

NeuroMET

Virtual MR Spectroscopy Workshop

L2: Localization Methods

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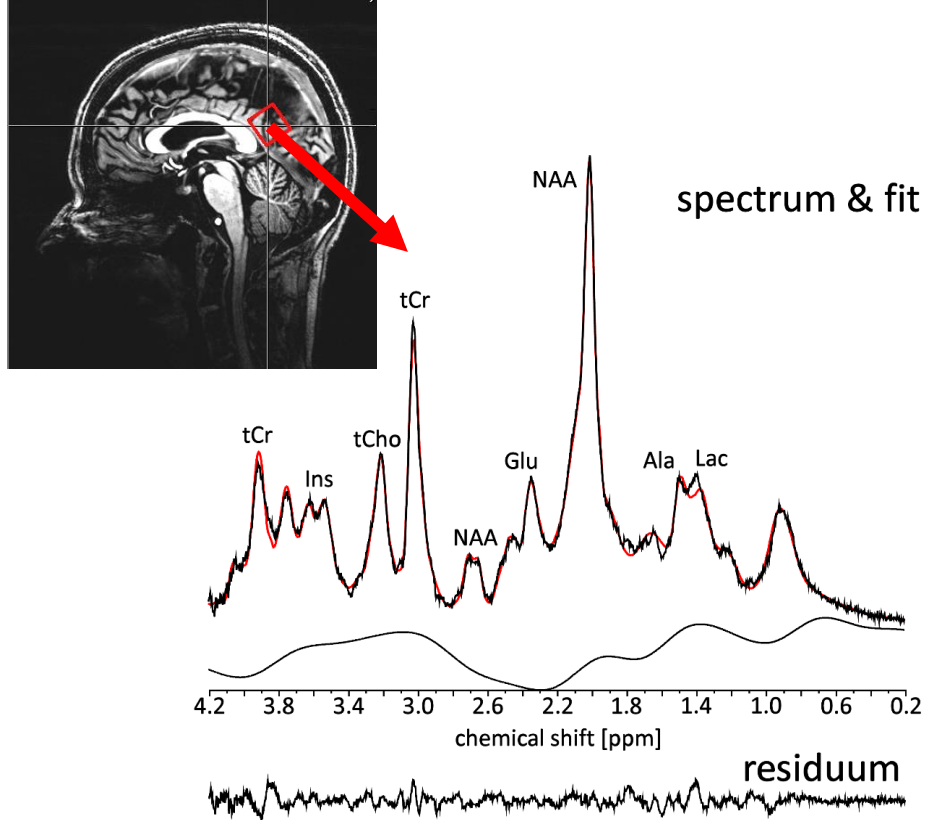
Physikalisch-Technische Bundesanstalt, AG «In-vivo MRT»



NeuroMET



How do we get there?



Content

1. SVS Localization Methods

1.1 PRESS



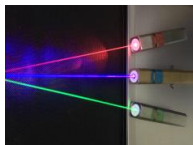
1.2 STEAM



1.3 ISIS \rightarrow ^{31}P MRS



1.4 (s)LASER



1.5 SPECIAL



2. Further Methods

2.1 MEGA

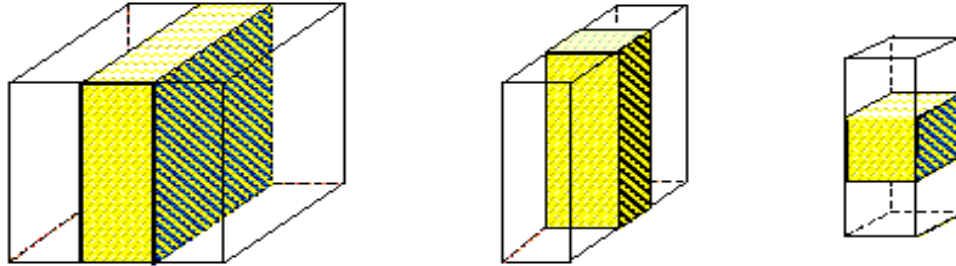


2.2 MRSI/CSI

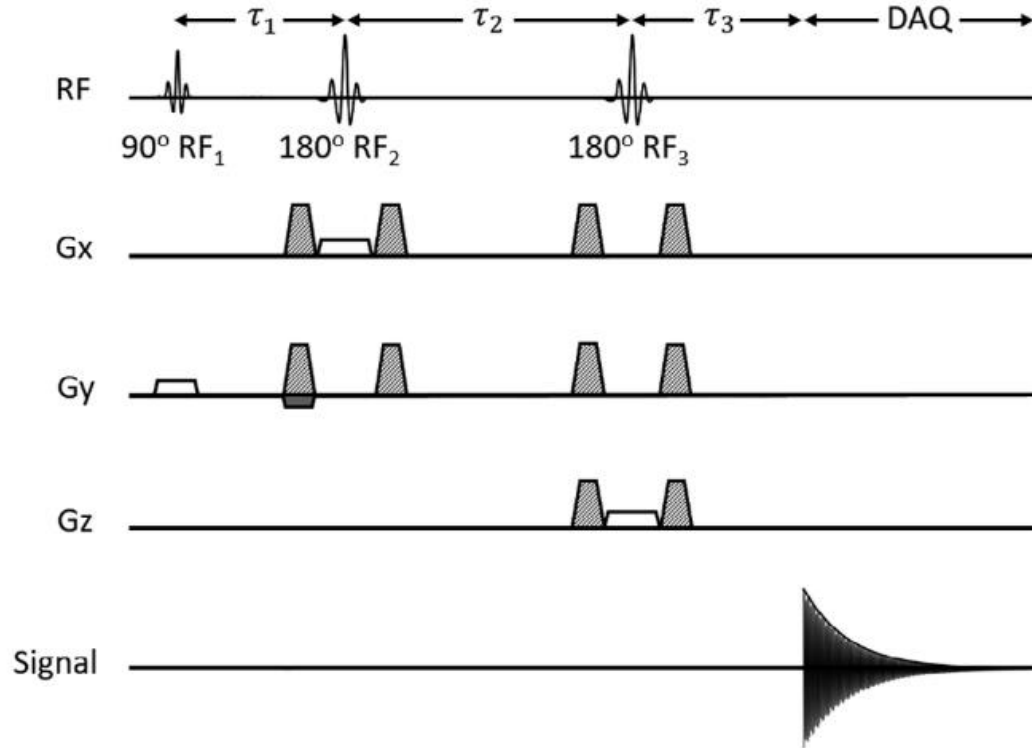


1. Localisation by gradients

in each direction a gradient field defines
one slice \rightarrow the intersection of these
slices is the selected volume of interest



