Designing Protocols for MR imaging and Spectroscopy

Benoit Boulat BIC – April 2005

Examples taken from the project proposed by and worked out with

P.T. NARASIMHAN

To combine imaging and spectroscopy in the study of the mouse brain using intermolecular multiple quantum coherences.

ACKNOLEDGEMENTS

Tim Hiltner
Xiaowei Zhang

Russ Jacobs Scott Fraser

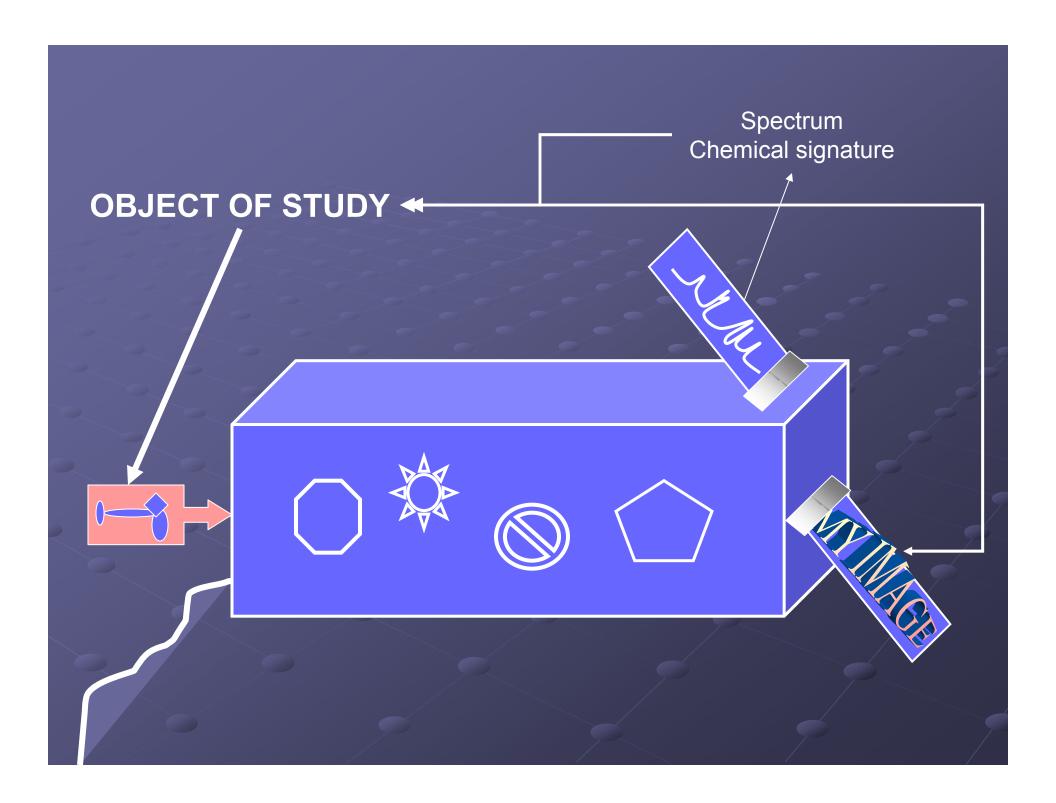
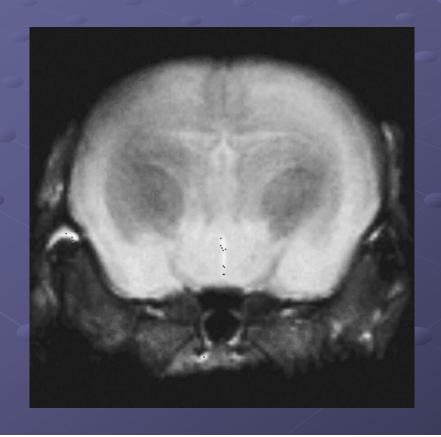


