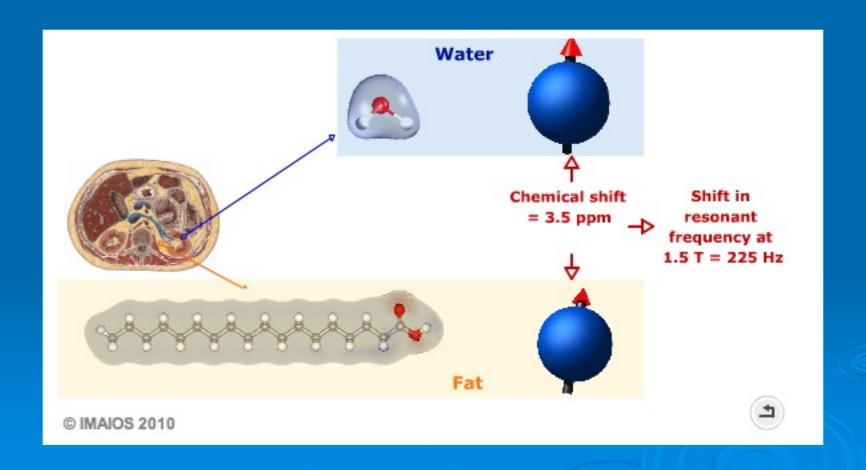
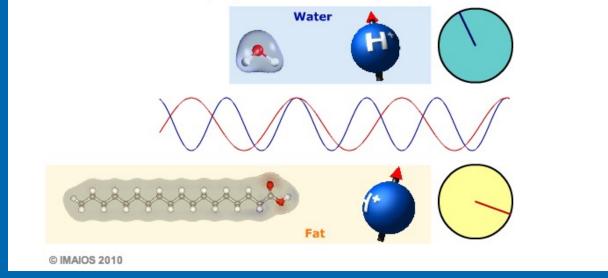
# MRS Magnetic Resonance Spectroscopy

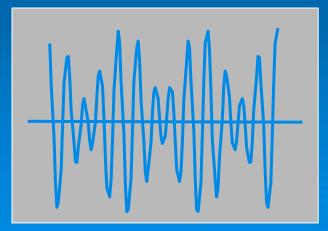
In vivo biochemistry

#### **Chemical shift**

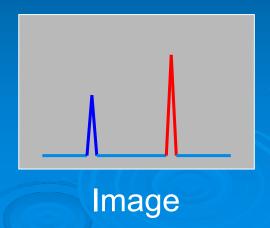


# Chemical shift and MR spectrum





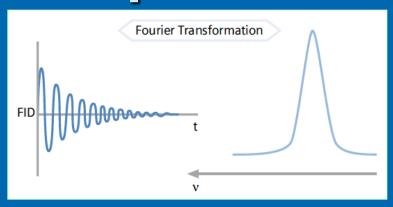
Fourier Transform

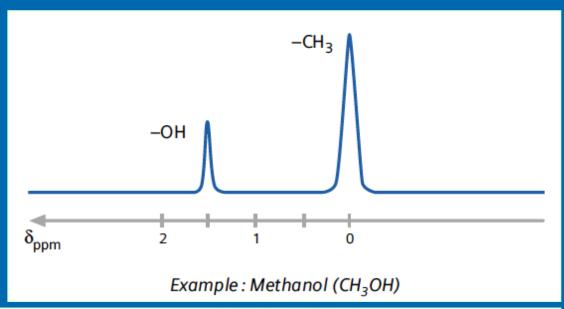


**Received Signal** 

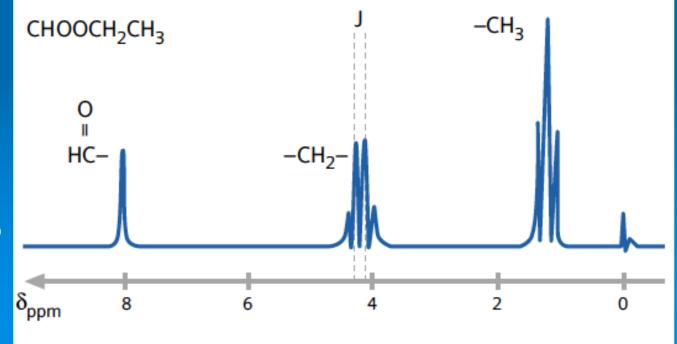
www.imaios.com/en/e-Courses/e-MRI/Image-quality-and-artifacts/chemical-shift

