

MRI in the wild

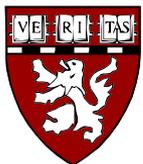
Matt Rosen, Ph.D

MGH/A.A. Martinos Center/Harvard University

Novel Methods for Phytosequestration

ARPA-E Workshop

Chicago, 23 July 2015

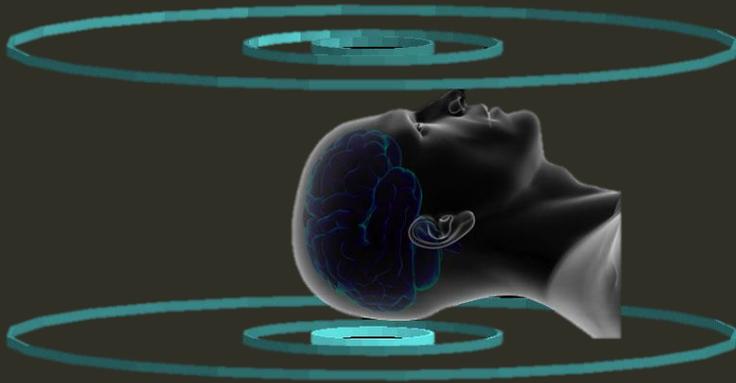


MGH/HST Athinoula A. Martinos
Center for Biomedical Imaging



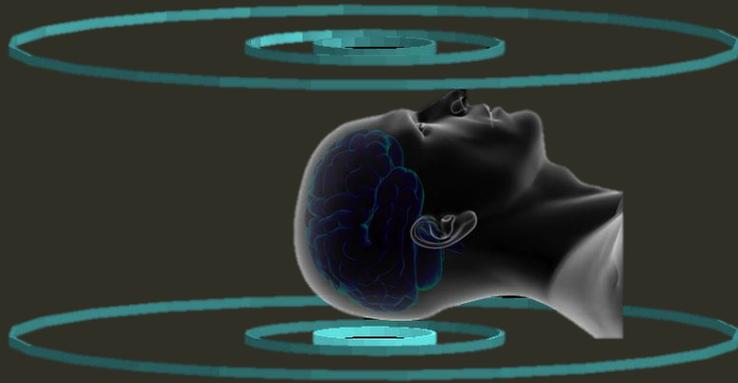
“Wild” MRI?

High-field MRI instruments are of *limited* utility in widely deployable contexts



“High-performance” ultra-low field MRI?

What would high-performance ULF MRI look like?



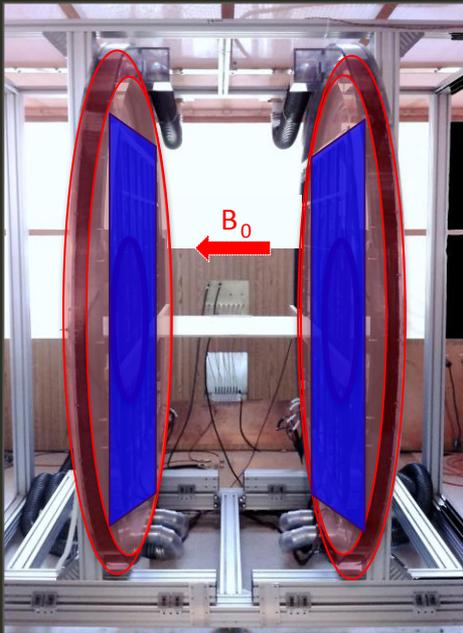
“High-performance” ultra-low field MRI?

What would high-performance ULF MRI look like?



MGH Connectome scanner : 24 MW
Los Angeles-class nuclear submarine

Ultra-low field MRI?



Custom made
Electromagnet
+
Gradients

6.5 mT

\$200k test-bed system



1.5 T – 3T

Cost: \approx \$1 million / Tesla

Weight: 7.3 tons

Cooling: Liquid cryogenics / **Availability?**

Shielding: RF/magnetic

Min. room size: 31m² ++

Ultra-low field MRI?

3 T



$\times 46^{3/2}$

0 SNR
($\times 10\,000$)

VS

0.0065 T



Acquisition time: seconds, minutes...