IMAGE -> CONTRAST

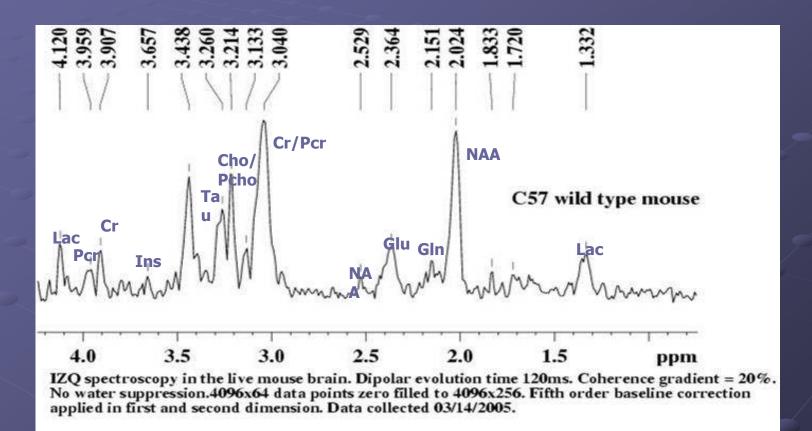






TO CARE FOR OR TO DESIGN IMAGING PROTOCOLS YOU NEED TO KNOW THE DETAILS OF THE TECHNIQUE THAT GIVES RISE TO THE CONTRAST(S).

SPECTRUM -> FREQUENCIES AND AMPLITUDES



YOU NEED TO KNOW THE DETAILS OF THE TECHNIQUE THAT GIVES RISE TO THE SPECTRUM.

SCHEMATIC OF THE BRUKER MR IMAGER

WORKSTATION

SOFTWARE CONTROL AND DEVELOPMENT

VISUALIZATION

VISUAL INTERPRETATION

SPECTROMETER

RF GENERATION AND RECEIVING

DIGITIZATION

MAGNET AND PROBES

SPATIO-TEMPORAL DISTRIBUTION OF MAGNETIC FIELDS

SPIN DYNAMICS

W

WORKSTATION (W)

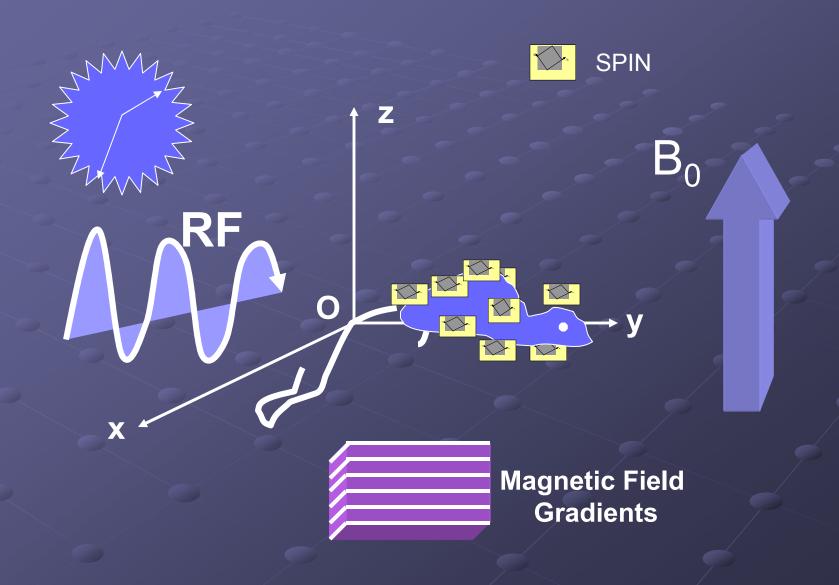
Software Control And Development MAGNET PROBES AND SPINS (MPS)