# <sup>1</sup>H NMR spectra

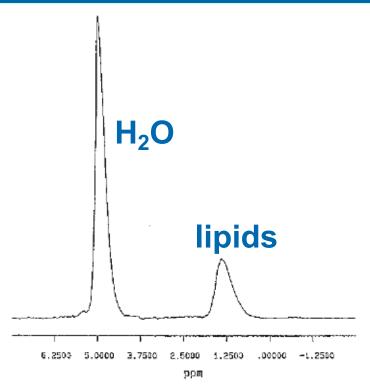




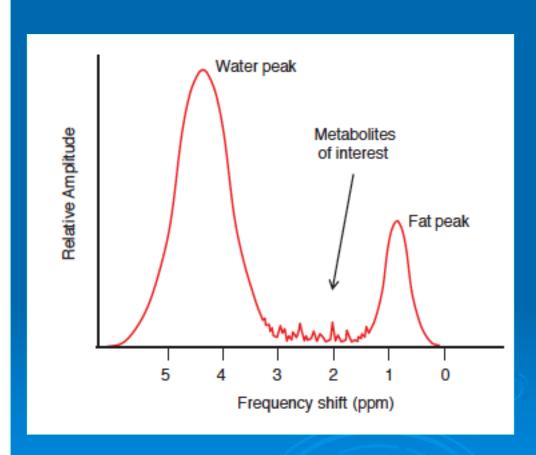
- ✓ Peaks pattern is footprint of molecules
- ✓ Peak area is proportional to the number of <sup>1</sup>H nuclei



#### Proton spectroscopy

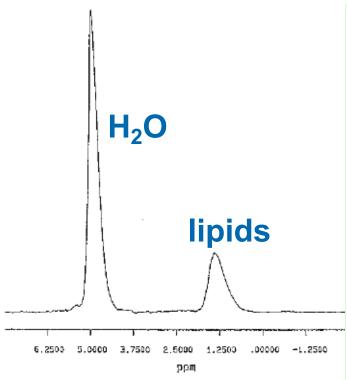


**Figure 1.** Typical in vivo H-1 spectrum from a steatotic liver (TR = 3 seconds, TE = 24 and 50 msec, four signals acquired, 262 cm<sup>3</sup> volume of interest). No signal filtering before Fourier transformation and no baseline correction were applied.