2005–2011



2D Gradient echo – 1 slice – acquired at ULF

Acq. time = 52 min / Voxel size = (3.9 x 7.8 x 15) mm³

Fast acquisition strategy for ULF MRI

Balanced Steady State Free Precession (b-SSFP):

- Fast & efficient but demanding
- Refocused quantum control-based acquisition

PHYSICAL REVIEW

VOLUME 112, NUMBER 5

DECEMBER 1, 1958

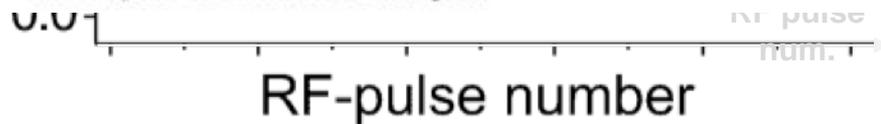
Steady-State Free Precession in Nuclear Magnetic Resonance*

H. Y. CARR

Department of Physics, Rutgers University, New Brunswick, New Jersey

(Received August 4, 1958)

A steady-state free precession technique for observing nuclear magnetic resonance is described. A mathematical analysis is presented for certain special conditions, and initial experiments verifying the results of this analysis are reported. This technique provides two opportunities for improving the signal-to-noise ratio. First, it provides a mechanism, similar to that of the "spin echo," for eliminating the effect of the inhomogeneity of the magnetic field on signal strength. This permits the effective use of larger samples. In the second place it provides a steady-state signal which can be observed with a narrow-band detector. Under certain conditions the technique has a broad response as a function of frequency or field. The upper limit to the width of this response is determined by the electronic apparatus supplying the rf pulses rather than the magnet or the nuclear sample.



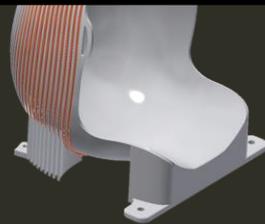
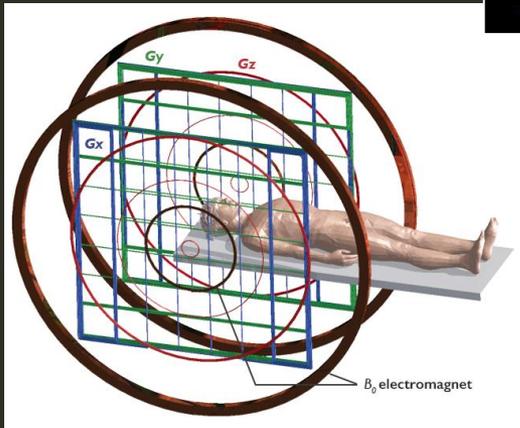
State of the Art: non-cryogenic ULF MRI (6.5 mT)

6 min,

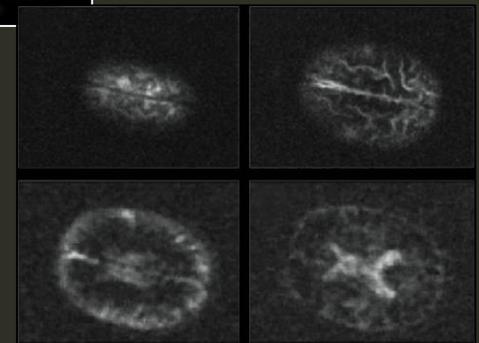
3.9 x 7.8 x 15 mm³
52 minutes, 2D GE

75x15

LANL 2015



Custom-fit single channel
spiral head coil (276 kHz)