1. Sequence diagram



1.1 Spin Echo Selection

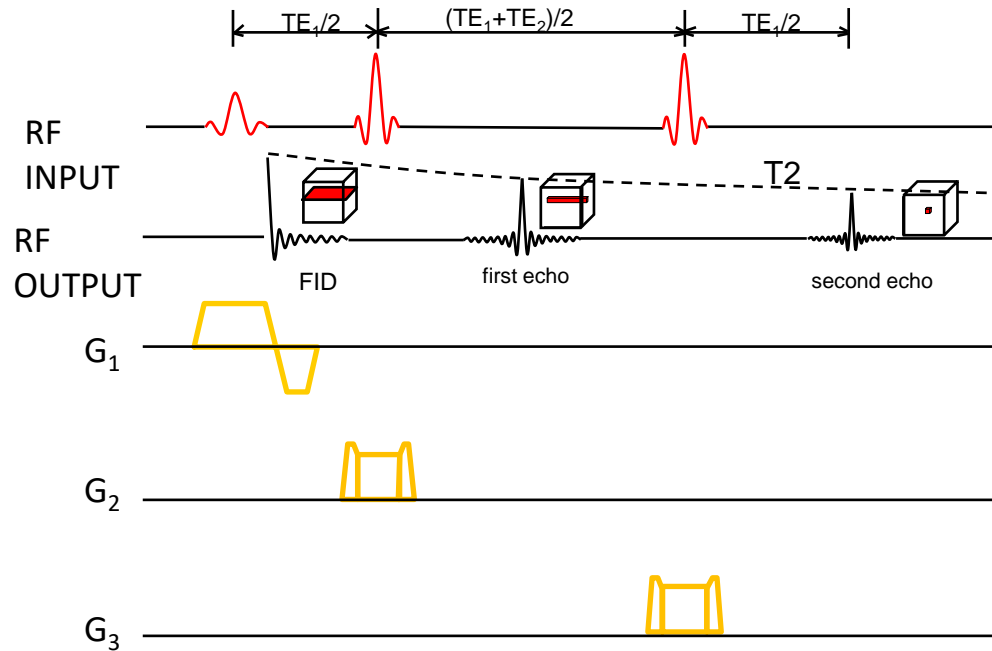


<https://www.drcmr.dk/BlochSimulator/>

1.1 Spin Echo Selection



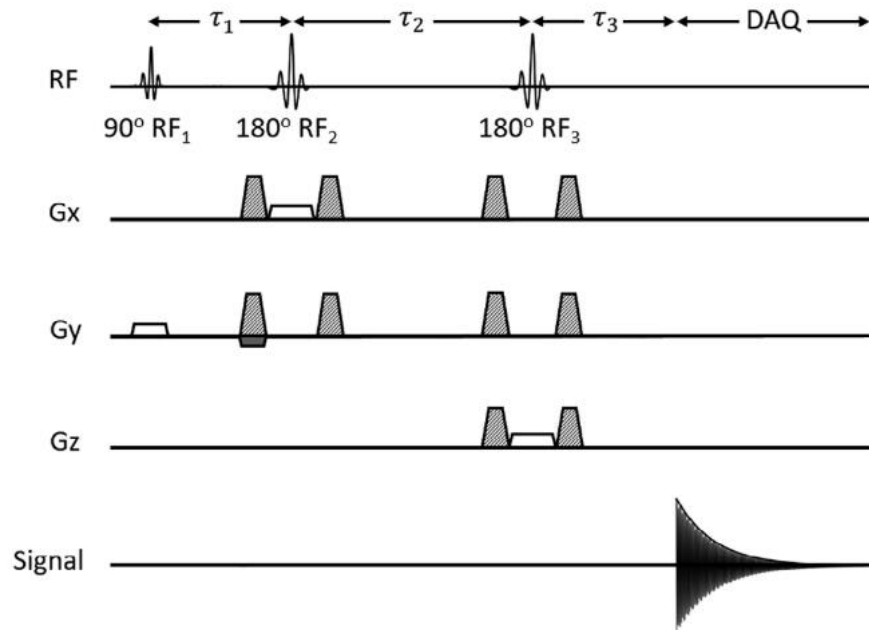
Method of choice for ^1H spectroscopy



1.1 PRESS



- Point **RES**olved **S**pectroscopy
- Double spin echo sequence: 90° - 180° - 180° - echo acquisition
- Spoiler/crusher gradients: eliminate unwanted echo signals from outside the selected volume



Landheer et al., Journal of MRI, 2019

1.1 PRESS



Advantages:

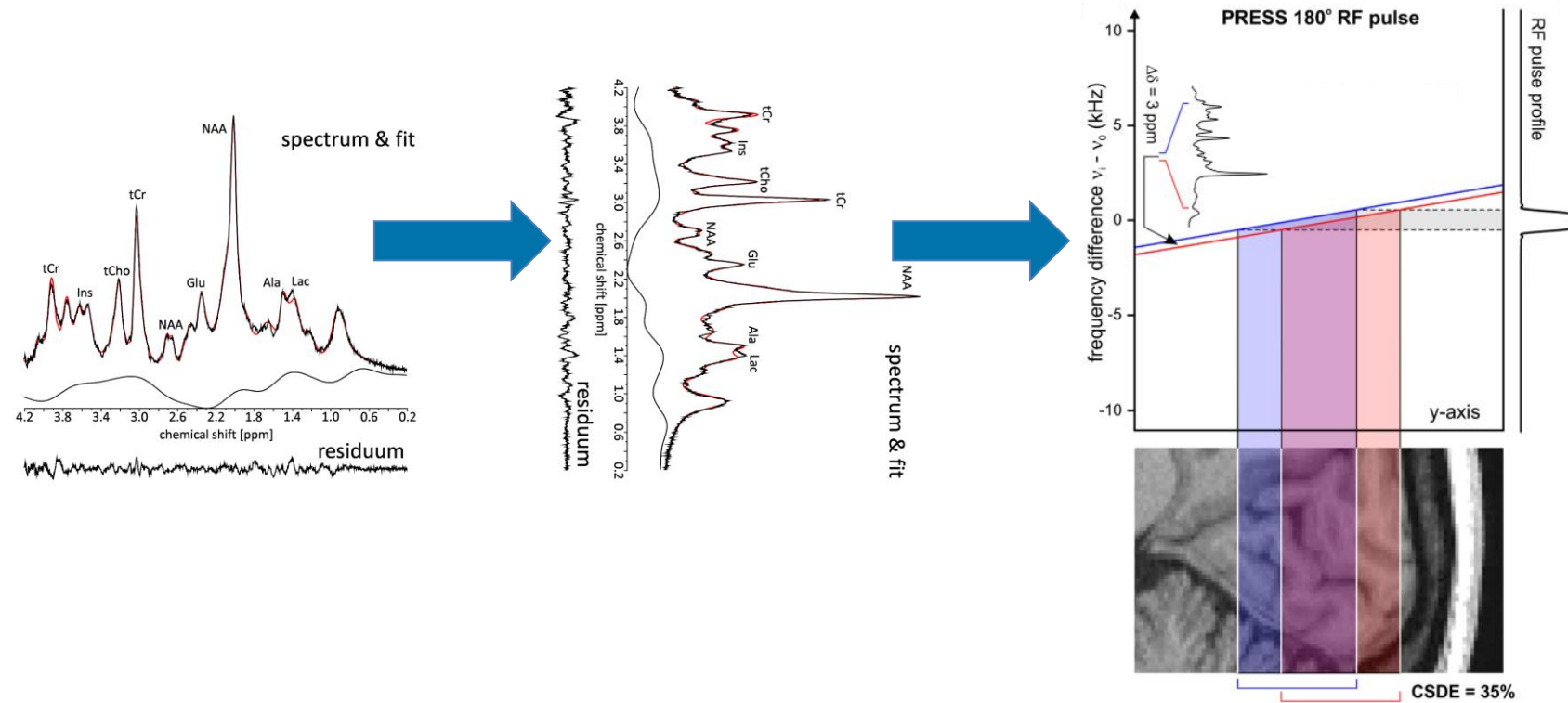
- Good localization and strong suppression of signals outside the selected volume in one measurement → widely used for ^1H spectroscopy
- Less motion sensitive than STEAM (in a minute 😊)
- Twice as much signal as with STEAM



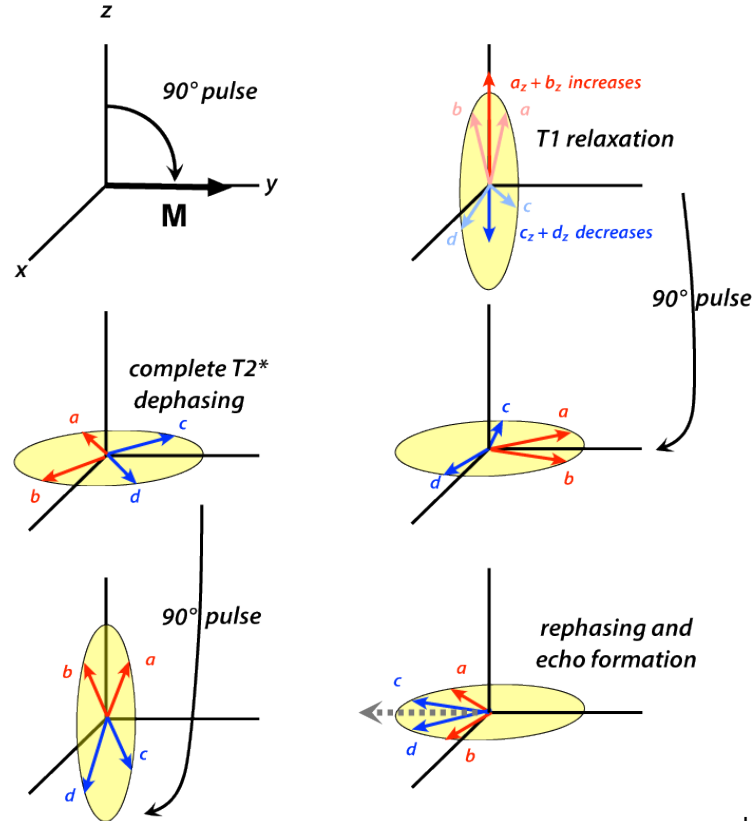
Disadvantages:

