex

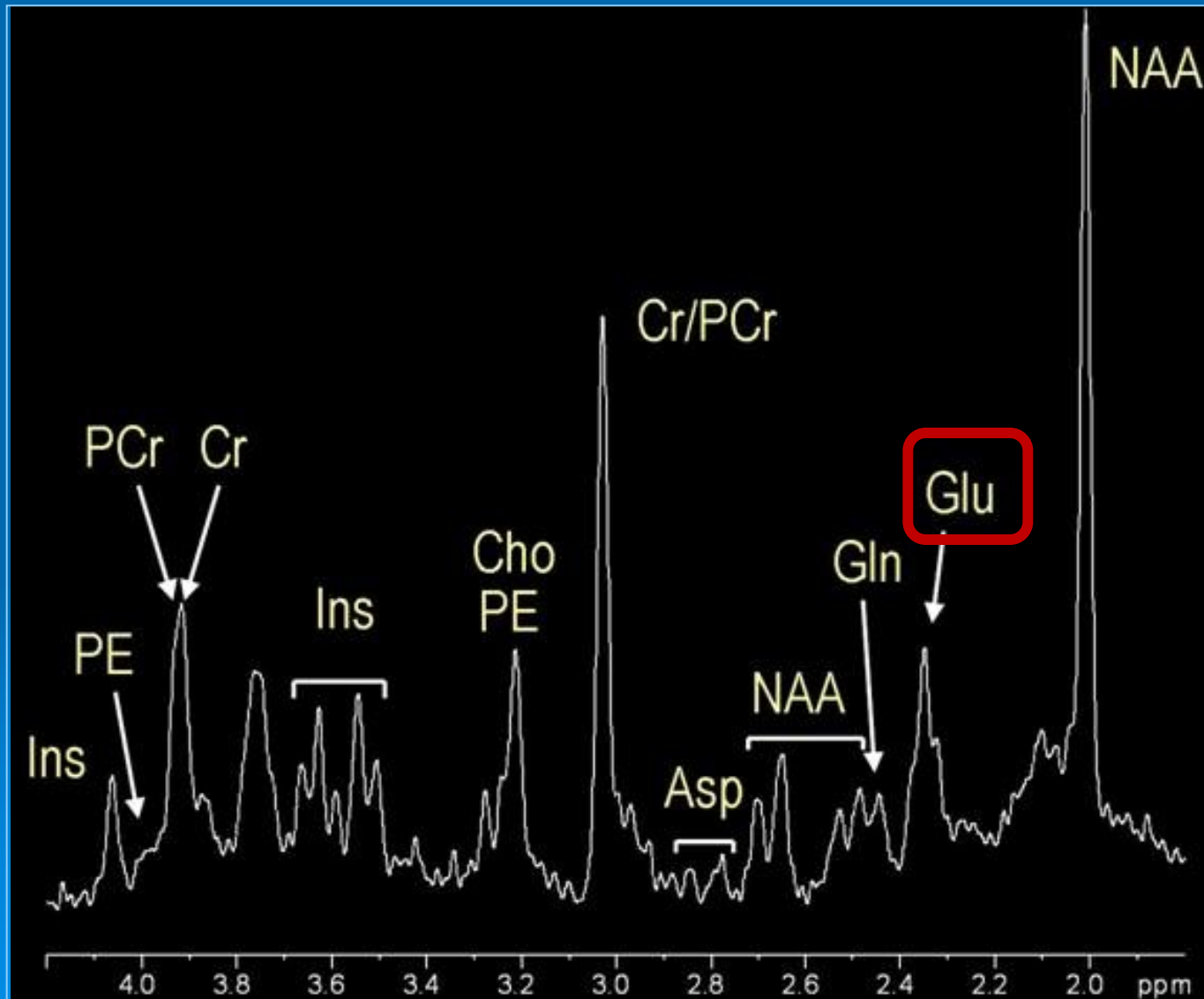
# *In vivo* biochemistry



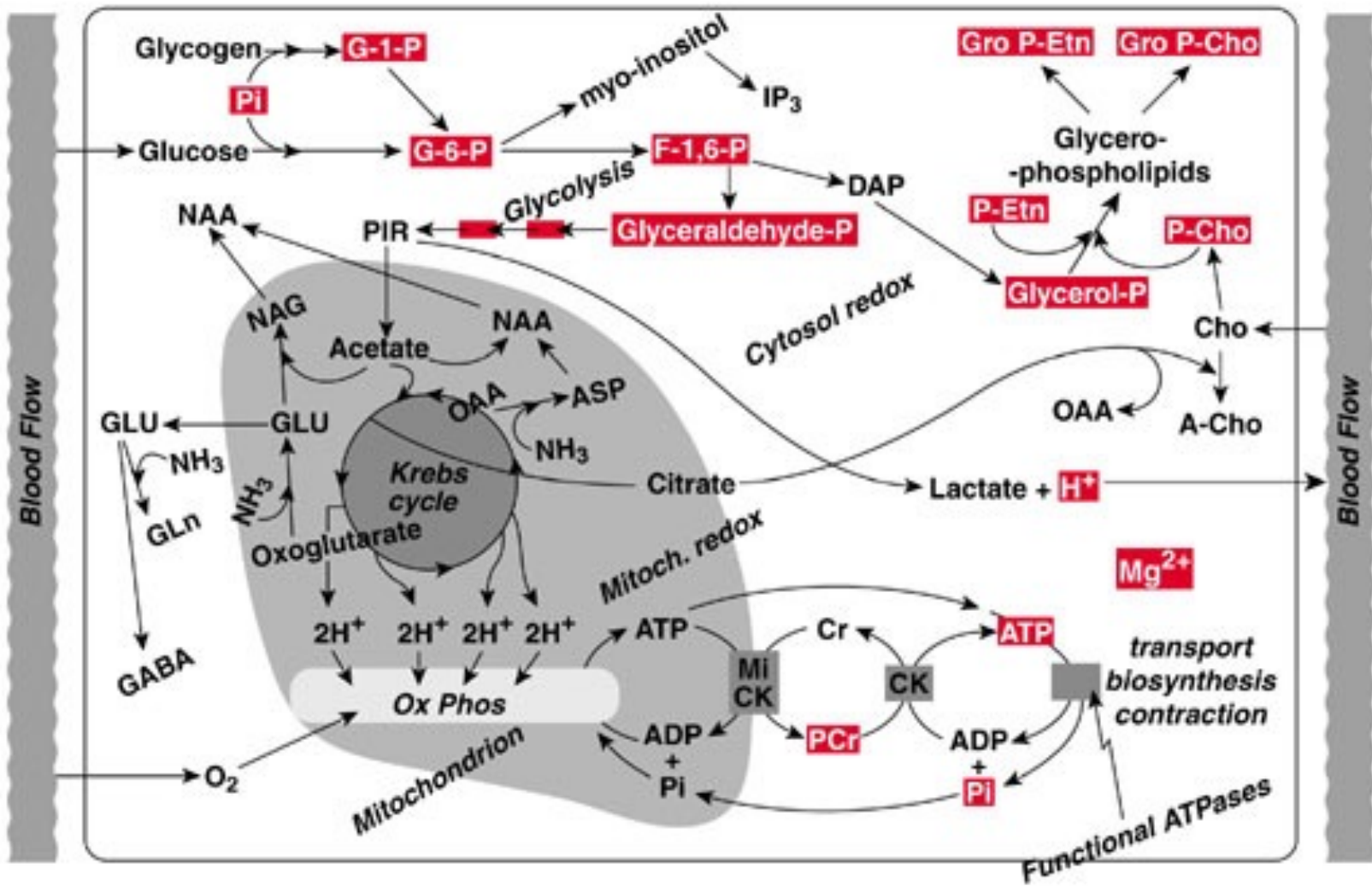
# <sup>1</sup>H spectroscopy



# $^1\text{H}$ spectroscopy

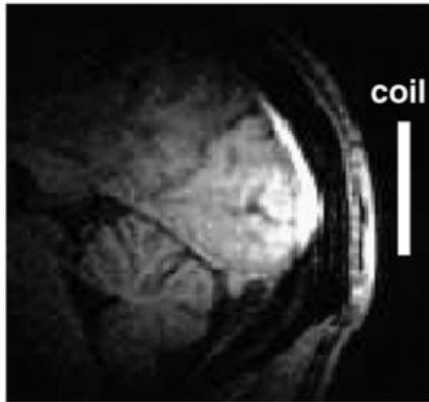


# <sup>31</sup>P spectroscopy



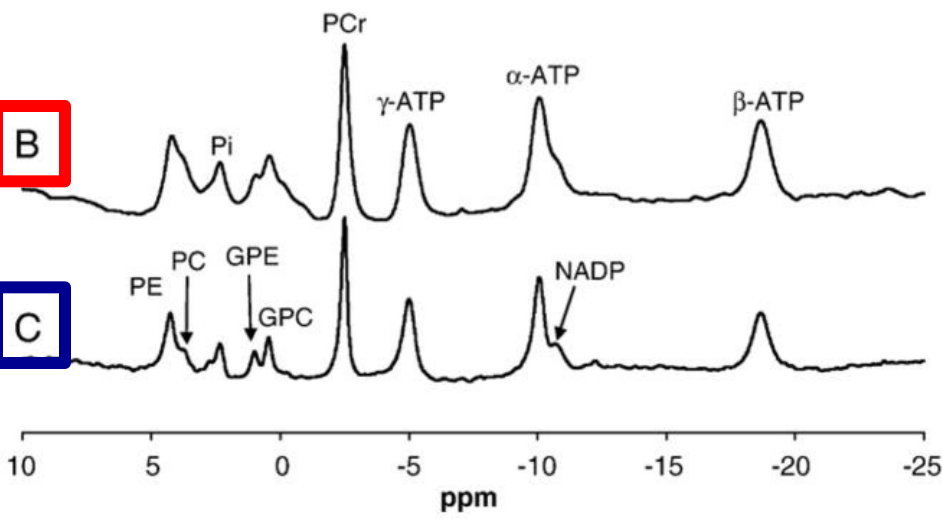
# $^{31}\text{P}$ MRS

A



B

C



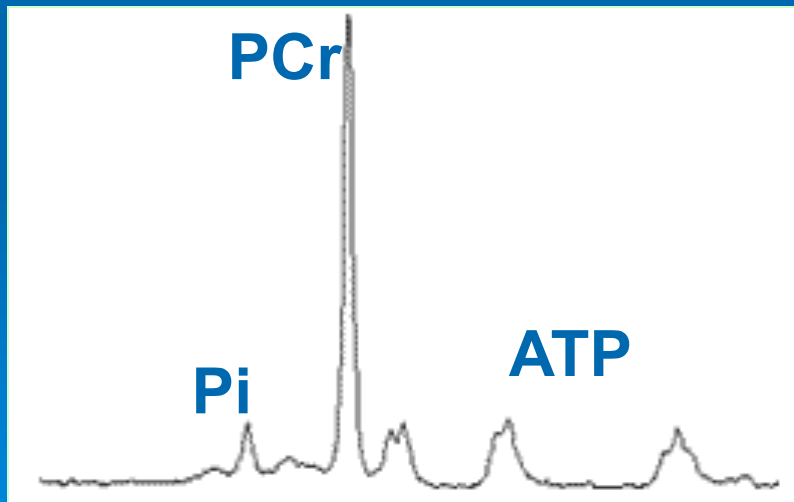
In vivo  $^{31}\text{P}$  spectra acquired from the human occipital lobe at (B) 4 T and (C) 7 T:

- PE phosphoethanolamine
- PC phosphocholine
- Pi inorganic phosphate
- GPE glycerophosphoethanolamine
- GPC, glycerophosphocholine
- PCr phosphocreatine
- ATP adenosine triphosphate
- NADP nicotinamide adenine dinucleotide phosphate