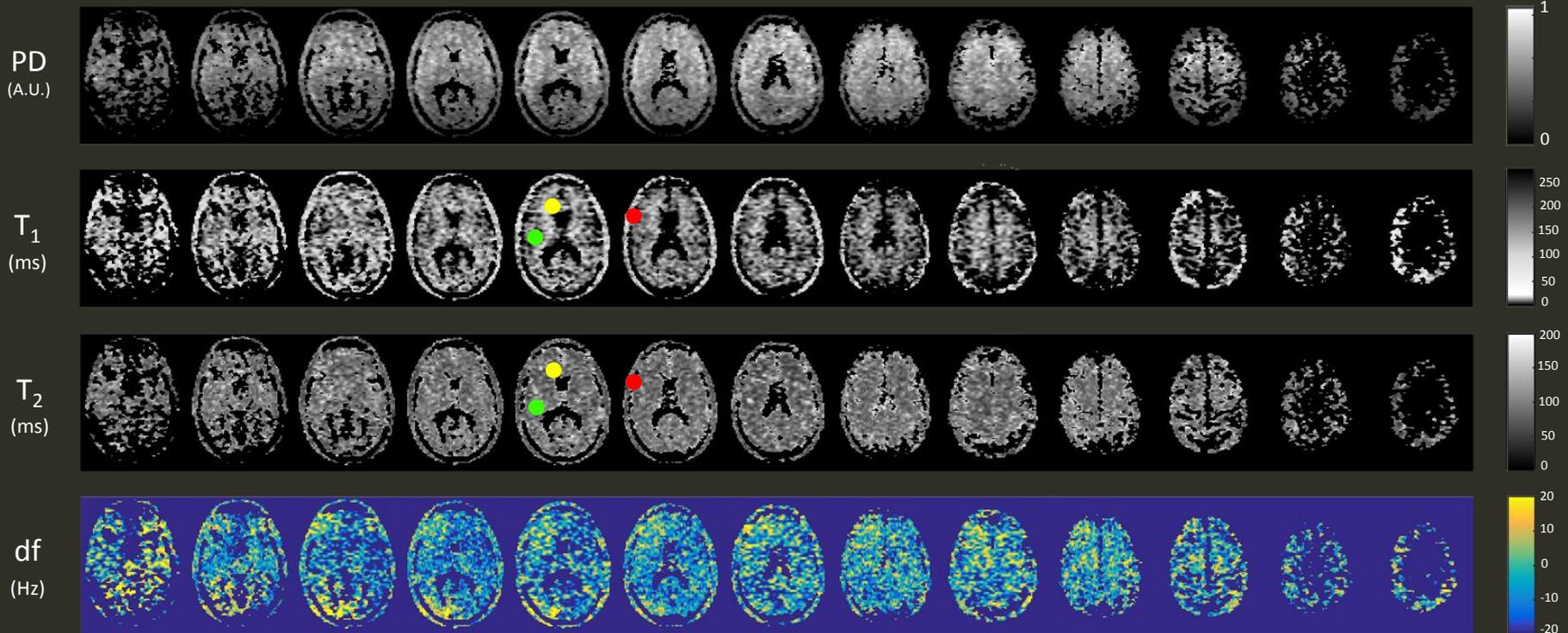


2 x 2.4 x 15 mm³ (5 slices)

67 min

3D MR fingerprinting *in vivo* at 6.5 mT

Stochastic Bloch trajectory-based imaging:



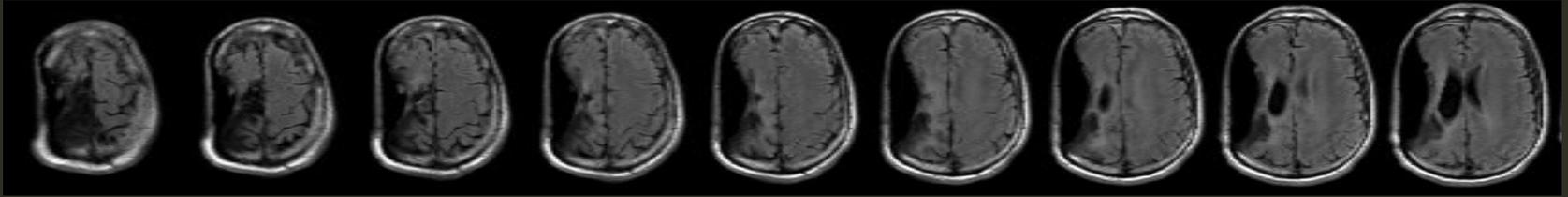
Optimized 20 point trajectory
3x3x10 mm³
NA: 1
Total acquisition time: 14min