- Minimal echo time: 19 ms (1.5T) / 31 ms (3T)
- 3T, 7T: large chemical shift displacement (CSD)

Chemical Shift Displacement (CSD)

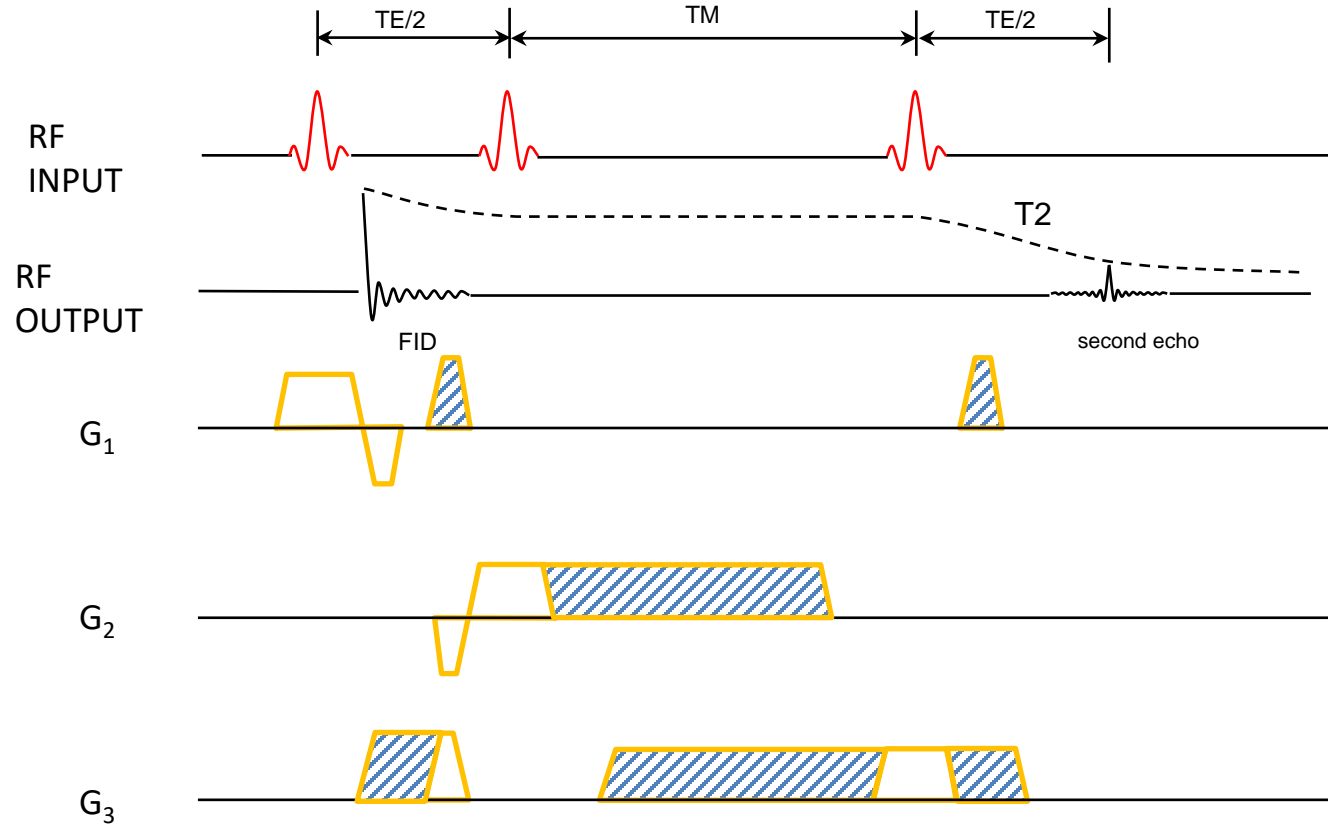


1.2 Stimulated Echo



<https://www.mriquestions.com/stimulated-echoes.html>

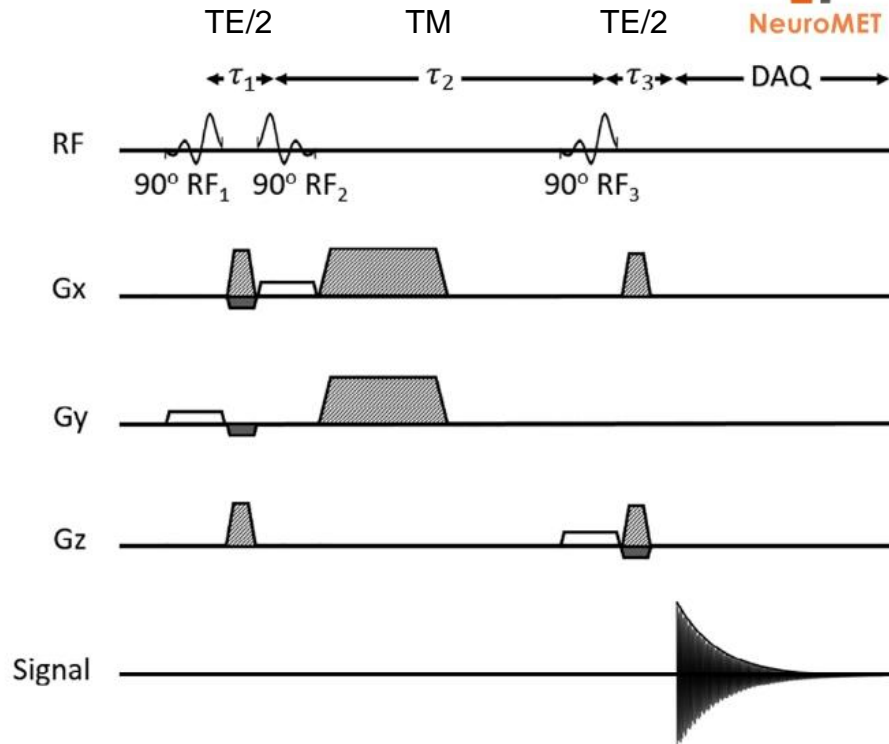
1.2 Stimulated Echo



1.2 STEAM



- Stimulated Echo Acquisition Mode
- Selective excitation
- Uses 3 slice selective 90° pulses forming a stimulated echo after TE + TM



Landheer et al., Journal of MRI, 2019

1.2 STEAM



Advantages:

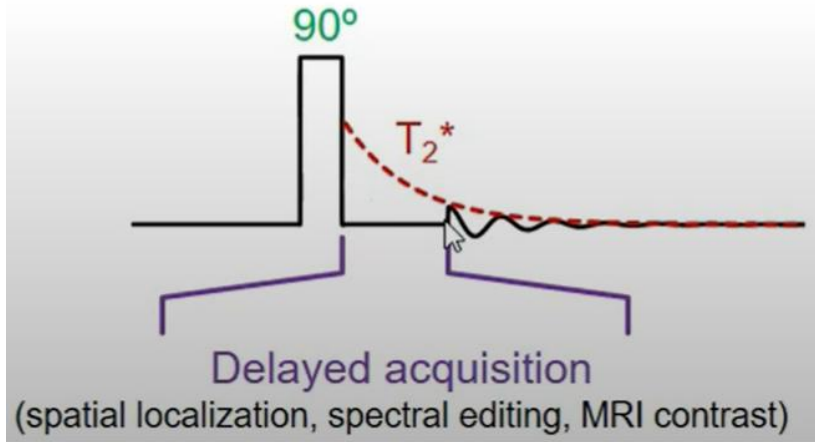
- Allows shorter echo times than PRESS (down to 5 ms)
- Less CSD due to using only 90° pulses



Disadvantages:

- Signal intensities are only **half** of those obtained with PRESS

How do we get a signal if the FID is gone?