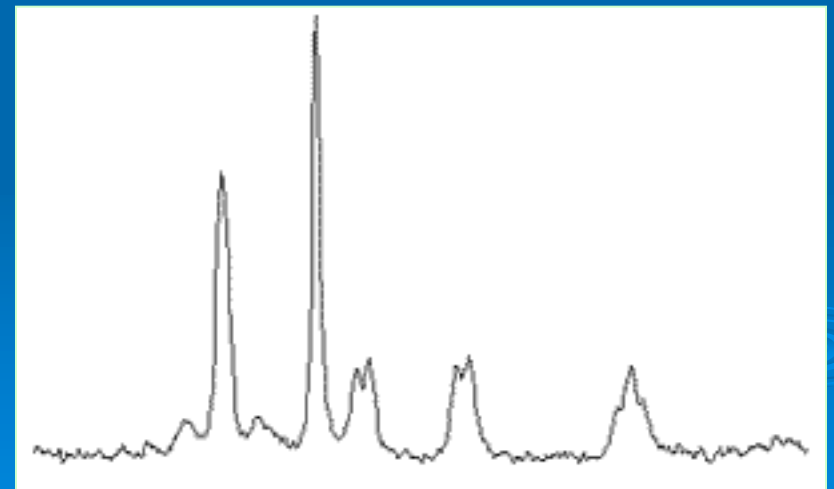
# muscle $^{31}\text{P}$ MRS

- Muscoli gastrocnemi normal subject

rest

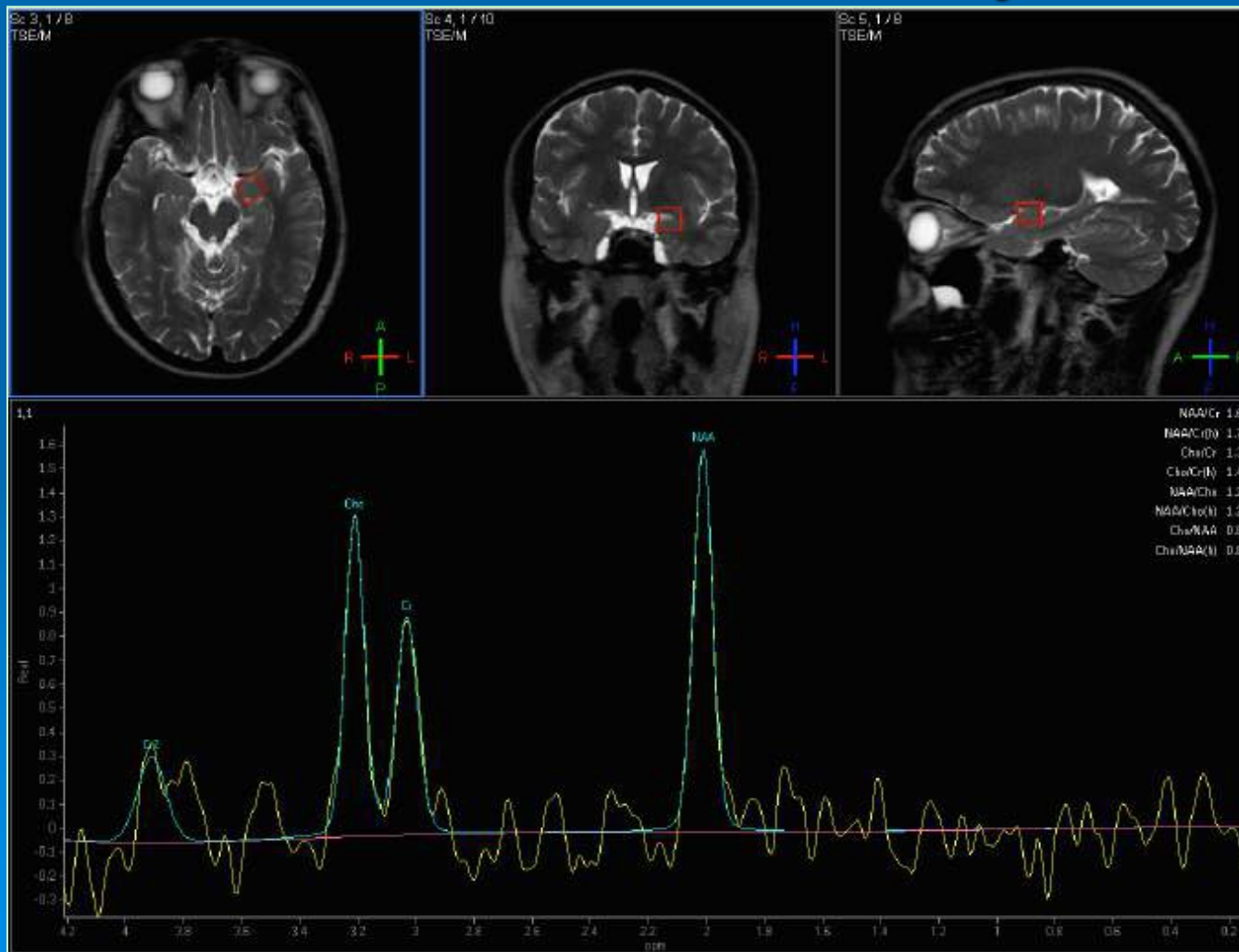


exercise



# Signal localization

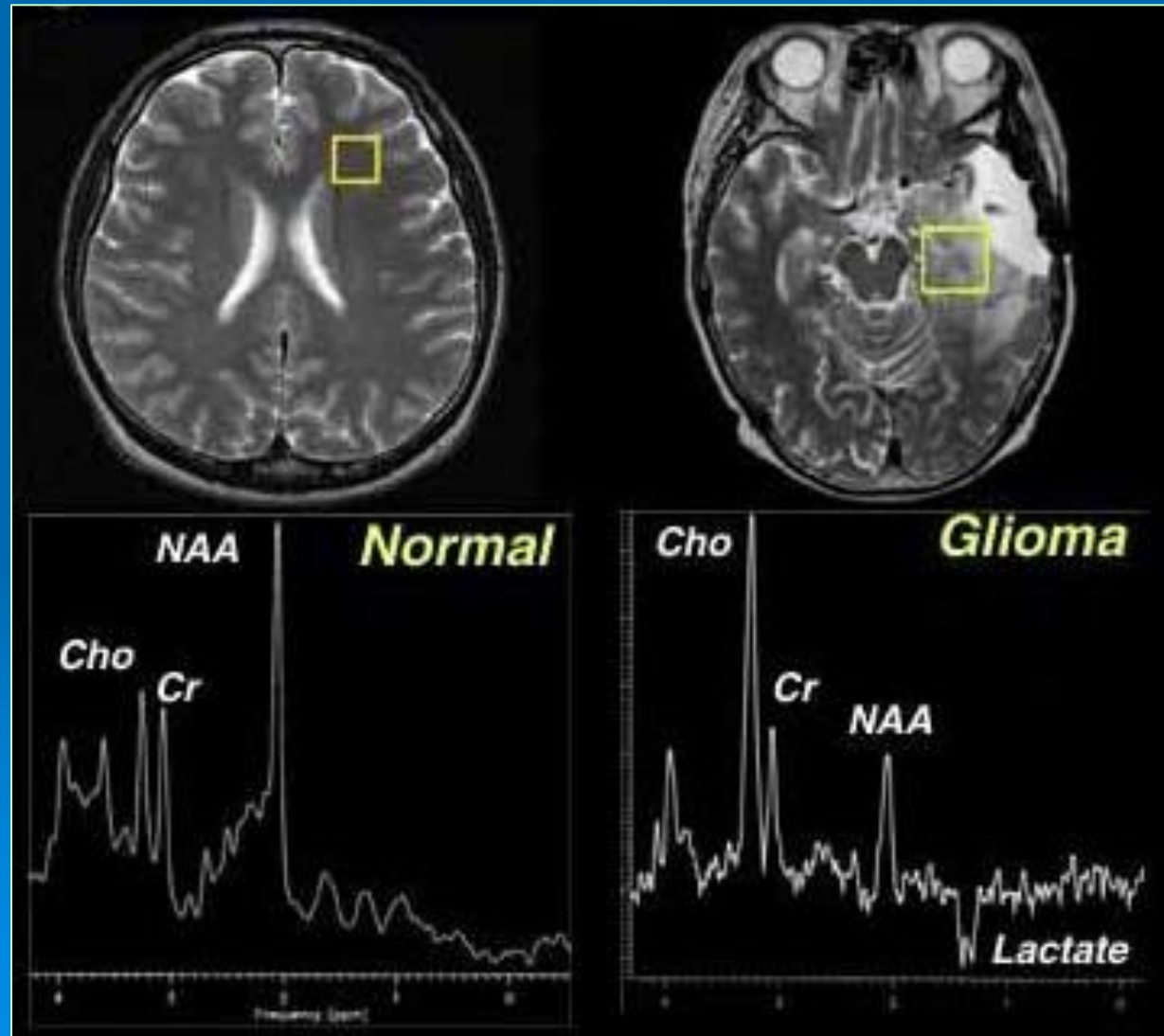