	T ₁	T ₂
● Gray matter	172ms	85ms
● White matter	127ms	76ms
● Scalp	91ms	69ms

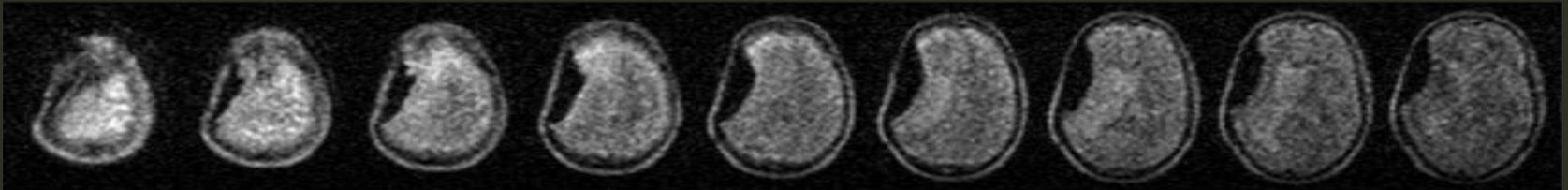
Quantitative image contrast!

Non-cryogenic ULF MRI (6.5 mT): Ultra-low field vs. high field in **clinical TBI**

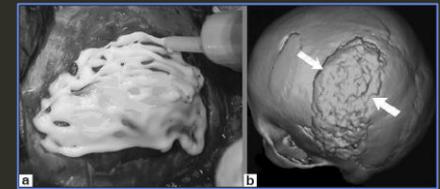
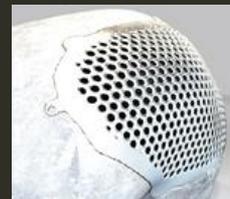
3 T



6.5 mT



Traumatic skull cranioplasty w/ **titanium mesh** & **MMA implant**



High field: T2 FLAIR

ULF: 3D b-SSFP, 3x3x8.5 mm, 15 slices

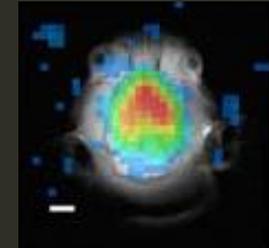
NA=140, $\alpha=84^\circ$, Matrix: 64x75x15, 28 min.

What do we measure in MRI?

- Nuclear (Boltzmann) polarization: $P \sim \mu B / k_B T$
- Magnetization per unit volume $\sim \rho \mu P$

Direct detection:

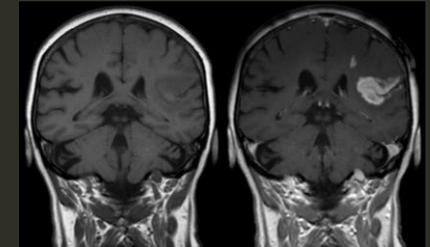
- Liquids, *solids* (thermally polarized)
- ...& gases (non-Boltzmann “hyperpolarization”)
- Not just ^1H !
- ^3He , ^{129}Xe , ^{13}C , ^{23}Na , ^{29}Si



Rosen et. al 1997

Indirect detection:

- Paramagnetic things (Gd) via nuclear relaxation
- **Free radicals** (Bohr magnetons) via polarization transfer (DNP)



BBB leakage in stroke

Physics: Overhauser techniques transfer electron polarization into long-lived nuclear polarization