Spatio-temporal Distribution of Magnetic fields

Spin dynamics

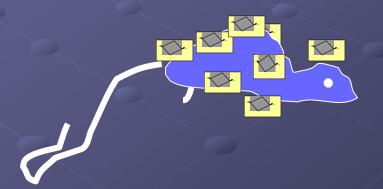
MPS

MAGNETIC RESONANCE

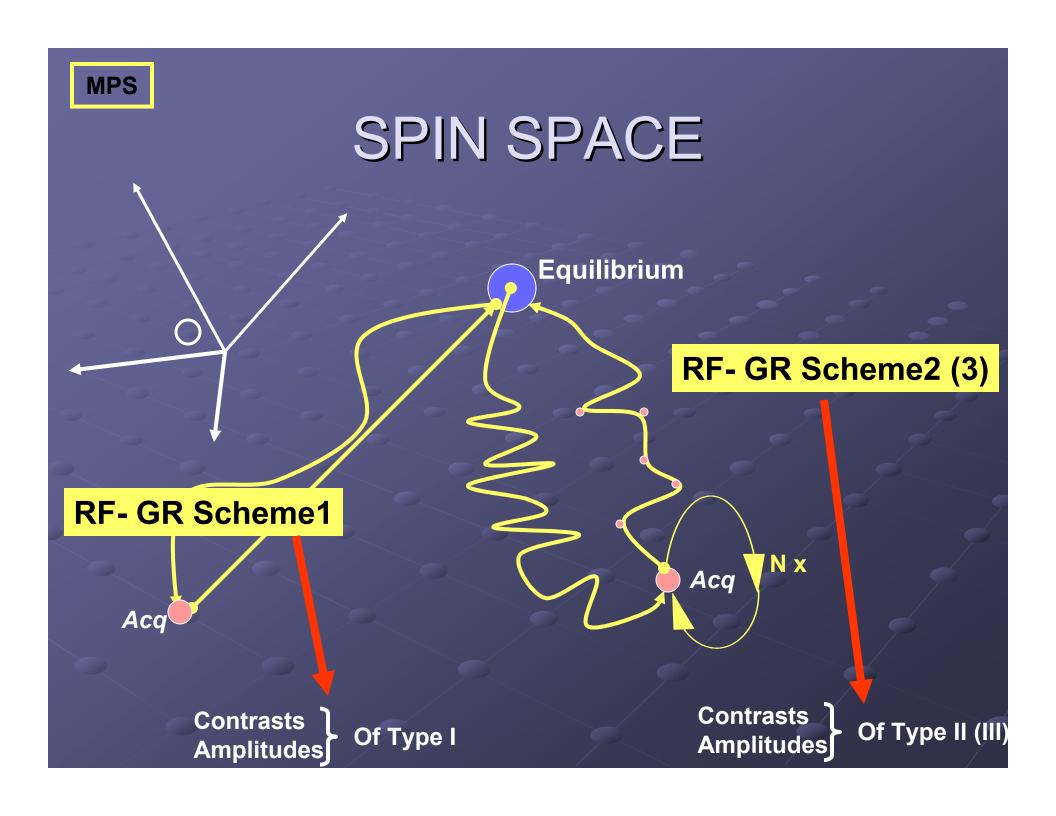


SPIN DYNAMICS

Ultimately the dynamics imposed on the nuclear spins by the rf and the gradients will be reflected in the contrasts of the image or in the amplitudes of the resonance lines in the spectrum.



At this point the precise material nature or shape of the object of study does not count much for us. Our objective is to precisely control the nuclear spins it carries.

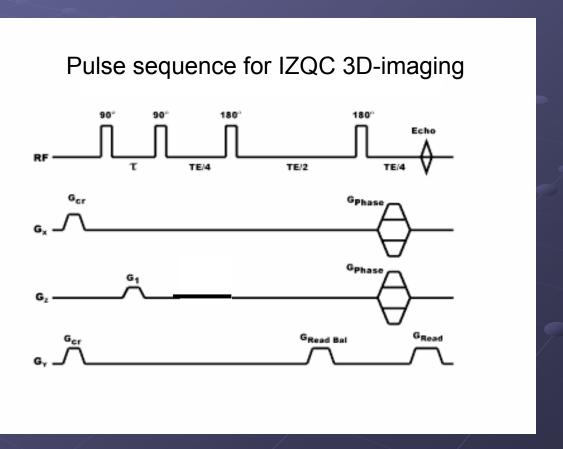


The spin manipulations are contained in the pulse sequence which is a summary of the procedures to be carried out to run the experiment. At this point the jargon used should relate to some model of spin dynamics the experimenter has in mind.