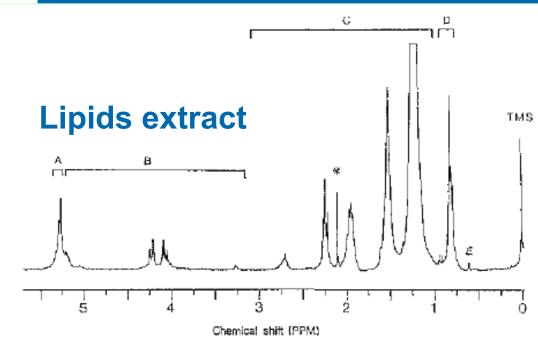


The water signal has to be suppressed

#### Proton spectroscopy



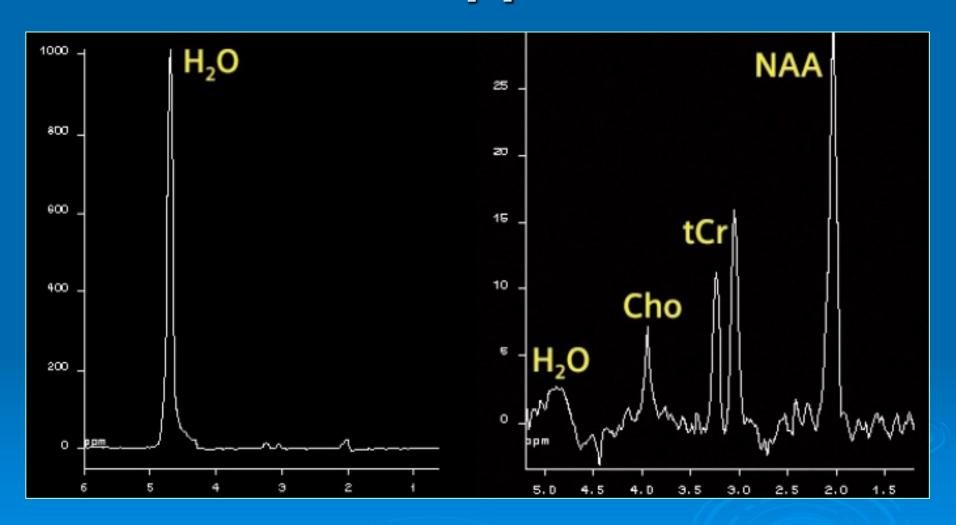
**Figure 1.** Typical in vivo H-1 spectrum from a steatotic liver (TR = 3 seconds, TE = 24 and 50 msec, four signals acquired, 262 cm<sup>3</sup> volume of interest). No signal filtering before Fourier transformation and no baseline correction were applied.



**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

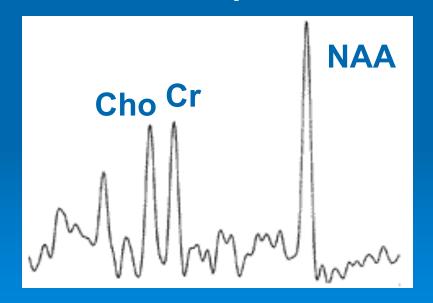
#### <sup>1</sup>H MRS metabolites

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	N-acetyl-∟-aspartate	2.01	Presence of intact glioneural structures
Lactate		1.33 (inverted)	Anaerobic glycolysis
Lipids	Free fatty acids	1.2-1.4	Necrosis

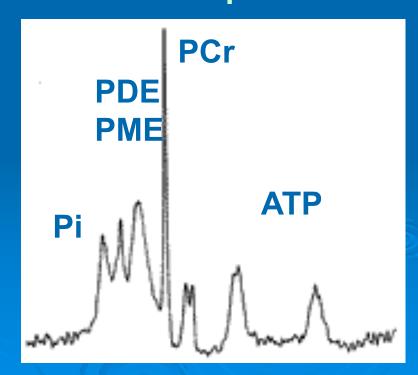
#### Brain spectroscopy

➤ The ¹H (or ³¹P) nuclei in different molecules have slightly different resonance frequencies

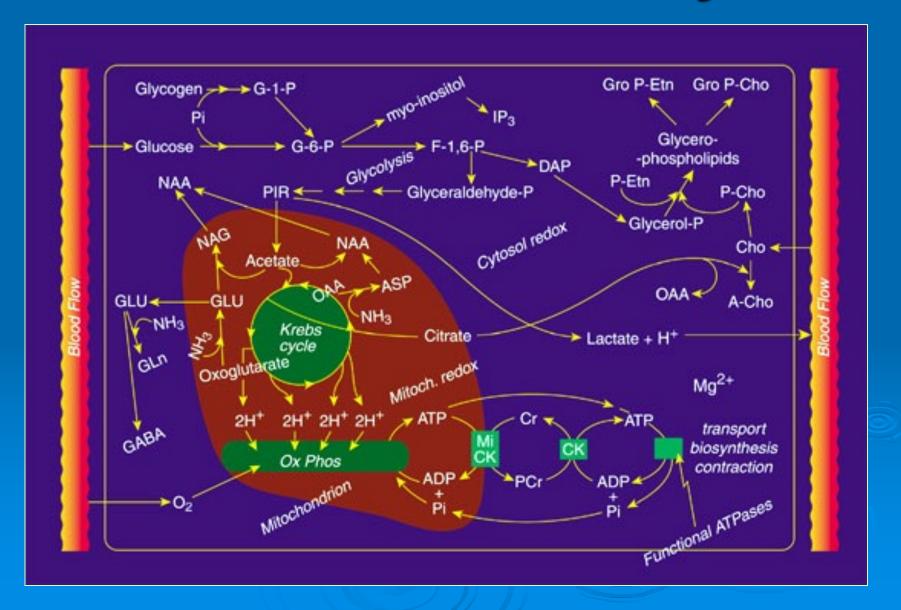
Each peak is related to a molecules (metabolite)
In vivo ¹H spectrum
In vivo ³¹P spectrum