- T_2^* occurs from macroscopic, reversible loss of phase coherence in addition to T_2 relaxation
- T_2 occurs from microscopic, irreversible loss of phase coherence
- $T_2 \gg T_2^*$

→ We have to compensate for T_2^* relaxation

<https://www.youtube.com/watch?v=A7Pa-rIVzao&list=PL471uBfQUs9qcODBkQGJTZkcbIIKrkjKry&index=5>

1.3 ISIS

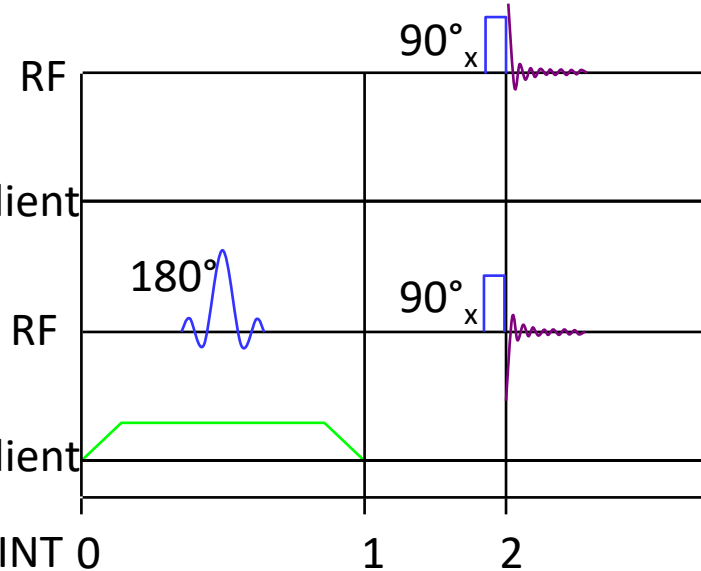


<https://www.mriquestions.com/isis.html>

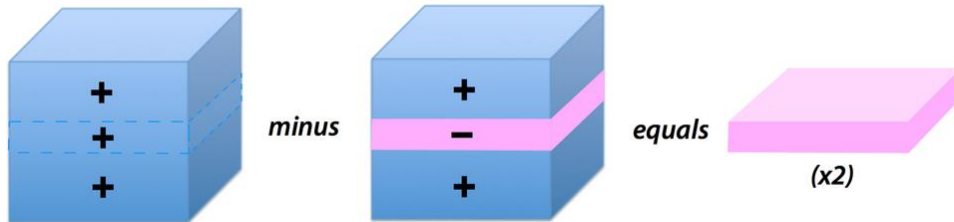
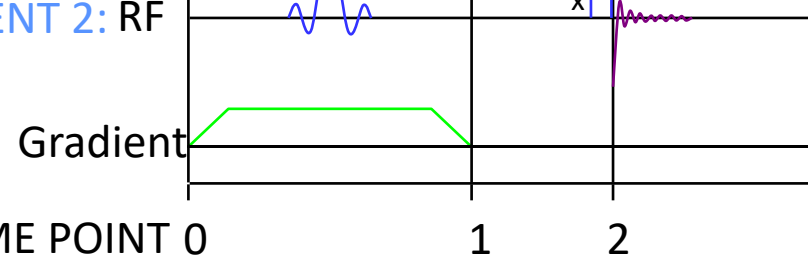


- Image **S**electe**d** **I**n vivo **S**pectroscopy (oldest SVS technique)
- Method of choice for ^{31}P , ^{19}F MRS

EXPERIMENT 1:



EXPERIMENT 2:



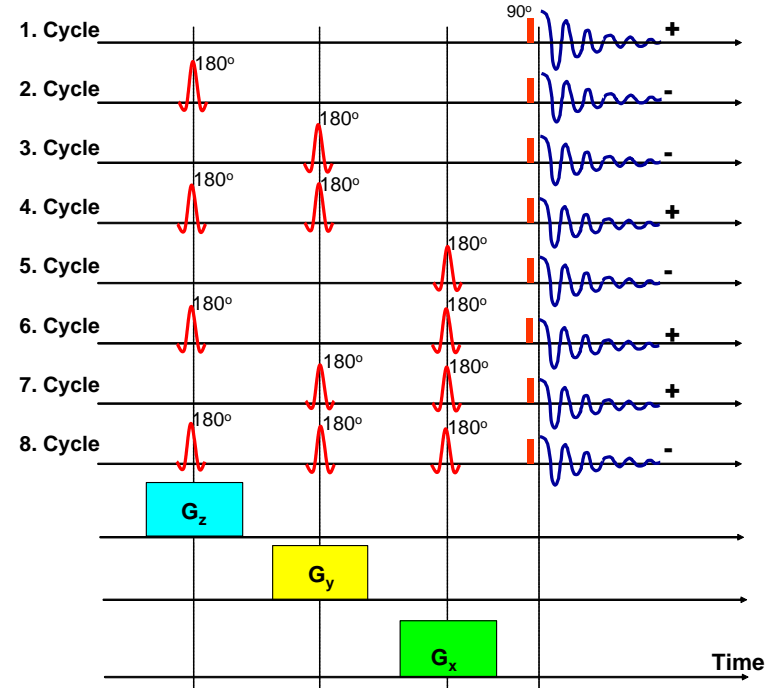
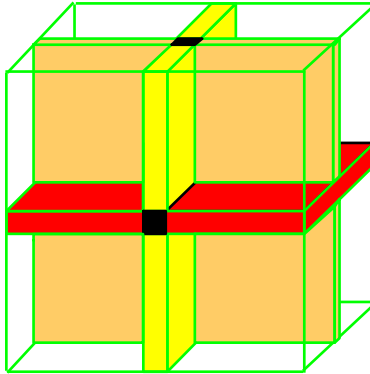
1.3 ISIS



<https://www.mriquestions.com/isis.html>



- Image **S**electe**d** **I**n vivo **S**pectroscopy (oldest SVS technique)
- Method of choice for ^{31}P , ^{19}F MRS
- 2D needs 4 experiments, 3D needs 8



1.3 ISIS



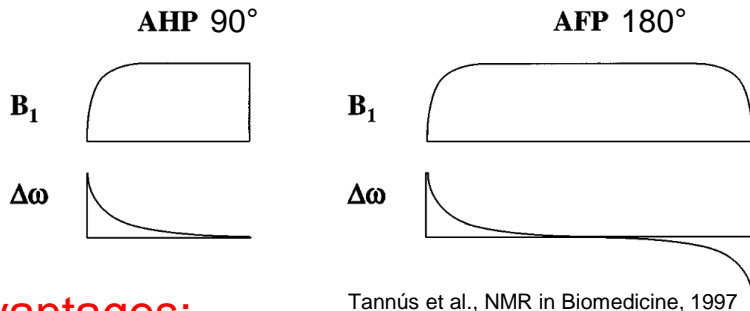
Advantages:

- FID acquisition starts immediately after excitation → no T_2 -loss, no J-modulation → primarily used for ^{31}P MRS because of short T_2 relaxation times
- Easily adjustable for 2 or 4 volumes with linear time increase
- The 8 needed phase cycles hardly increase the measurement time, as *in vivo* ^{31}P spectroscopy needs multiple signal acquisitions

Disadvantages:

- Large signals are subtracted from each other → sensible to instrumental instabilities, motion, T_1 saturation
→ Not used for ^1H spectroscopy anymore