- Ideally we want to build a software tool that will initialize a maximum of these values with the aim of allowing the method to be used by people who may not be experts in spin dynamics.
- We want to optimize the method with regard to the speed of acquisition, especially if applications to living systems are sought.

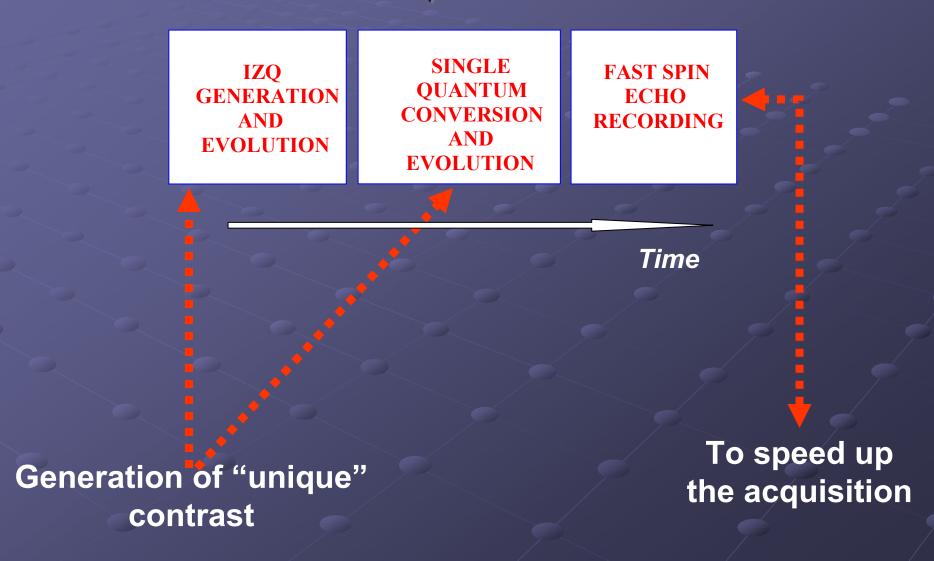
What we are given or have come up with:

```
sgrad r2d
start, 10u
slice, 10u
             fq8b:f1
           rgrad r2d
           groff
           fq1:f1
           ph11 ; excitation pulse
           grad\{(t10) | (t10) | (t10) \}; coherence filter on
     2m
                       ;coherence filter off
           groff
                      ;adjustment for zq
           ph12 ;45/135 degree pulse
     d30
           grad\{ (t4) | (0) | (0) \}
    d11
           groff
echo, d3
           grad{ (t15) | (t15) | (t15) }
    d5
           grad{ (t2) | (t2) | (t2) }
    p1:sp1 ph1
           grad{ (t15) | (t15) | (t15) }
     d4
           groff
```