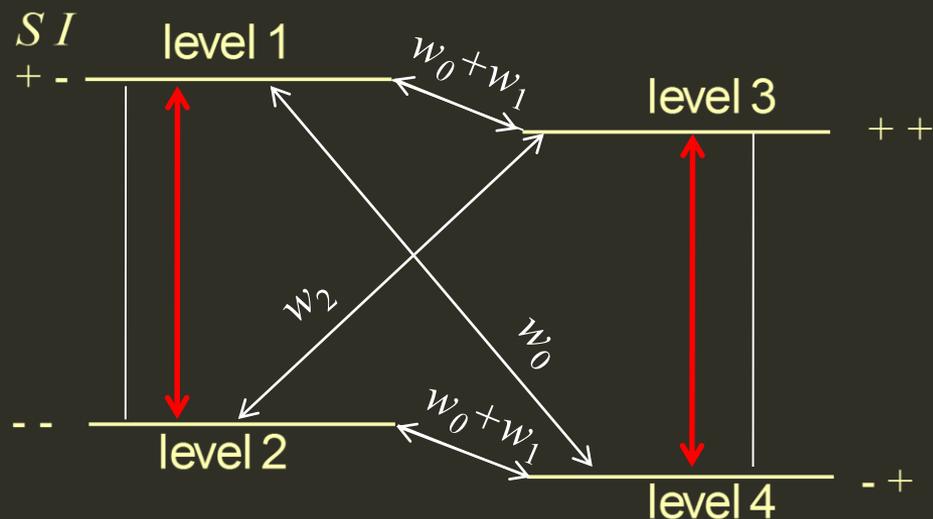
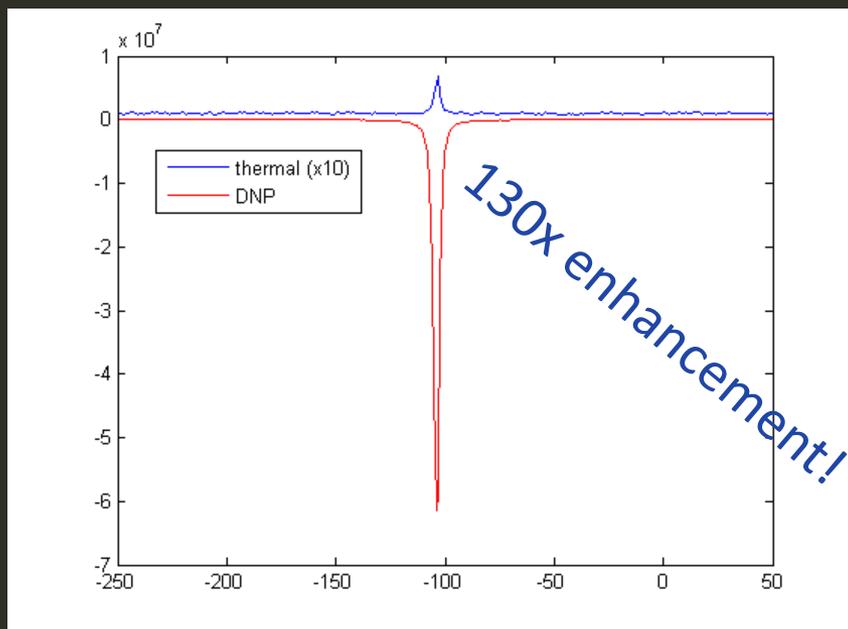
The Overhauser effect

$$H = \underbrace{g_S (\mathbf{S} \times \mathbf{H}_0) + g_I (\mathbf{I} \times \mathbf{H}_0)}_{\text{Zeeman interaction}} + g_S g_I \underbrace{\left[\frac{\hbar}{3} |\gamma(0)|^2 (\mathbf{I} \times \mathbf{S}) - \frac{3(\mathbf{I} \times \mathbf{r})(\mathbf{S} \times \mathbf{r})}{r^5} + \frac{\mathbf{I} \times \mathbf{S}}{r^3} \right]}_{\text{Scalar Time dependent coupling} + \text{Dipolar}}$$