1.4.1 Adiabatic Pulses



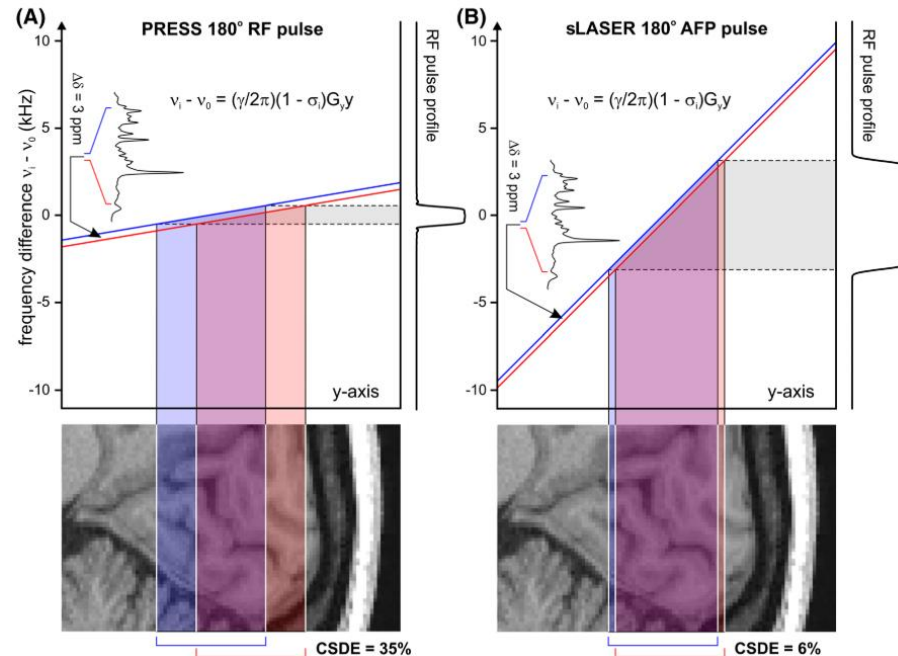
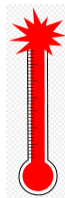
Youtube: Robin de Graaf, 'Basics of in vivo NMR'

Advantages:

- Insensitive to B_1 inhomogeneities
- Smaller CSD
- Larger bandwidth \rightarrow insensitive to B_0 inhomogeneities

Disadvantages:

- Increased specific absorption rate (SAR)
- Long pulse duration

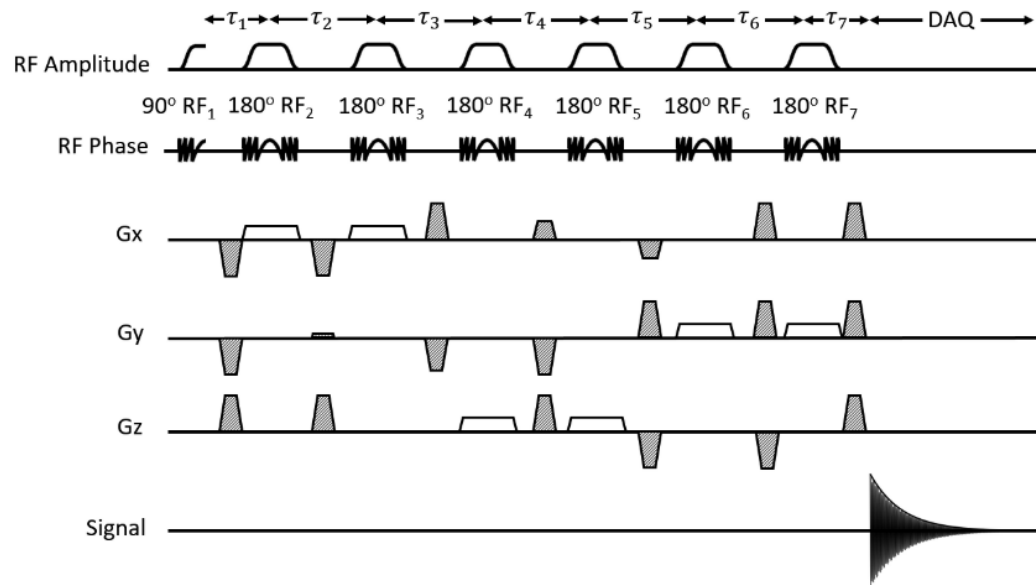


Öz et al., NRM in Biomedicine, 2020

1.4.1 LASER



- Localization by **A**diabatic **S**elective **R**efocusing



- Non-selective excitation
- Uses 3 slice selective 180° pulse pairs for slice selection

Landheer et al., Journal of MRI, 2019

1.4.1 LASER



Advantages:

- Reduced CSD compared to PRESS
- Reduced B_1 sensitivity



Disadvantages:

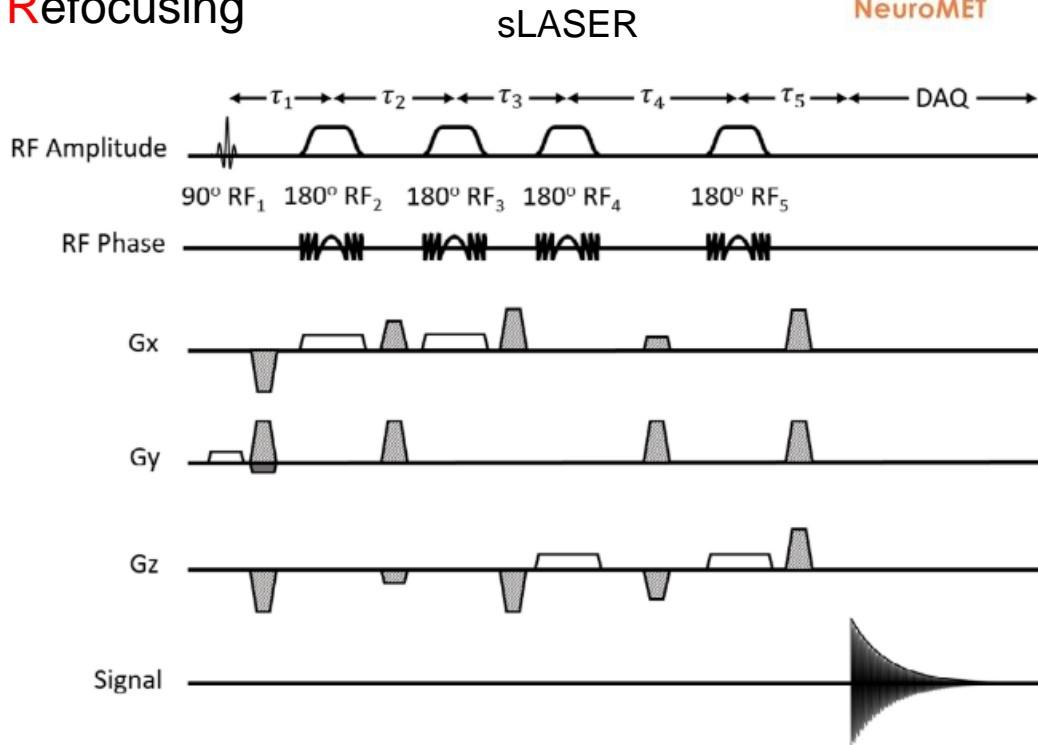
- Nonlinear phase profile of adiabatic pulse → paired pulses
→ increased TE compared to PRESS
- Higher SAR due to three refocussing pulse pairs

1.4.2 sLASER



- Semi Localization by Adiabatic Selective Refocusing

- Uses one slice-selective excitation pulse, followed by two pairs of adiabatic refocussing pulses (three pulses replaced from LASER: one non-selective excitation and two slice-selective refocussing)



Landheer et al., Journal of MRI, 2019

1.4.2 sLASER



Advantages:

- Smaller TE than LASER (7 T: 24 ms) → good for editing (section 2...stay thrilled)
- Lower SAR than LASER

Disadvantages:

- Excitation pulse sensitive to B_1 inhomogeneities and lower bandwidth because no adiabatic pulse