Build a method to initialize the parameters

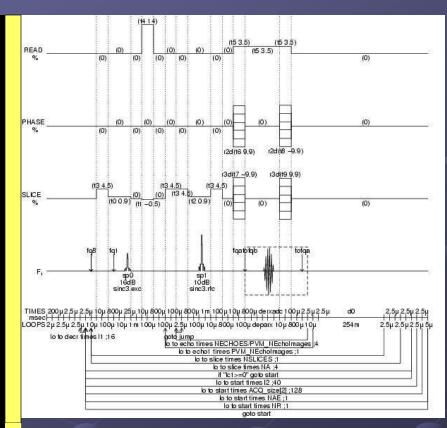
IZQ - FSE





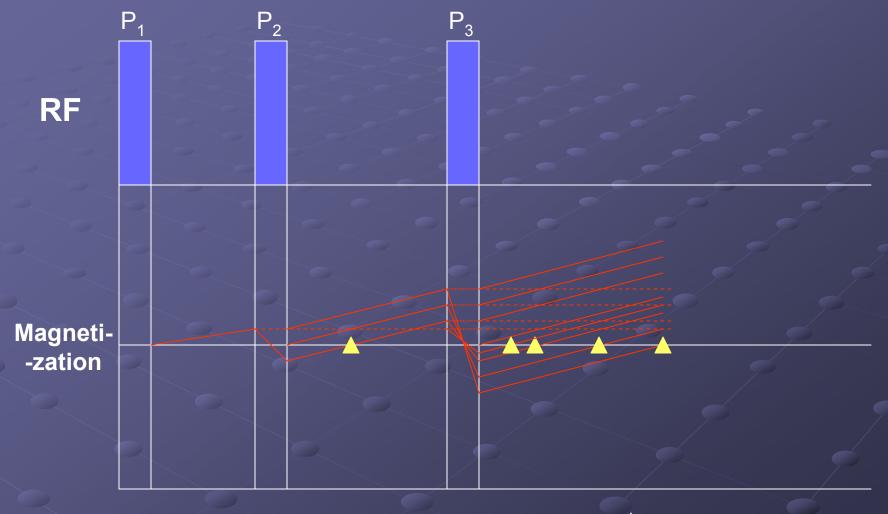
Fast Spin Echo (RARE)

```
slice, 10u fg8b:f1
      d4 grad{(0)|(0)|(t3)}
              -----slice selection-----
    d4 grad{(0)|(0)|(t0)}
    d3 fq1:f1
     d8 gatepulse 1
    p0:sp0
    -----slice rephase-----
    d4 grad{(t4)|(0)|(t1)}
    d11
             ----- refocusing group ------
echol, d4 grad{(0)|(0)|(t3)}
    rgrad r2d
    goto jump
echo, d4
          grad{(0)|(0)|(t3)}
jump, d5
            grad{(0)|(0)|(t2)} gatepulse 1
      d3
      p1:sp1 ph1
      d3
      d4
            grad{(0)|(0)|(t3)}
               ----- refocusing group - end ------
   -----read on + phase encoding----
if( ACQI cur dim == 1 )
                                    ; no phase encoding
      d4 grad{(t5)|(0)|(0)}
if( ACQI cur dim == 2 )
    d4 = grad\{(t5) | r2d(t6) | (0)\}; 2nd dim only
if ( ACQI cur dim == 3 )
         grad\{(t5)|r2d(t6)|r3d(t7)\}; 2nd and 3rd dims
    ADC INIT B(ph0, ph1)
           ----read on + phase encoding----
if( ACQI cur dim == 1 )
      d4 grad{(t5)|(0)|(0)}
if( ACQI cur dim == 2 )
```



A method exists to initialize the parameters

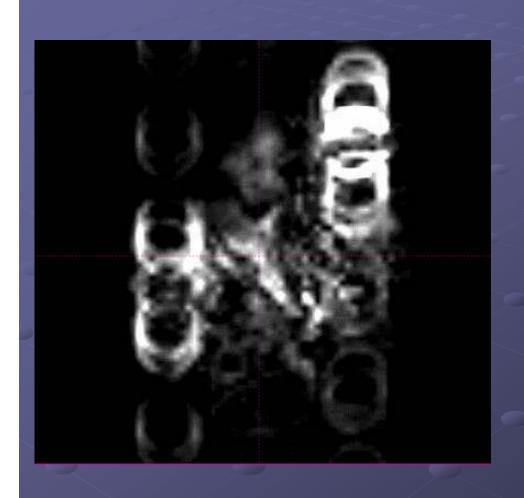
FAST SPIN ECHO: SOME DETAILS



▲: Echo



Fast Spin Echo





MODULE A (IZQ) MODULE
B
(FSE)



MODULE A + MODULE B

(IZQ-FSE)

ModuleA (IZQ) --- ModuleB (FSE)

Solutions found to solve problems of the stand alone module B do not necessarily work when some module A is placed in front of it, since the initial conditions to module B have been changed.