Zeeman interaction

Scalar
Time dependent coupling

Dipolar



Rates depend on dynamics, H_0 , & $[e^-]$

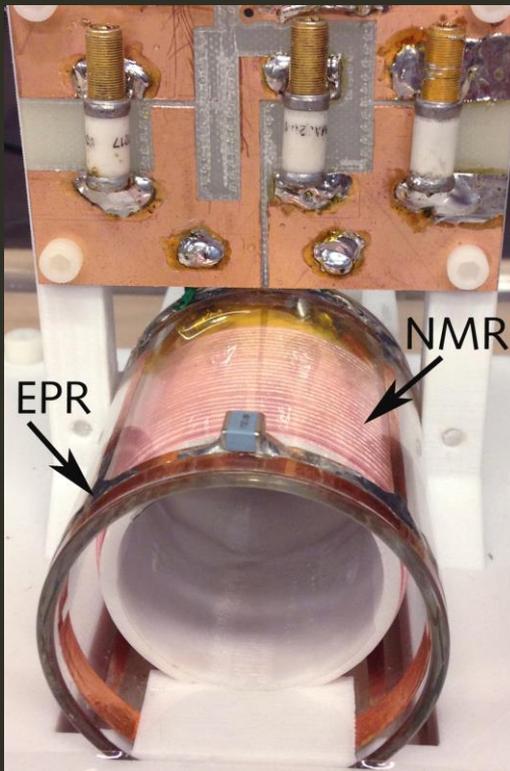
Hardware: free radical imaging at 6.5 mT

NMR (276 kHz)

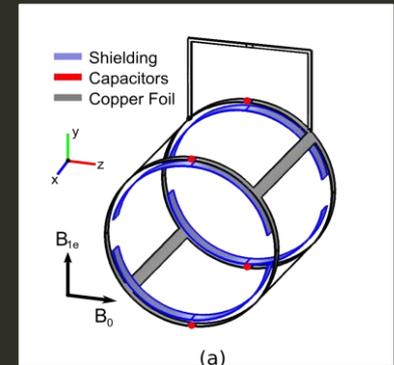
Johnson noise dominated

EPR (141 MHz)

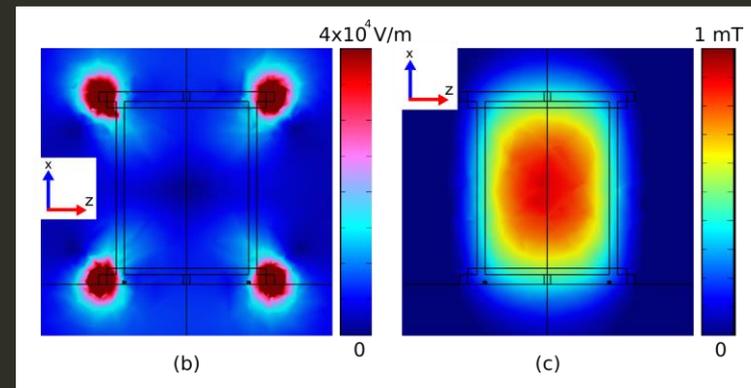
Modified A-G resonator



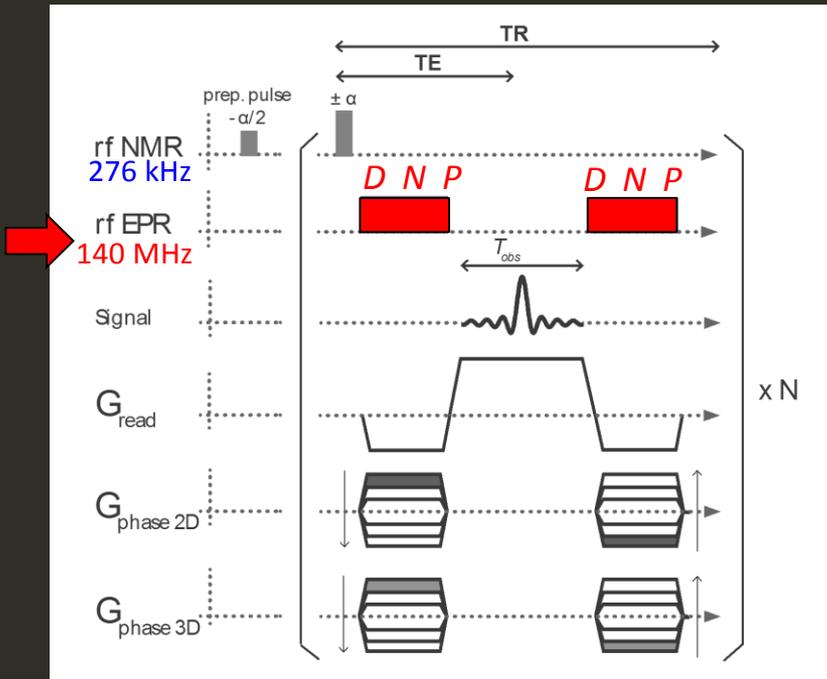
- High ESR homogeneity
- Suppressed E-field (SAR)