In *in vivo* MRS the signal localization is mandatory





# Signal localization

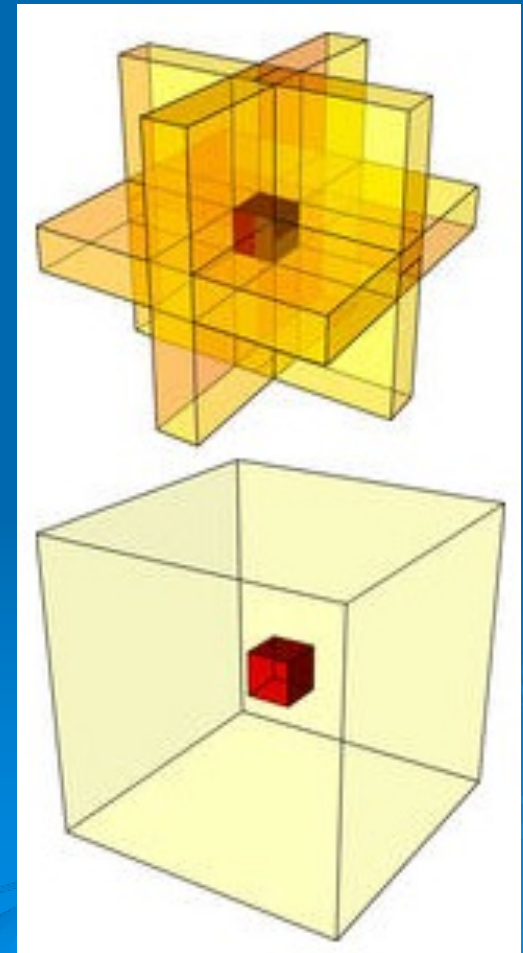
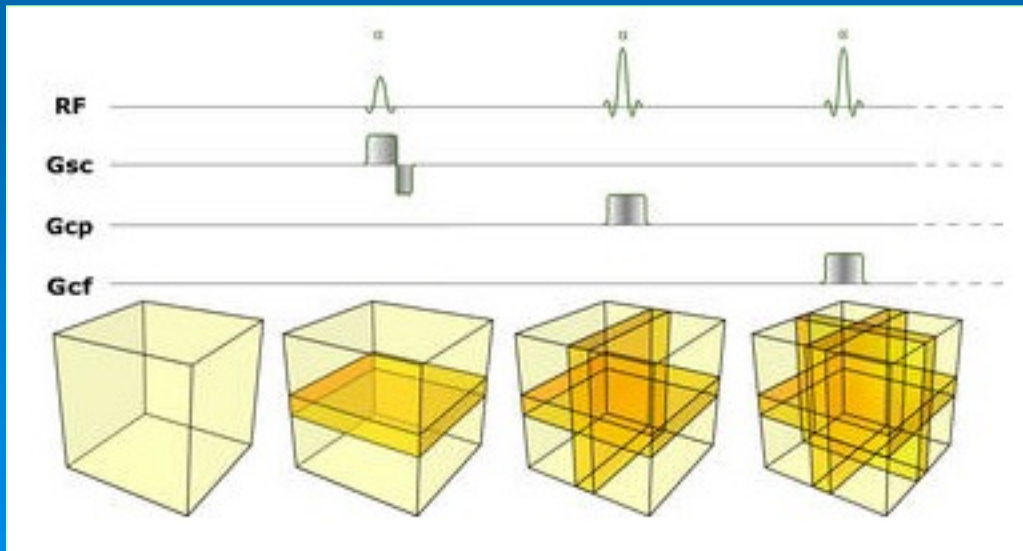
- In in vivo MRS the signal localization is mandatory