END of IZQ MODULE

MPS

FSE

M_L Amplitudes wound in a helix along B₀

 M_{T}

To preserve the "unique" contrast of IZQ we do not want to mix M_L and M_T inside FSE

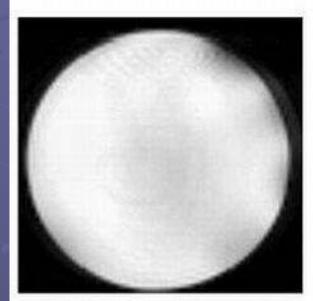
Need to work out a new solution for



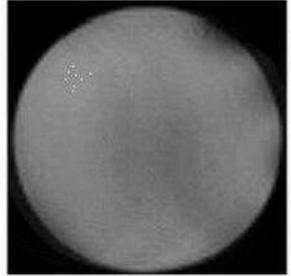
MODULE A + MODULE B



BENCHMARK TO TEST MODIFICATIONS



Gradient along BO



Gradient orthogonal to B0



Gradient at the magic angle relative to BO

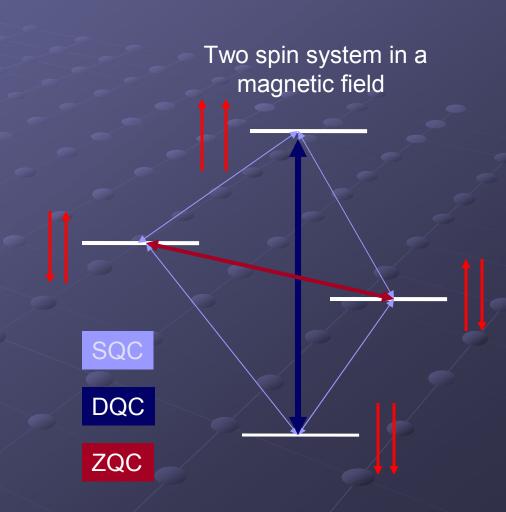
SPECTROSCOPY

- Spectroscopy can provide a wealth of information about living systems
- To obtain a good resolution of the spectral lines one needs to perform a good shimming of the main static field. Shimming is still pretty much an art. Only a few automated methods exist and they do not work in all situations.
- At best, shimming procedures yield a uniform field over small volumes only.
 Thus signal intensity is proportionately reduced.
- Field drift can be taken care of by a "field frequency lock" which is usually not available in imaging systems.
- "Block" averaging can mitigate the situation only partly and suffers other drawbacks.
- We explored the possibilities of implementing IZQ spectroscopy in our imaging system to overcome the shimming and drift problem.

Spin Systems

One spin System in a magnetic field

Single Quantum Coherence



Zero Quantum Coherences Spin A – Spin X ω_{A}

MPS

$$B_0 + \Delta B(R)$$

$$B_0 + \Delta B(P)$$

$$SQC_A: \omega_A + \delta\omega_A(P)$$

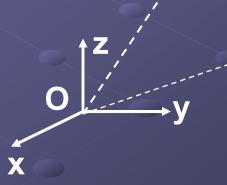
$$SQC_{x}: \omega_{x} + \delta\omega_{x}(P)$$