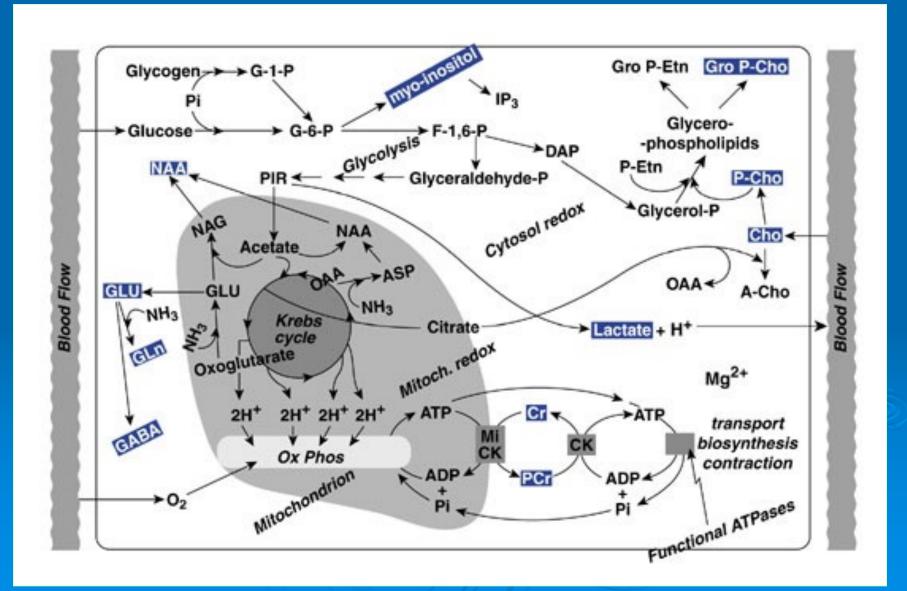
Occipital cortex



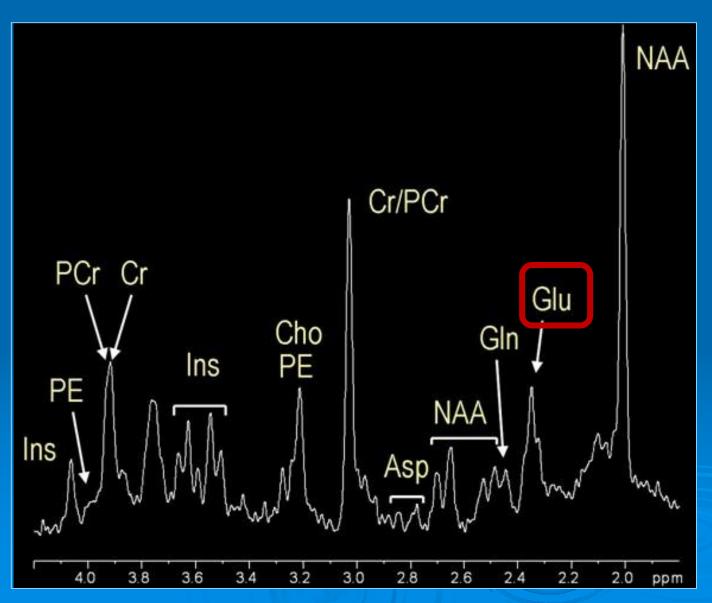
#### In vivo biochemistry



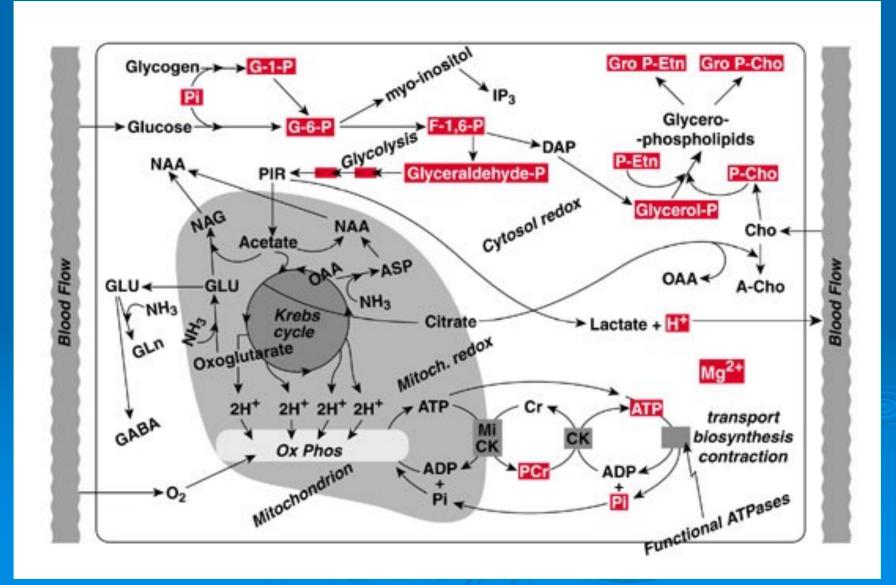
#### <sup>1</sup>H spectroscopy



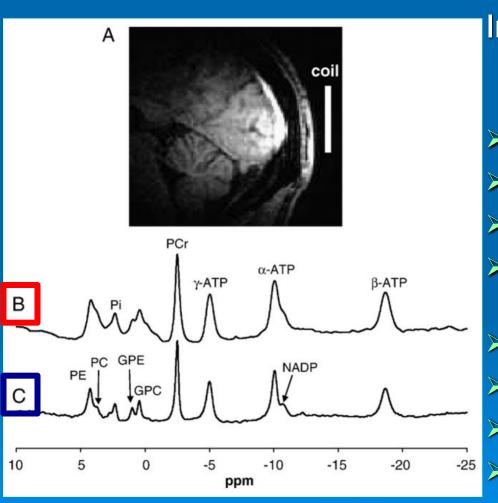
#### <sup>1</sup>H spectroscopy



#### <sup>31</sup>P spectroscopy



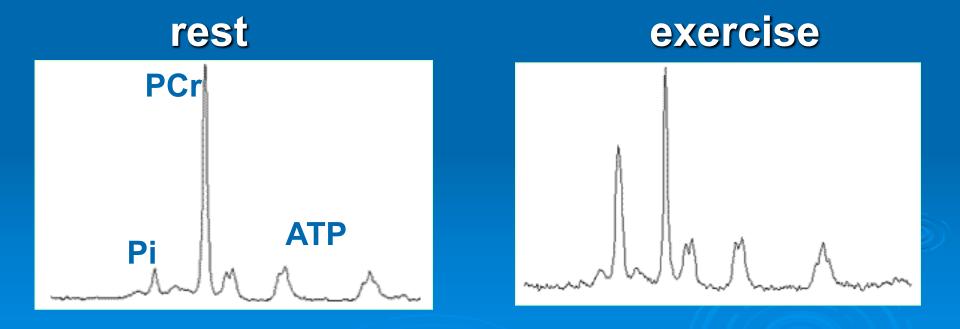