<sup>1</sup>H MRS

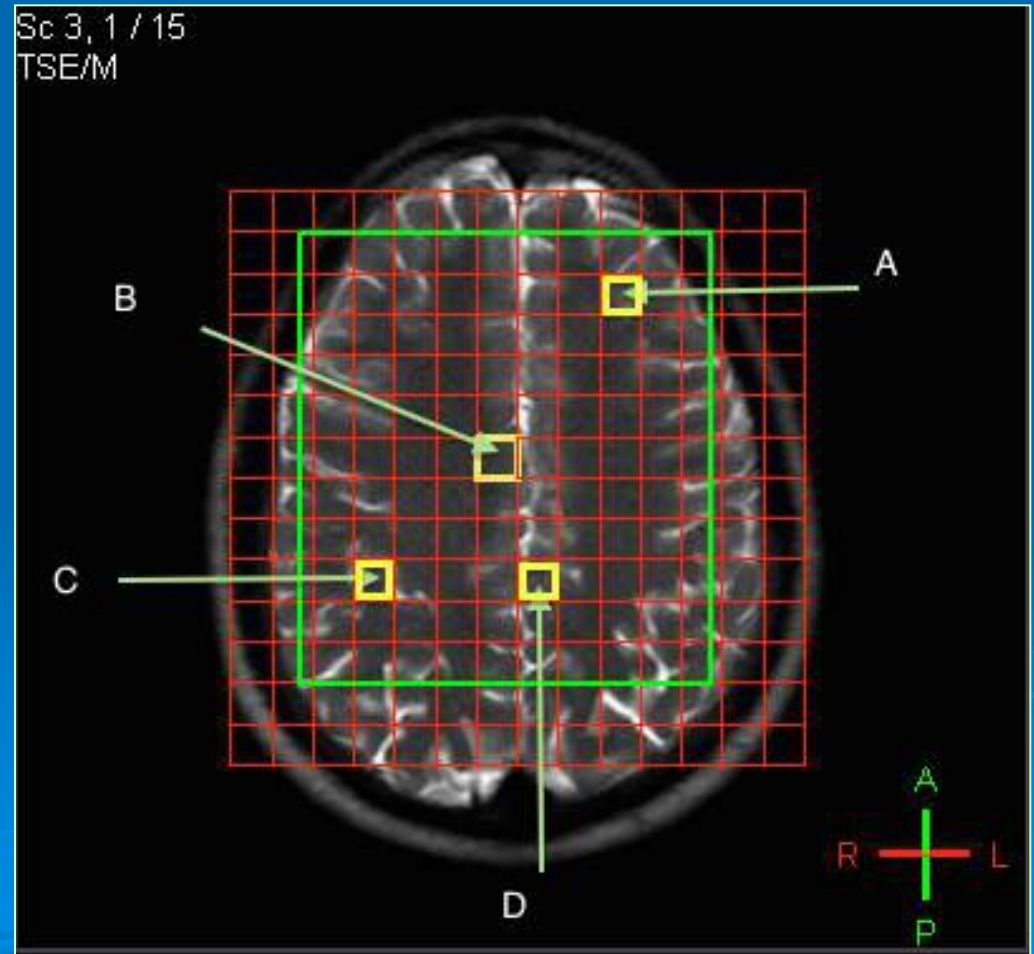
# Volume selection

- ✓ The localization of the MRS signal is essential for in vivo application
- ✓ Gradients are used localized spectroscopy
- ✓ The simplest localization technique is the use of the surface coil



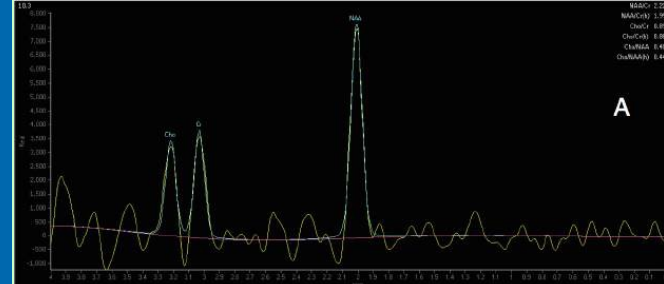
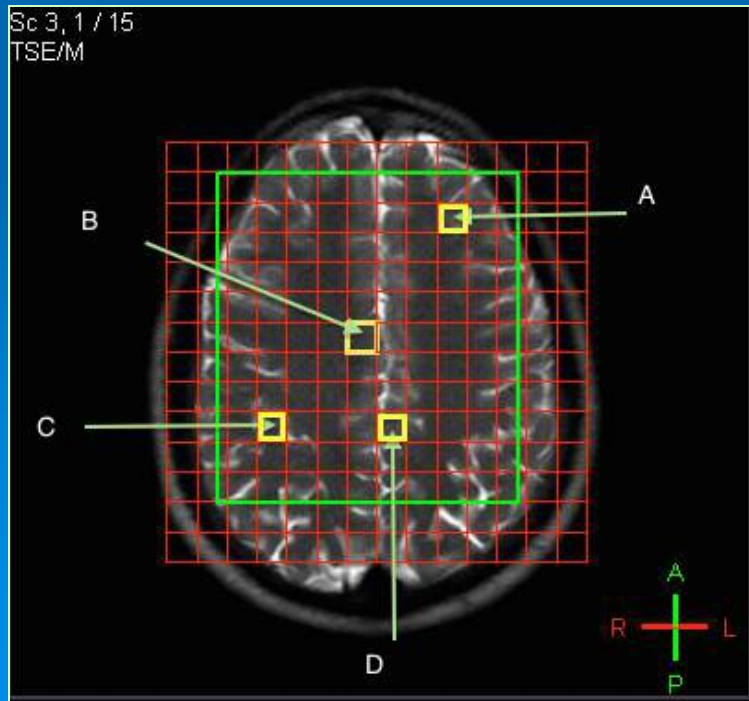
# Spectroscopic imaging (Chemical shift imaging CSI)

- ✓ CSI is an acquisition sequence that allows the acquisition of a spectrum per each voxel
- ✓ The acquisition time is large (> 10 minutes)



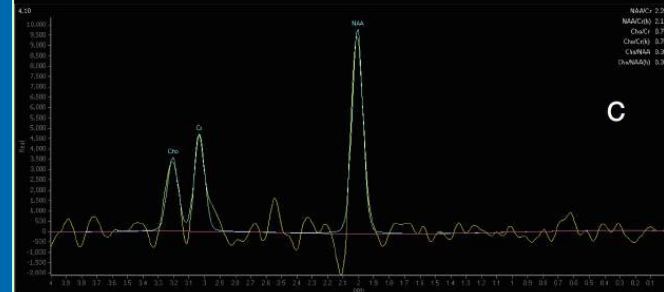
# Spectroscopic imaging

- ✓ CSI is an acquisition sequence that allows the acquisition of a spectrum per each voxel
- ✓ The acquisition time is large
  - > 10 minutes