 ZQC_{AX} : $\omega_A - \omega_X$

$$SQC_A$$
: $\omega_A + \delta\omega_A(R)$

$$SQC_x: \omega_{x+} \delta\omega_x(R)$$

$$ZQC_{AX}$$
: $\omega_A - \omega_X$



ZQCs are insensitive to field inhomogeneities

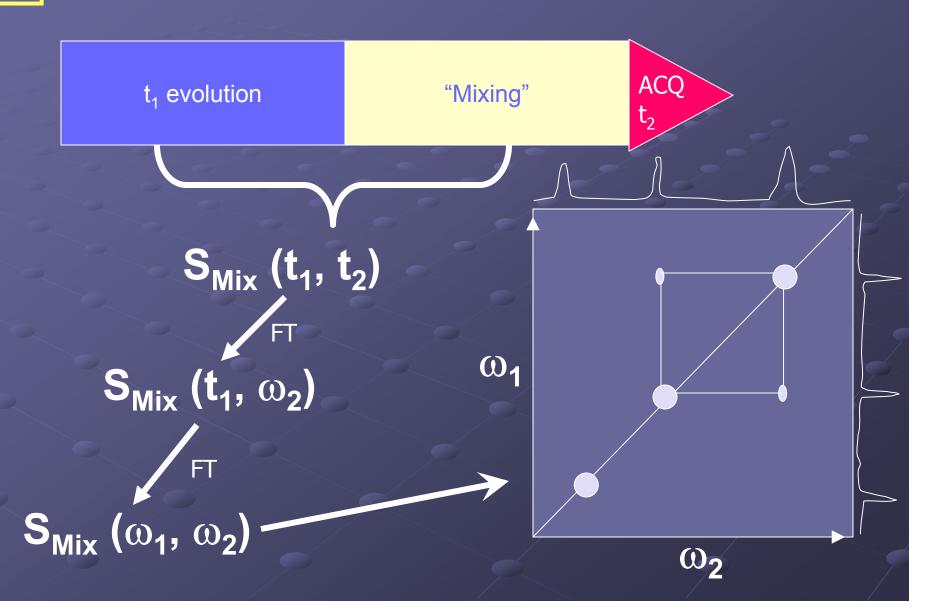
 One cannot directly detect through the RF coil coherences of order different from one.

 We therefore need to detect zero quantum coherence indirectly.

• We do this by using the framework of 2D spectroscopy, labeling zero quantum frequencies along the indirect dimension.



2D SPECTROSCOPY





2D IZQ SPECTROSCOPY

t₁ evolution IZQC

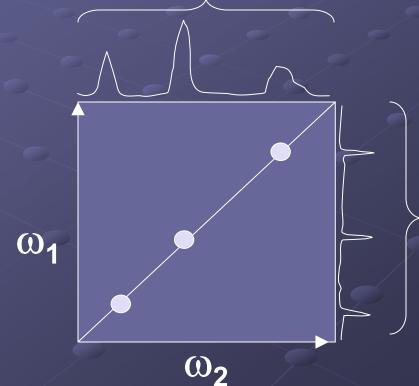
Evolution under the influence of distant dipolar field created by the spins of the solvent

ACQ

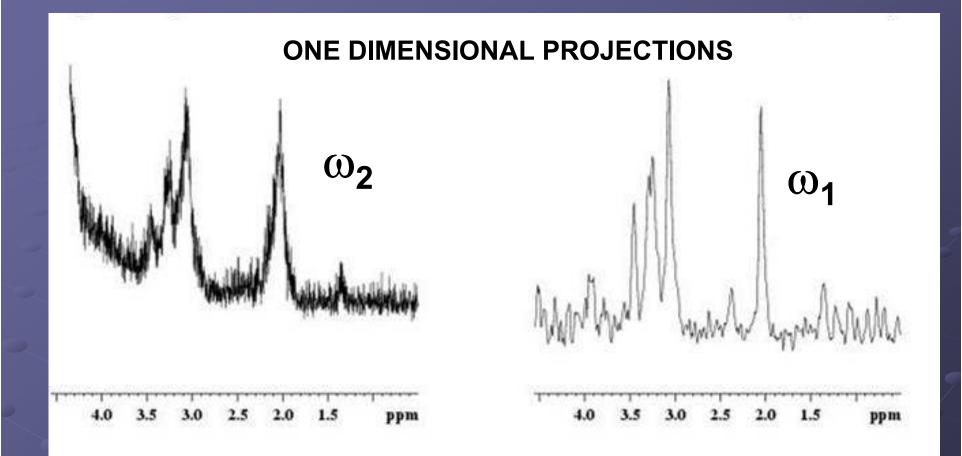
Insensitive to static magnetic field inhomogeneities