1.5 SPECIAL



- **SP**in **EC**ho, full **I**ntensity **A**cquired **L**ocalized

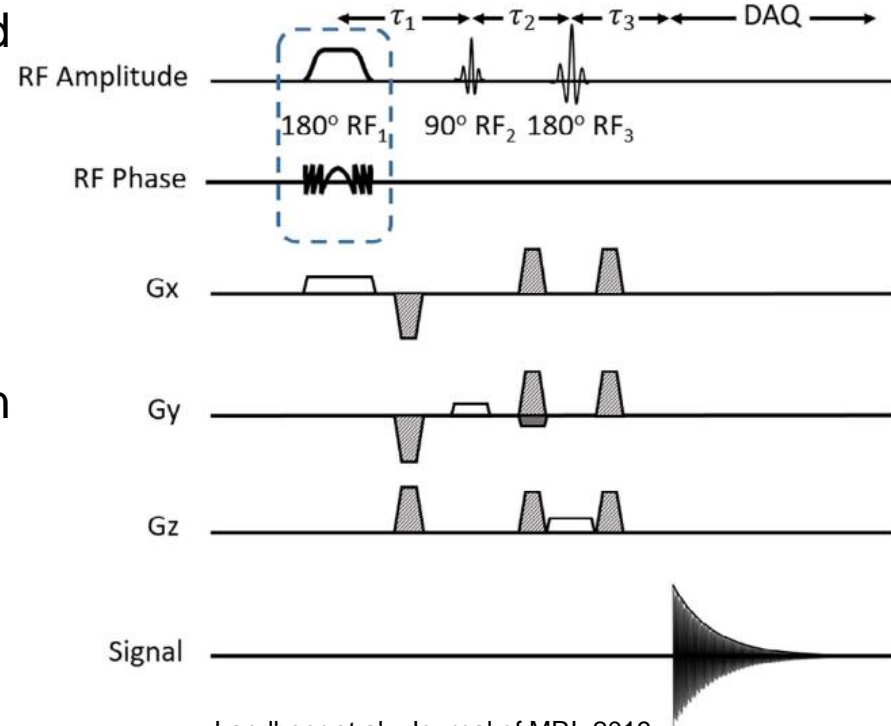
- Combination of



and



- Uses an on/off selective adiabatic inversion pulse, a 90° excitation and a 180° refocussing pulse



Landheer et al., Journal of MRI, 2019

1.5 SPECIAL



Advantages:

- Very short TEs ($3 T/7 T < 6/10$ ms) due to 1D ISIS before excitation

Disadvantages:

- Strong CSD in refocussing dimension → SPECIAL-sLASER
- two-cycle schemes → systematic variations between cycles manifest in their difference spectrum
- strong lipid contamination → lipid suppression techniques necessary

1. Conclusion



1. Conclusion

PRESS

Spin echo → longer TE_{\min}

best SNR

more CSD

→ 'standard' MRS sequence

STEAM

stimulated echo → shortest TE

only half the signal of PRESS

less CSD

„classical“ MRS
sequences

→ No adiabatic
pulses

ISIS

Historically oldest MRS sequence

Add/Subtract scheme

→ ^{31}P MR spectroscopy

1. Conclusion

LASER

Non-selective excitation with
adiabatic refocusing pulses
Lowest CSD

sLASER

Slice-selective excitation + LASER
Smaller SAR and TE than LASER

SPECIAL

1D ISIS (adiabatic pulse) + PRESS
Very short TEs
 $\rightarrow TE_{\text{SPECIAL}} < TE_{\text{sLASER}} < TE_{\text{LASER}}$

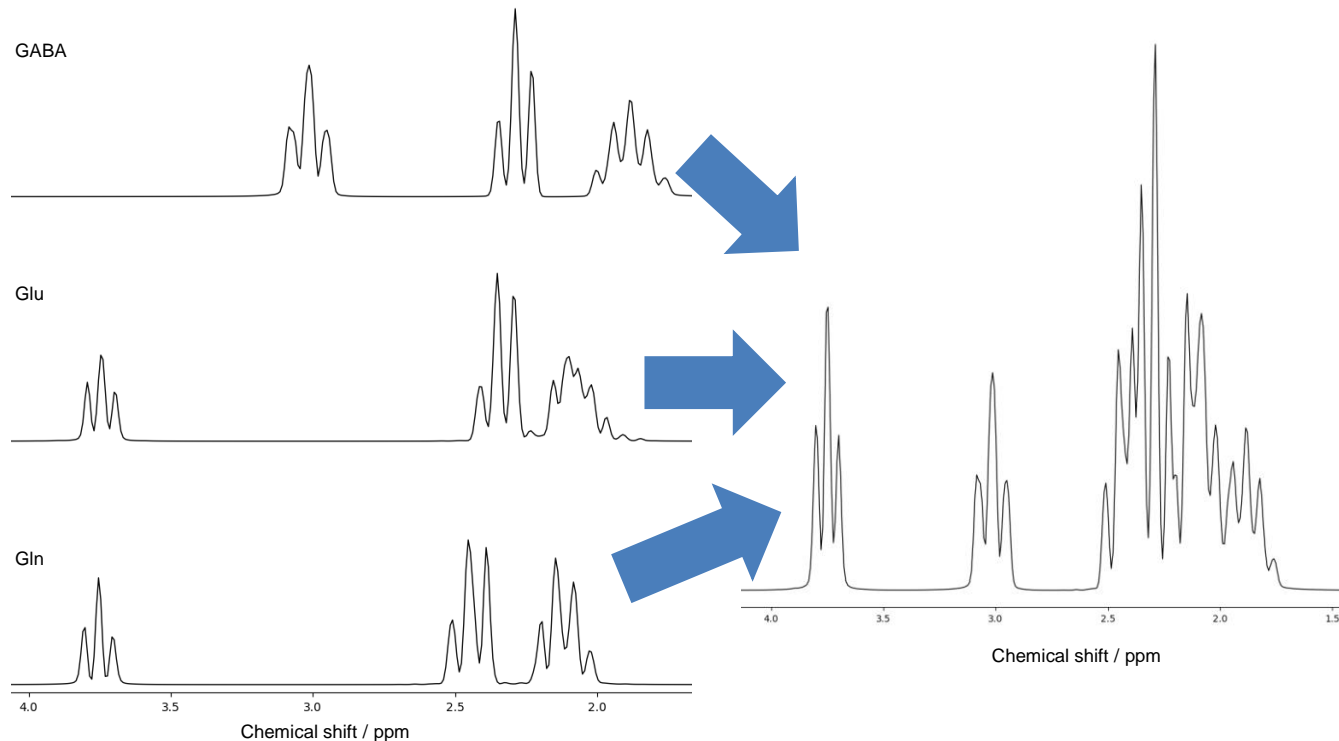
most of the latest
technical
developments