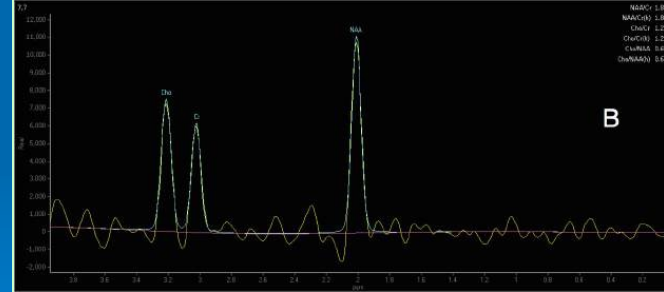
Spectro Results

Metab	Position	SNR	W088	Height	FWHM	Area	AreaCr	Metab	Position	SNR	W088	Height	FWHM	Area	AreaCr
NAA	2.007	3213	0.884	77266	1.996	7291	3.228	Cr	3.001	1634	0.076	3883	1.000	3258	1.000



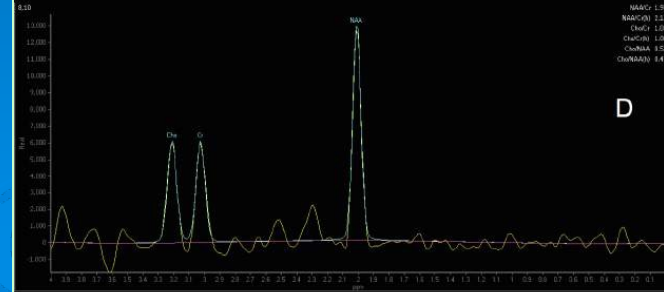
Spectro Results

Metab	Position	SNR	W088	Height	FWHM	Area	AreaCr	Metab	Position	SNR	W088	Height	FWHM	Area	AreaCr
NAA	2.005	2864	0.882	68869	2.212	8177	2.991	Cr	3.002	1458	0.076	4638	1.000	4129	1.000



Spectro Results

Metab	Position	SNR	W088	Height	FWHM	Area	AreaCr	Metab	Position	SNR	W088	Height	FWHM	Area	AreaCr
NAA	2.000	4262	0.867	11557	1.796	8387	1.825	Cr	3.003	2274	0.066	6235	1.000	4919	1.000

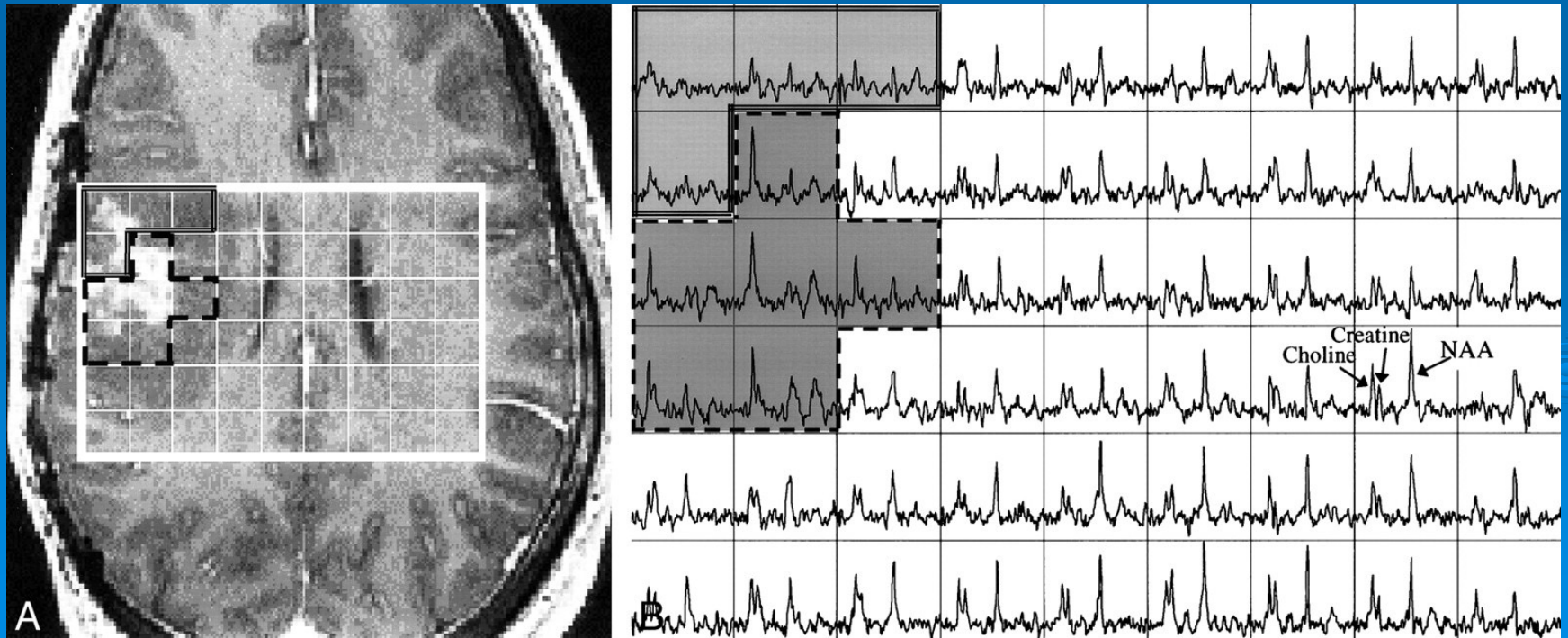


Spectro Results

Metab	Position	SNR	W088	Height	FWHM	Area	AreaCr	Metab	Position	SNR	W088	Height	FWHM	Area	AreaCr
NAA	2.000	3514	0.862	22923	2.120	8122	1.971	Cr	3.004	713	0.067	4983	1.000	4526	1.000