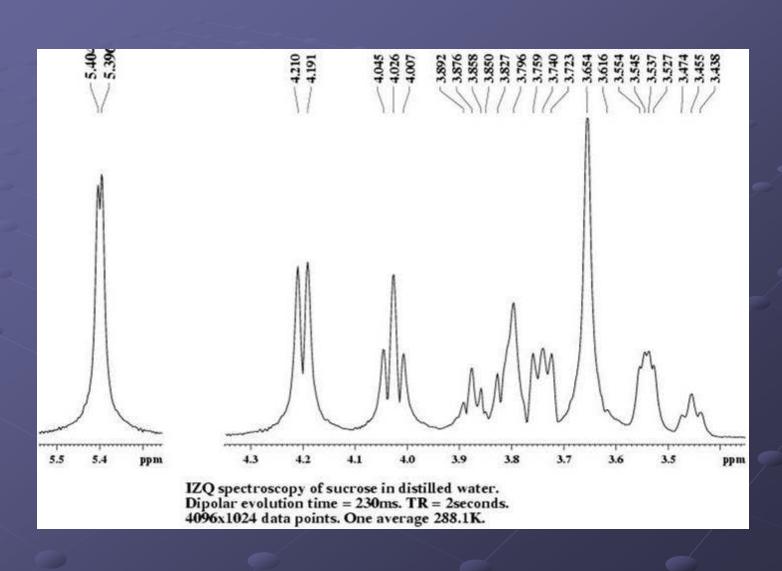


Narrow Lines



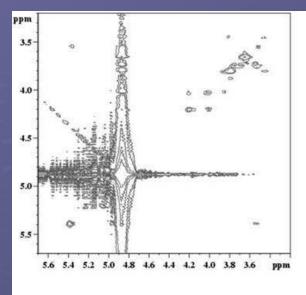
IZQ spectroscopy in the live mouse brain. Dipolar evolution time = 120ms. Relaxation delay = 5s. 4096x512 data points zero filed to 4096x1024. Micro2.5. 20mm birdcage coil. Two averages per t1 increments.

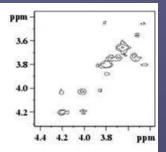
BENCHMARK TEST I



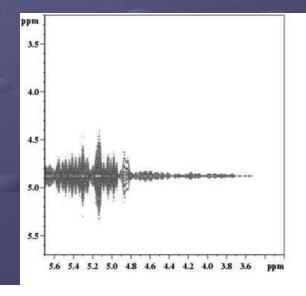


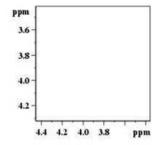
BENCHMARK TEST II



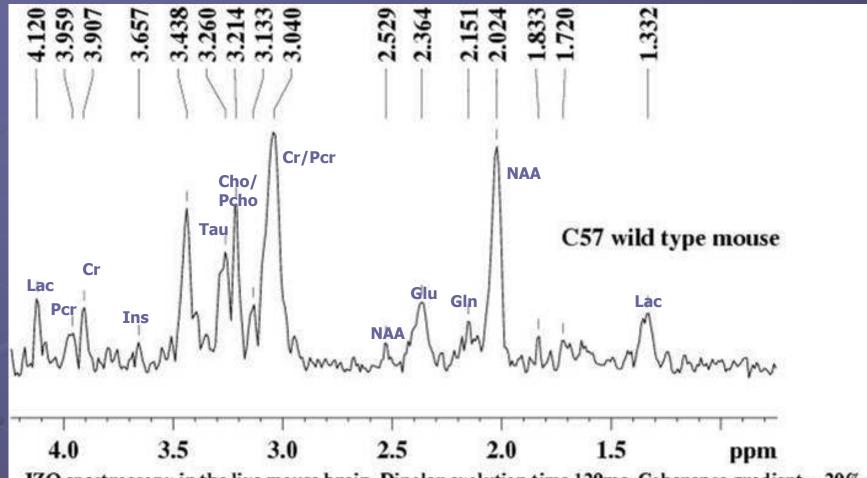


IZQ_COSY of sucrose in distilled water. 1024x300 data points. Dipolar evolution time = 60ms. 288.1K. Double quantum filtered with gradient selection prior to acquisition.





Coherence gradient at the Magic angle



IZQ spectroscopy in the live mouse brain. Dipolar evolution time 120ms. Coherence gradient = 20%. No water suppression.4096x64 data points zero filled to 4096x256. Fifth order baseline correction applied in first and second dimension. Data collected 03/14/2005.