#### <sup>31</sup>P MRS



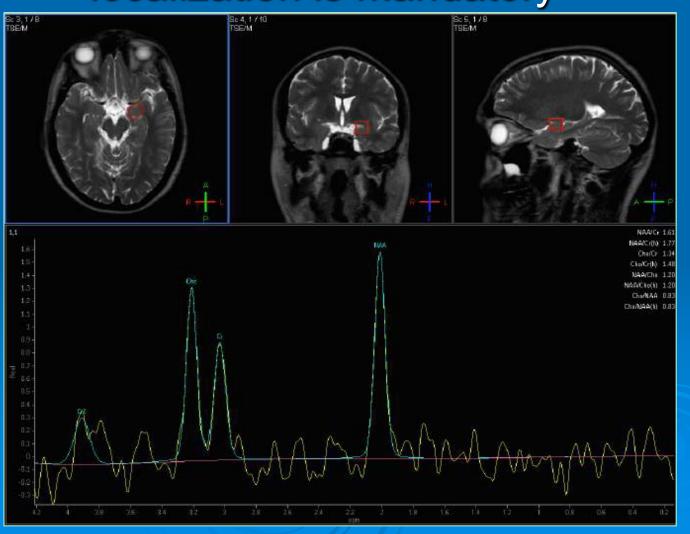
- In vivo <sup>31</sup>P spectra acquired from the human occipital lobe at (B) 4 T and (C) 7 T:
- > PE phosphoethanolamine
- > PC phosphocholine
- Pi inorganic phosphate
- GPE glycerophosphoethanolamine
- GPC, glycerophosphocholine
- PCr phosphocreatine
- ATP adenosine triphosphate
- NADP nicotinamide adenine dinucleotide phosphate