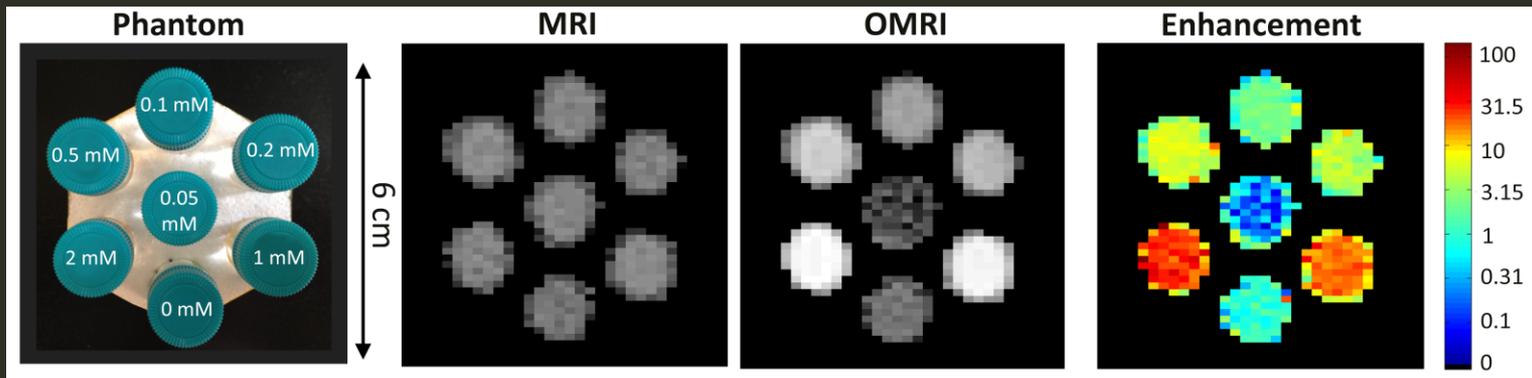
Litz wire
BW=3KHz
Q=92



High speed 3D free radical imaging



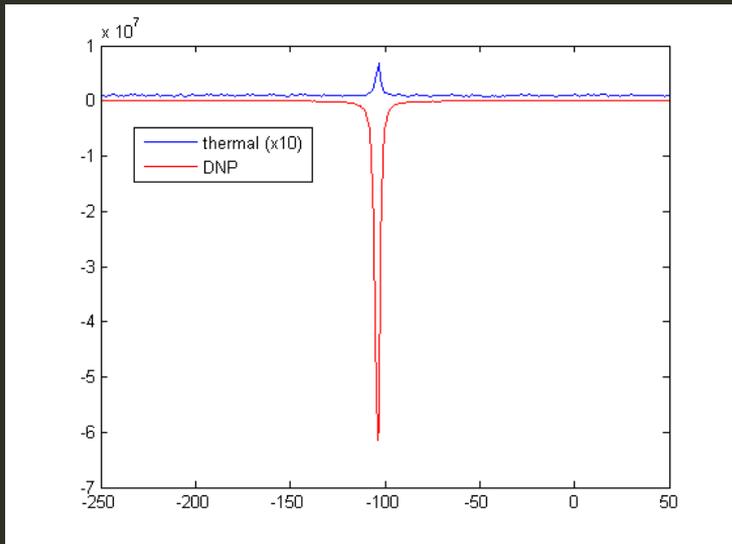
b-SSFP based OMRI
Embedded ESR pulse
No additional sequence time