„modern“ MRS sequences
 \rightarrow adiabatic pulses

2.1 MEGA

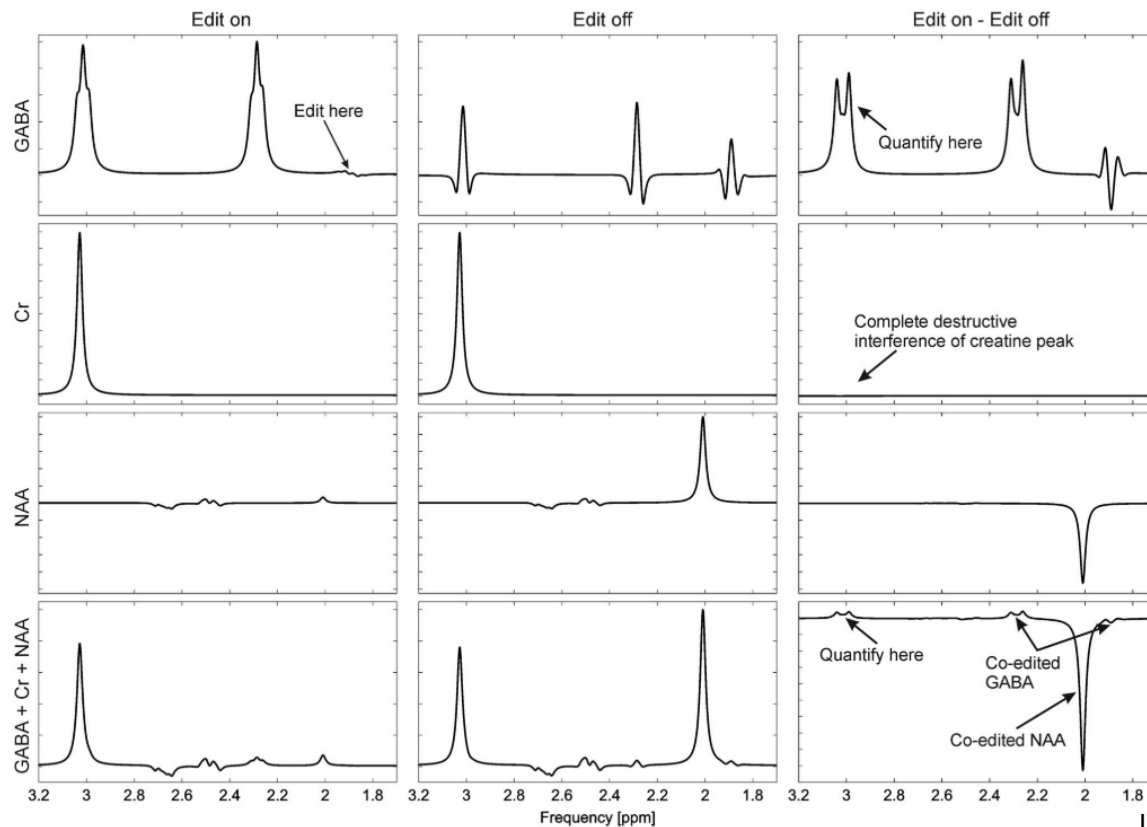


- **ME**sher and **GA**rwood (first and last author of first publication about this topic)



- Necessary to quantify single metabolites out of overlapping peaks
→ glutamate, GABA, glutamine

2.1 MEGA

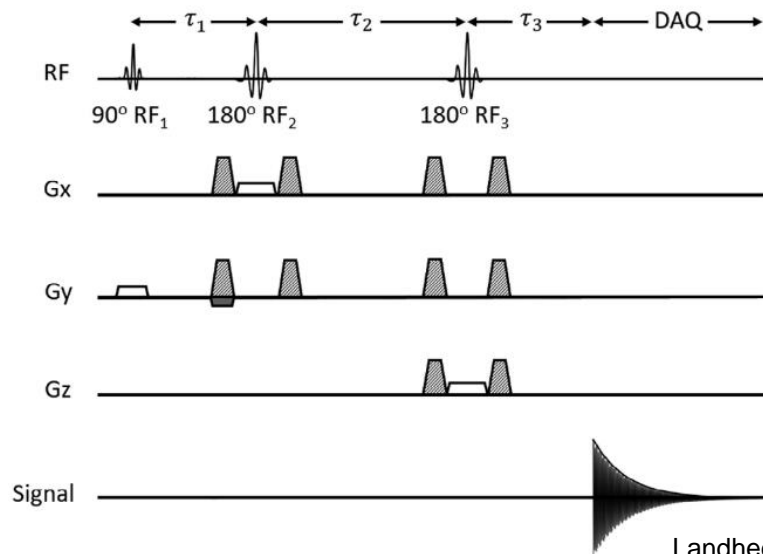


2.1 MEGA

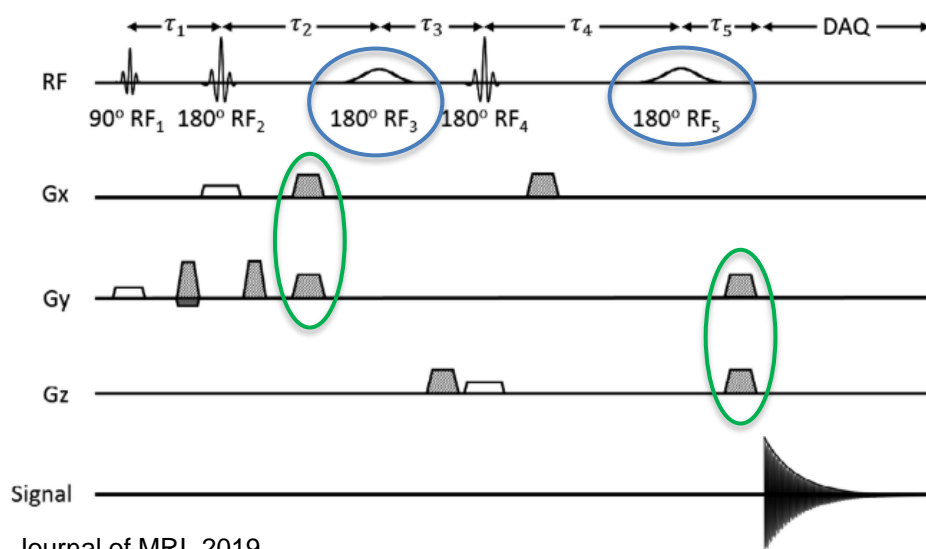


- two frequency-selective pulses designed to invert and, with the use of crushers, dephase unwanted resonances while not affecting desired resonances

PRESS



MEGA-PRESS

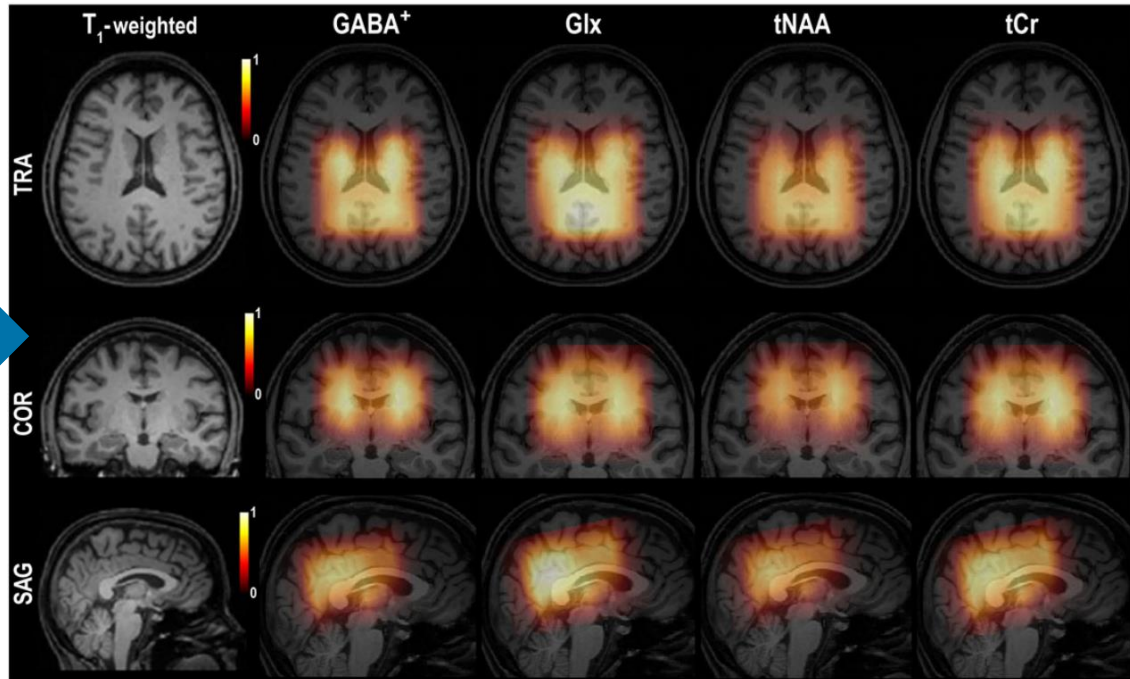
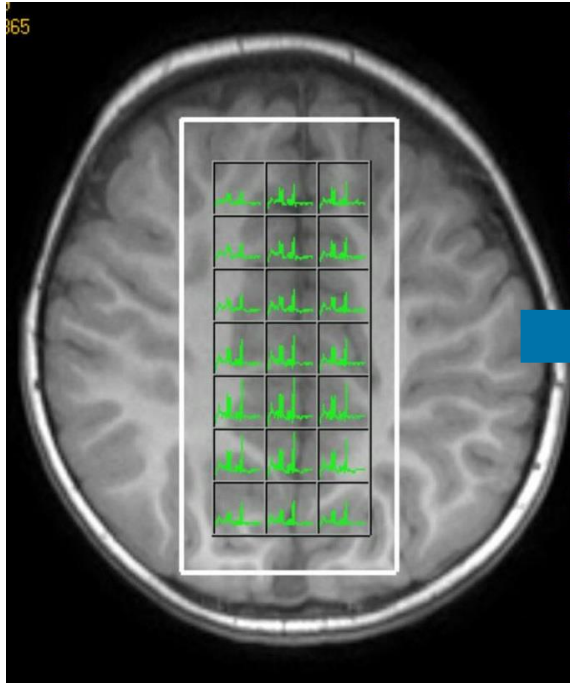