#### muscle 31PMRS

Muscoli gastrocnemi normal subject

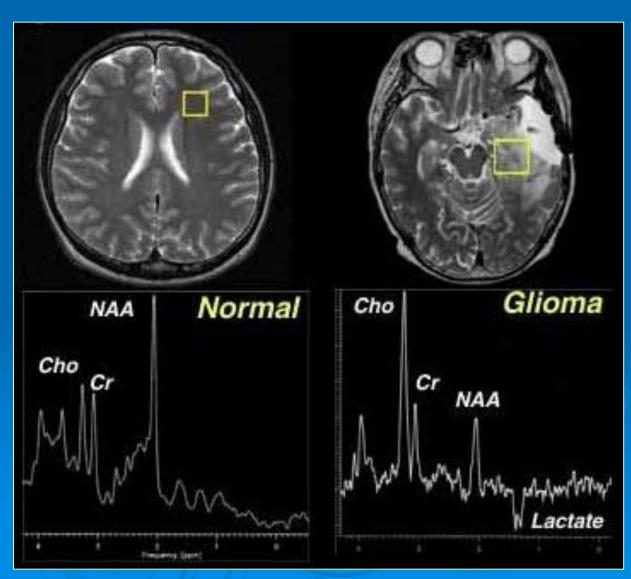


# Signal localization In in vivo MRS the signal localization is mandatory



#### Signal localization

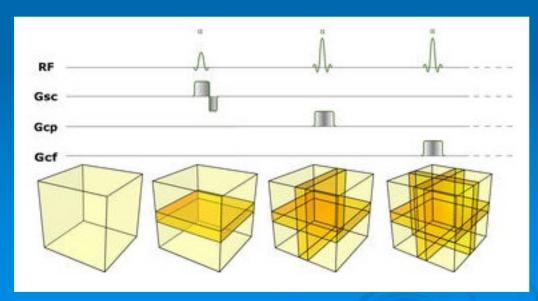
In in vivo
MRS the
signal
localization is
mandatory