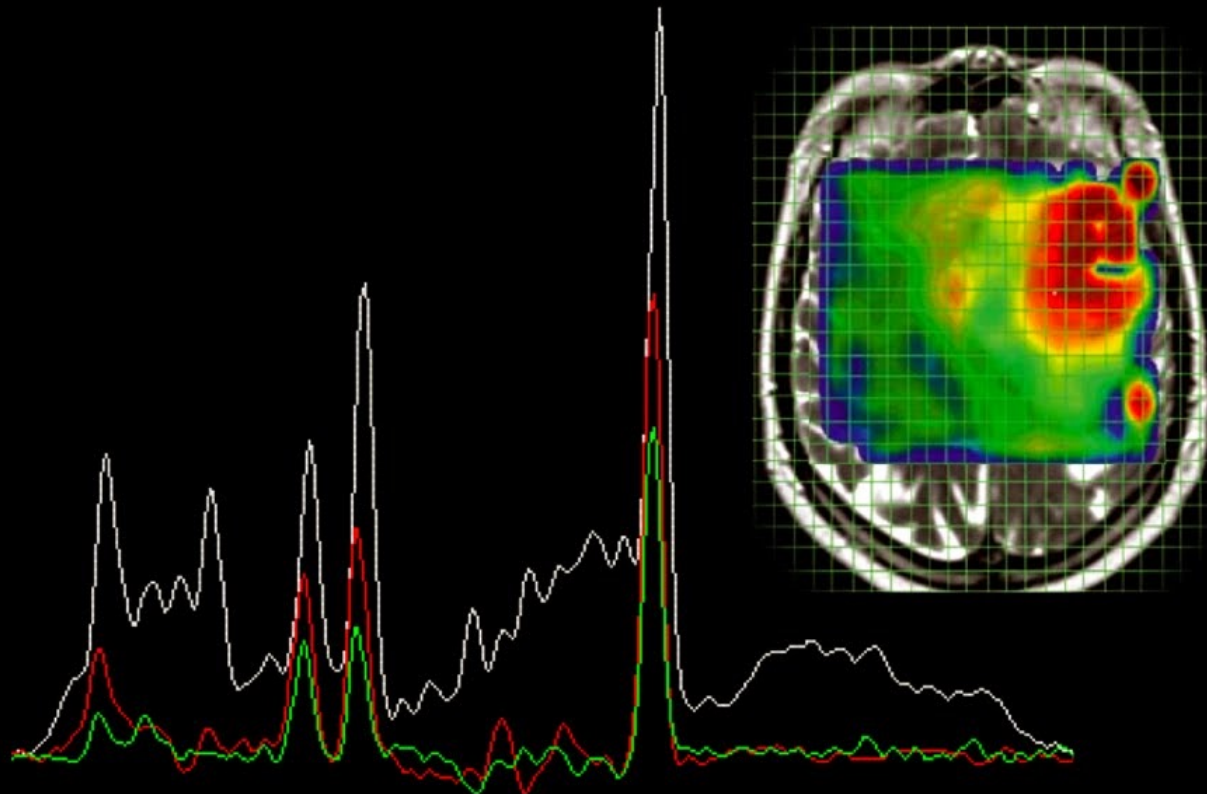
# Spectroscopic imaging (Chemical shift imaging CSI)

- ✓ CSI is an acquisition sequence that allows the acquisition of a spectrum per each voxel
- ✓ The acquisition time is large (> 10 minutes)



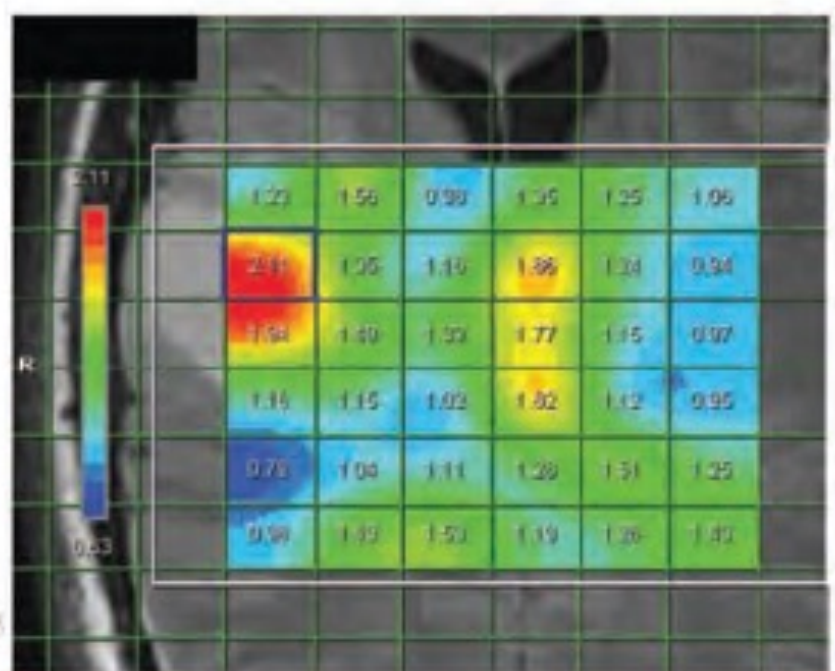
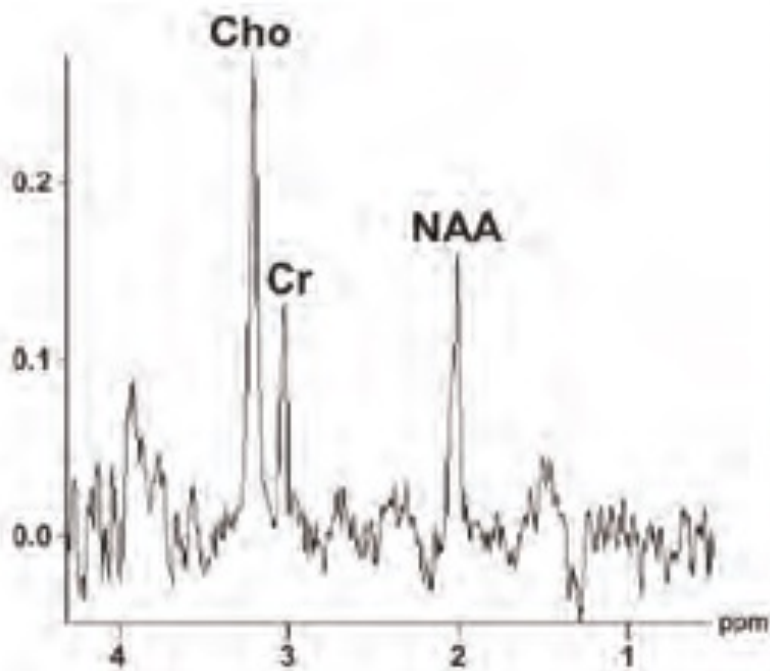
# Spectroscopic imaging (Chemical shift imaging CSI)

