Introduction to Magnetic Resonance - a general overview

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# Nuclear spin

- Spin: basic property of particles
	- Angular momentum and magnetic dipole moment
- Discrete quantum states
- In MR, spins are flipped between the two states
- Photons with frequency proportional to  $\Delta E$ will be emitted





## Equilibrium, excitation, signal

• In equilibrium, distribution of spins given by the Boltzmann equation

$$
\frac{N_{down}}{N_{up}} = e^{-\frac{\Delta E}{k_B T}}
$$

• At 3T:  $\frac{\Delta N}{N}$  $N_{\boldsymbol{u} \boldsymbol{p}}$ +N $_{down}$  $= 10$  ppm

- RF fields can flip spins between the two states
- After excitation, magnetization will oscillate and can induce current in receive coil



## Description with classical physics



## Repeated vs single excitation

Magnetization: sum of all magnetic dipoles



Signal sampling

#### Precession frequency and angles



$$
Signal = sin(\alpha)
$$
  
Remaining  $M_z = cos(\alpha)$ 

$$
Precession frequency f = \frac{\gamma}{2\pi} B_0
$$

Precession phase: given by  $f$  duration and RF phase

$$
\varphi = \varphi_0 + f \cdot t
$$

Often both *x*- and *y*-projections are sampled

## Spectroscopy





- Electrons shields nuclei slightly
	- Chemical shielding
- Frequency at nuclei determined by specific electronic structure
- Larmor relation  $f = \frac{\gamma}{2\pi}B_0$
- Sampling of temporal signal variation
	- Fourier Transformation
	- Signal on frequency axis
- Signal amplitude given by molecular concentration

## Principles of MR imaging

• Encoding spatial information in MR signal



## Frequency encoding



## 3 dimensions and k-space



K-space

- Fourier inverse to normal space
- Images can be decomposed to weighted sum of spatial frequencies
- Each spatial frequency symbolized by point in k-space
- In MR, the amplitude of spatial frequencies are sampled
	- Spatially dephased magnetization corresponds to one spatial frequency





- Signal is sampled in k-space
- Using gradients to move around in k-space
- After having sampled the entire k-space, images can be reconstructed by 2D inverse Fourier Transformation

#### Different ways to sample k-space



## K-space filtering



Image **K-space** 

## Frequency + spatial information



- Frequency adds one extra information dimension
	- Time consuming
	- Lower spatial resolution



# Signal amplitude and noise

- Signal amplitude given by
	- Density of nuclei
		- Proton density is high
	- Gyromagnetic ratio  $\gamma$ 
		- Signal  $\propto \gamma^3$
	- Polarization
		- For standard MR around 10 ppm
- Noise
	- Mainly from patient
- $SNR = \frac{signal}{noise}$ noise
- Signal on scanner is not absolute

### Functional MR methods

- Static or dynamic imaging
	- Various contrast choices: T1w, T2w, T2\*w, etc
- Functional MRI methods
	- Diffusion weighted MRI
	- Blood perfusion measurements
	- Phase contrast flow measurements
	- BOLD based fMRI
	- Cell metabolism with hyperpolarized bioprobes

## Diffusion weighted MRI



Free diffusion: Water molecular motion only limited by collision with each other



Restricted diffusion: In biological tissue. Water molecular motion limited by collisions with cells and macromolecules

#### Diffusion measurement in stroke

Acute 2 hours 24 hours 24 hours



Infarction area

## Assessment of anisotropic diffusion



- Can also use more advanced models
	- Neurite density
	- Neurite orientation dispersion index
	- Kurtosis
	- IVIM

#### White matter tractography example



### Blood perfusion measurements





- Injection of contrast agent
	- DSC, DCE
- Labelling arterial blood
	- ASL
- Quantitative parameters obtained using models
- Hyperpolarized MR
	- Urea, pyruvate

## fMRI



- Mapping of brain activity
- Based on blood flow variations
- BOLD: differences in T2\* of oxy- and de-oxy blood
- Activation or resting state based
- Low signal differences
	- Multiple averages and statistical measures needed

#### Phase contrast, vessel blood flow



- Blood flow in vessels
- Measuring global cardiac function
- Assessing heart valve integrity and vessel stenoses



Mean flow<sub>vessel</sub> = 
$$
\frac{\sum_{frames} Flow_{vessel}}{N_{frames}}
$$

$$
Stroke\ vol = \frac{Mean\ flow}{Cycle\ time}
$$

# Hyperpolarized MR

- MR molecular imaging
- In vivo study of various metabolic pathways
- $[1-13C]$  pyruvate
	- Glycolysis, aerobic vs an-aerobic metabolism, pH
	- Cancer probe, monitor of therapy response
- $[2^{-13}C]$  pyruvate
	- Study of TCA cycle pathways
- $[1,4$ -<sup>13</sup>C<sub>2</sub>] Fumarate
	- Detection of cell necrosis
- <sup>129</sup>Xe Xenon gas
	- Assessment of lung function

## History of MR

- ~1943: Spin and interaction with radiofrequency fields discovered
- ~1973: Paul Lauterbur made the first MR images
- 1980 ->: MR being used for clinical non-invasive imaging
- 1953: Overhauser predicted the possible manipulation of nuclear Boltzmann distribution transferring polarization from electrons
	- Initially doubted by Bloch, Rabi, Ramsey etc
- 1953: Experimental verification by Carver and Slichter
- 1994: Hyperpolarized Helium and Xenon introduced for lung imaging
- 2013: Ardenkjær introduces dDNP and showed a factor >10000 signal enhancement for 13C

## Summary

- Traditional MR
	- Frequency of signal: information about chemical structure
	- Spatial information: enables imaging
	- Allows measurements with different contrasts and functional measurements
	- Signal is low: some methods on the edge of useful sensitivity
- Hyperpolarization
	- 10000 to 50000 fold signal enhancement
	- Enables new measures of metabolism
	- Much larger signal but sensitivity still an issue