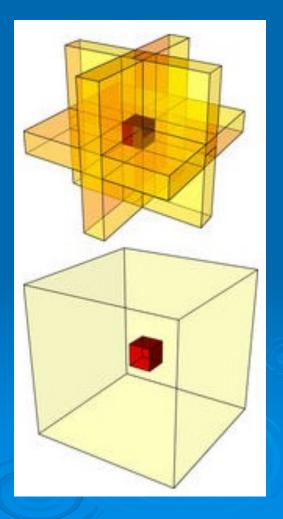


<sup>1</sup>H MRS

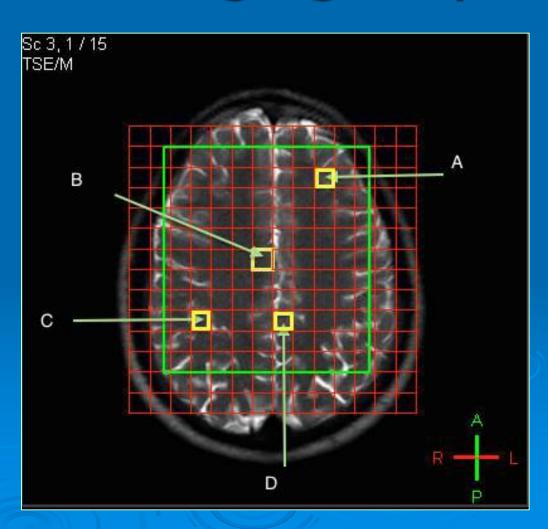
#### Volume selection

- √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



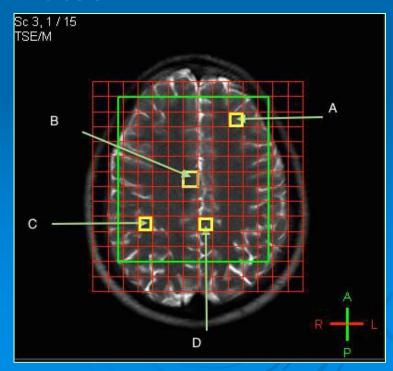


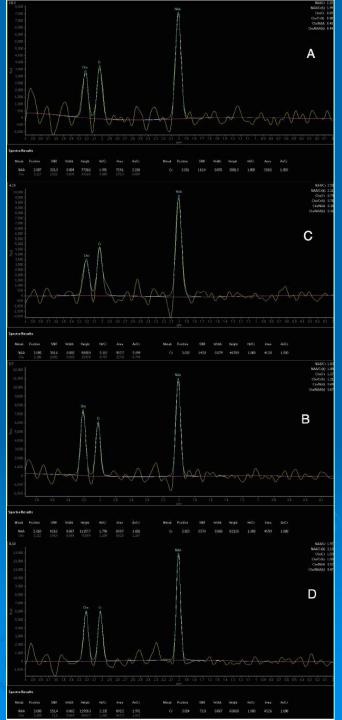
- ✓ CSI is an acquisition sequence that allows the acquisition of a spectrum per each voxel
- √The acquisition time
  is large (> 10 minutes)



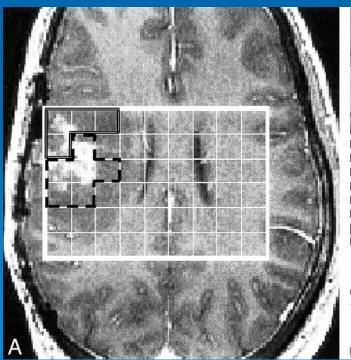
# Spectroscopic imaging

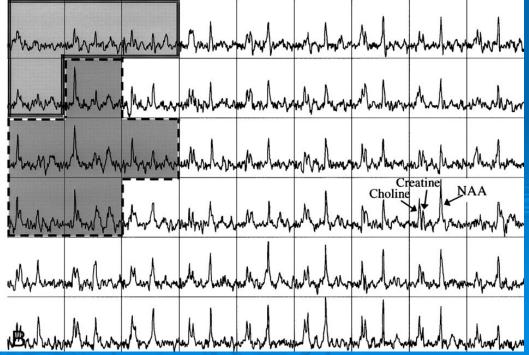
- ✓ CSI is an acquisition sequence that allows the acquisition of a spectrum per each voxel
- √The acquisition time is large
  - > 10 minutes