## Magnetic Resonance Spectroscopy



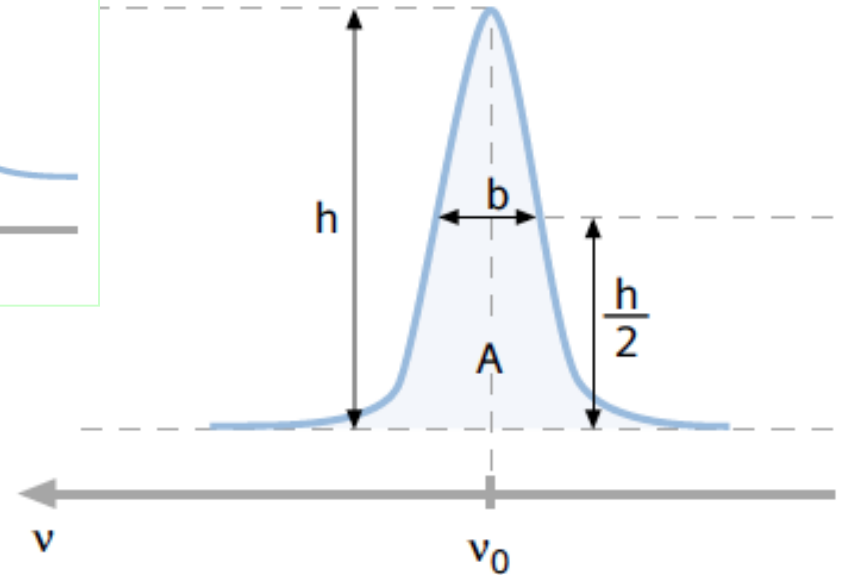
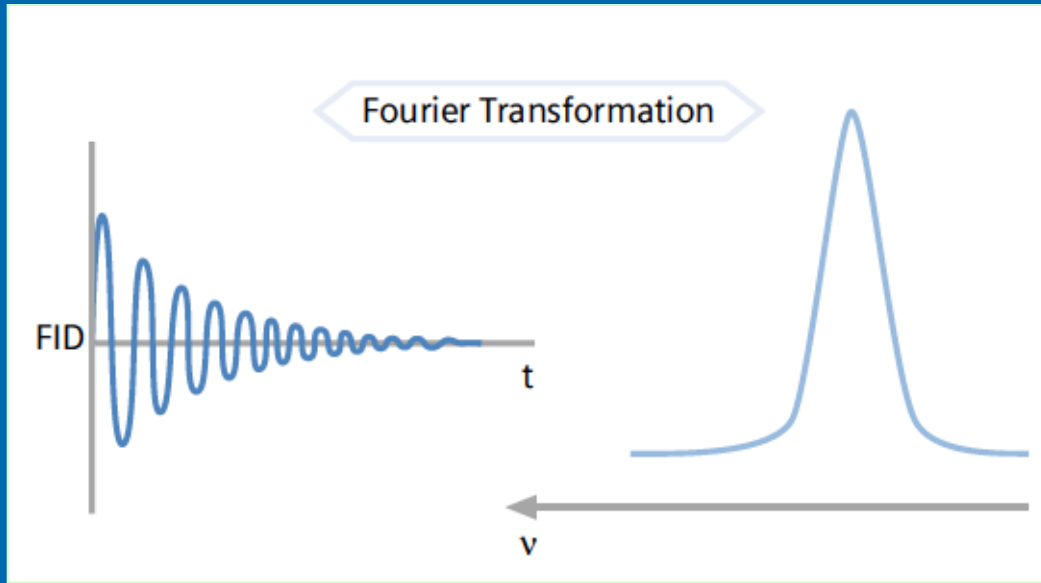
# Spectroscopic imaging (Chemical shift imaging CSI)

RadioGraphics 2006; 26:S173-S189



MR Spectrum from anaplastic oligoastrocytoma Choline / Creatine ratio map

# Quantitative data analysis



- Resonance frequency:  $v_0$
- Height of peak:  $h$
- Full Width at Half Maximum, FWHM:  $b$
- Integral:  $A$



# Applications



## Functional Magnetic Resonance Spectroscopy: The “New” MRS for Cognitive Neuroscience and Psychiatry Research

Jeffrey A. Stanley<sup>1\*</sup> and Naftali Raz<sup>2,3,4</sup>

<sup>1</sup>Department of Psychiatry and Behavioral Neurosciences, School of Medicine, Wayne State University, Detroit, MI, United States, <sup>2</sup>Department of Psychology, Wayne State University, Detroit, MI, United States, <sup>3</sup>Institute of Gerontology, Wayne State University, Detroit, MI, United States, <sup>4</sup>Center for Lifespan Psychology, Max Planck Institute for Human Development, Berlin, Germany

Preliminary evidence of the ability of <sup>1</sup>H fMRS to detect changes in glutamate during various perceptual, motor, and cognitive tasks.

# Applications

**Magnetic resonance spectroscopy assessment of brain injury after moderate hypothermia in neonatal encephalopathy:**

**a prospective multicentre cohort study**

*Peter J Lally et al for the MARBLE consortium*

*Lancet Neurol 2019; 18: 35–45*

## Summary

**Thalamic proton MRS measures acquired soon after birth in neonatal encephalopathy had the highest accuracy to predict neurodevelopment 2 years later.**

# Applications

**Magnetic resonance spectroscopy assessment of brain injury after moderate hypothermia in neonatal encephalopathy:  
a prospective multicentre cohort study**

*Lancet Neurol 2019; 18: 35–45*

## Methods

- ✓ **3.0 Tesla scanner**
- ✓ **single  $15 \times 15 \times 15 \text{ mm}^3$  voxel centred on the left thalamus**
- ✓  **$^1\text{H}$  MRS metabolite peak area ratios (7 min)**
- ✓  **$^1\text{H}$  MRS metabolite absolute concentrations (25 min)**
- ✓ **diffusion weighted MRI (DW MRI; 7 min)**