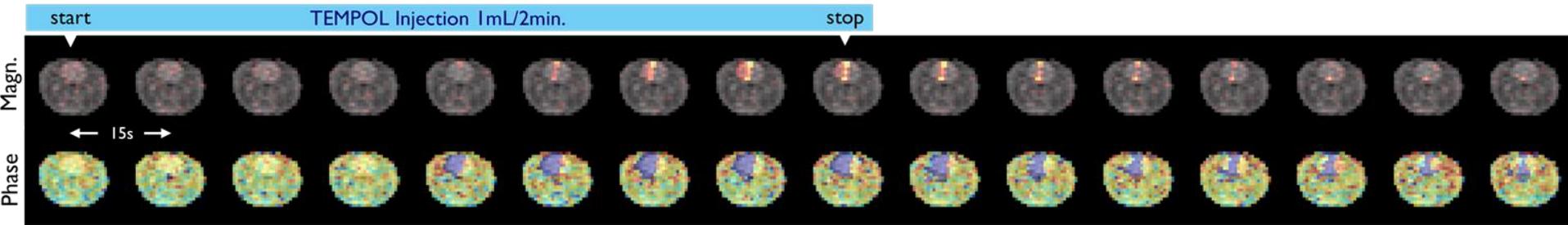
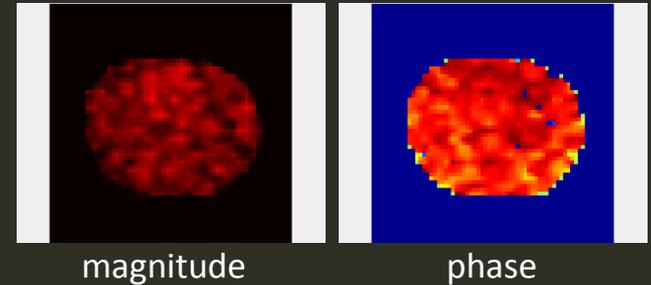
Images acquired at 6.5 mT: NMR: 276 kHz, ESR: 140 MHz, 2 mM TEMPOL in H₂O

Dynamic *in vivo* OMRI

Proof of concept in healthy rat



OMRI
Central slice in rat head – axial view



Matrix : 128 x 35 x 11

Resolution: 1.7 x 2.2 x 4.2 mm³

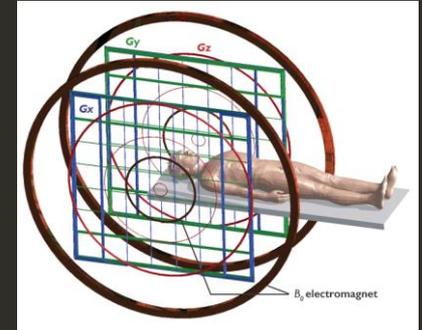
1 full 3D OMRI scan (11 slices) = 10 s + 5 s pause

16 dynamic scans: 4 min total imaging time

Carotid injection: 1 mL in 2 min, 150 mM TEMPOL

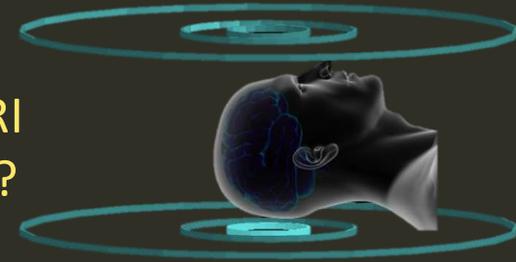
Conclusions

- MRI is possible outside the scanner suite
 - Not limited to brain imaging!
 - Not limited to existing scanner footprint
 - Understand impact of time/resolution tradeoff
 - Next revolution in health care? **Cost!**



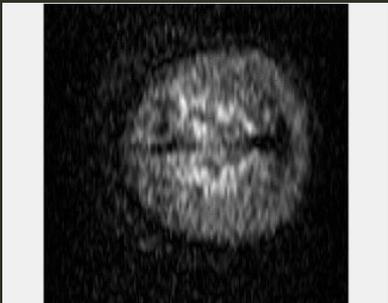
Eg. highly optimized tradeoff between resolution, speed, specificity, cost?

- Free radical sensitive MRI is possible **Stethoscope (\$50)**
 - Hyperpolarization via *in vivo* DNP & high speed ULF MRI
 - Free radical tracer agents? Redox/metabolic processes?



High-speed, inexpensive “medium resolution” MRI in the wild:

How would **you** use a tool like this?



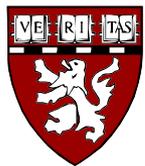
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Funding: DoD/DMRDP
NIH NINDS



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