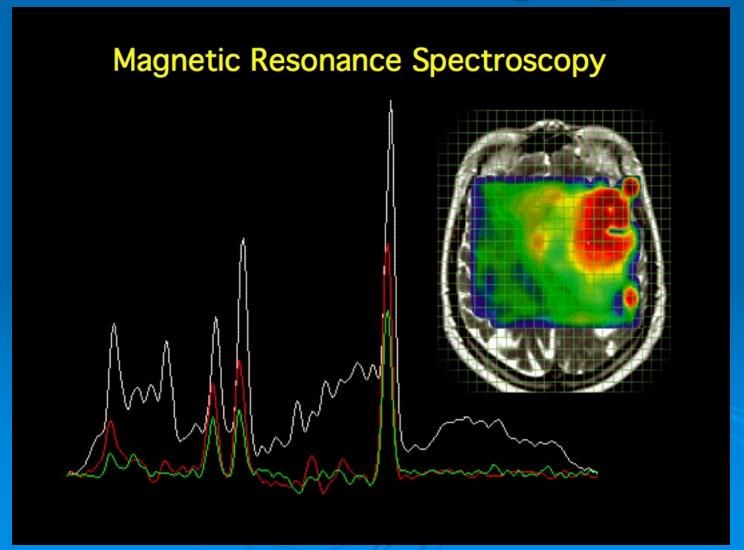


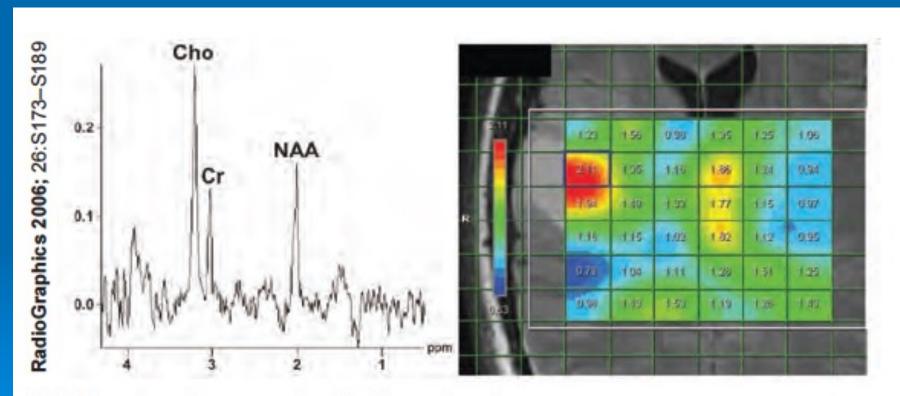


✓ CSI is an acquisition sequence that allows the acquisition of a spectrum per each voxel
 ✓ The acquisition time is large (> 10 minutes)



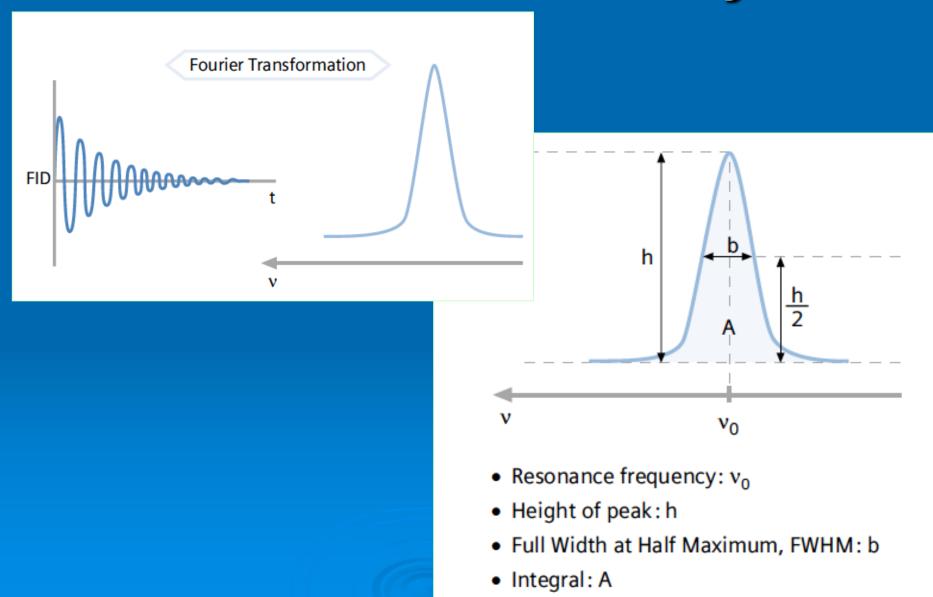






MR Spectrum from anaplastic oligoastrocytoma Choline / Creatine ratio map

#### Quantitative data analysis



#### **Applications**



REVIEW

published: 12 March 2018 doi: 10.3389/fpsyt.2018.00076



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

Jeffrey A. Stanley1\* and Naftali Raz2,3,4

<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 <u>glutamate</u> 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 15×15×15 mm³ voxel centred on the left thalamus
- √ ¹H MRS metabolite peak area ratios (7 min)
- ✓ 1H MRS metabolite absolute concentrations (25 min)
- √ diffusion weighted MRI (DW MRI; 7 min)