Landheer et al., Journal of MRI, 2019

2.2 MRSI



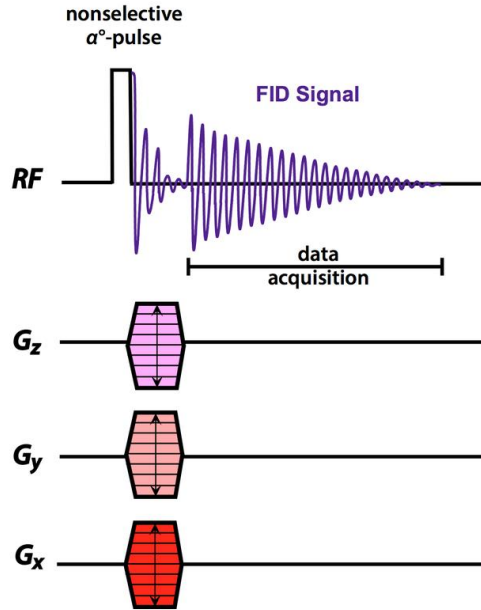
- Magnetic Resonance Spectroscopic Imaging or Chemical Shift Imaging (CSI, old)



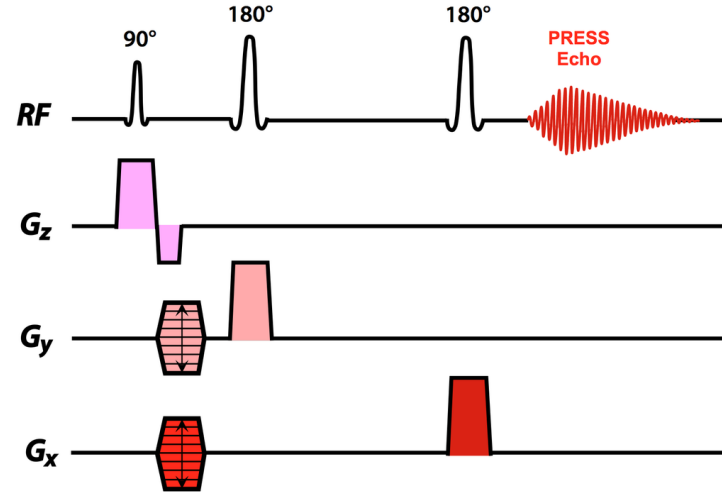
<https://www.mriquestions.com/csi.html>

Hniliová et al., NMR in Biomedicine, 2016

2.2 MRSI



3D MRSI sequence using nonselective volume excitation and stepped phase-encoding gradients along all three axes
→ ^{31}P



2D-PRESS MRSI sequence with slice-selective excitation pulses in three planes with stepped phase-encoding gradients along 2 axes → ^1H brain

<https://www.mriquestions.com/csi.html>

2.2 MRSI



Advantages:

- Spatial distribution of measured signal

Disadvantages:

- Larger point spread function → increased signal bleeding
- B_0 Shim not optimized
- Long measurement time



It is QUIZ time



<https://www.menti.com/95vrtaxcce>

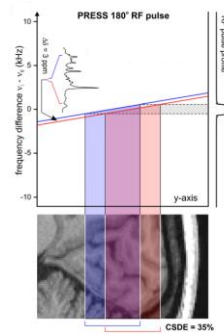
Menti.com, Code: 2800 3789

Take Home Messages



- ,classical' SVS MRS sequences without adiabatic pulses
- ,modern' ones with adiabatic pulses and combination with ,classical' sequences

- Trade-off between SAR , CSD and TE/measurement time



- MEGA technique → increasing metabolic specificity J-Difference editing
- MRSI if spatial distribution of the metabolite signal required, while SVS easier to acquire and easier to correctly quantify

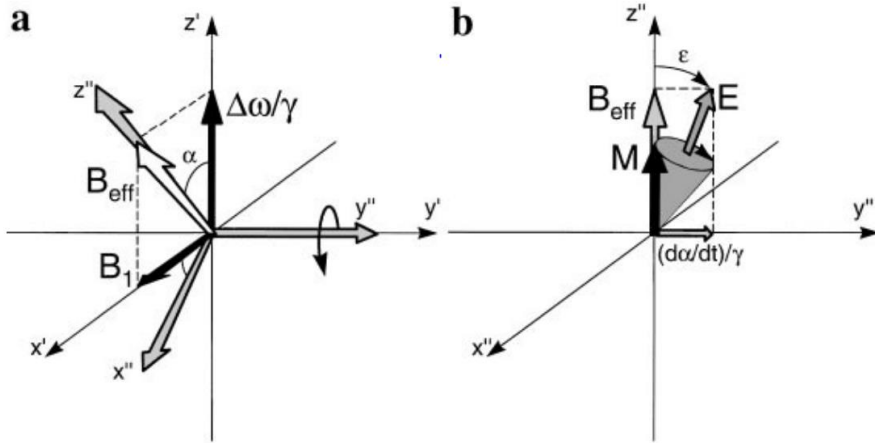
Thank you!



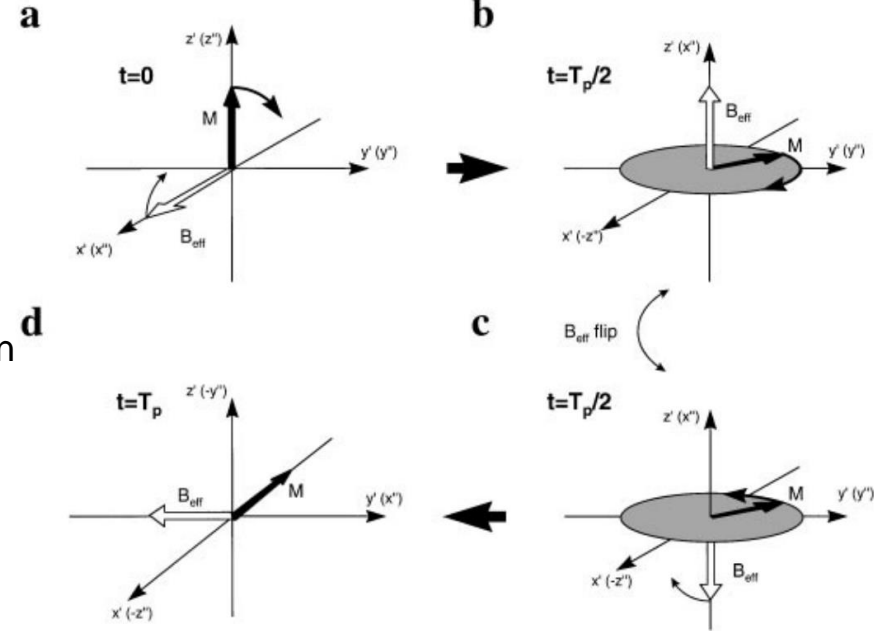
Layla.riemann@ptb.de

<https://www.dreamstime.com/illustration/cartoon-questions.html>

More adiabatic pulses (yippie 😊)



- Produces a nearly perfect inversion of net magnetization (**M**) that is relatively tolerant to **B**₁-field inhomogeneities
- Continuous RF-excitation swept over a range of frequencies from far below to far above the resonance frequency
- Provided the **B**₁-field was strong enough and applied slowly enough (the **adiabatic condition**), the net magnetization (**M**) could be nutated with a complete inversion by the end of the sweep



A closer look on stimulated echos

