

Parallel MRI Acceleration of Dynamic and High Resolution Hyperpolarized ^{13}C MRI

L Friesen Waldner^{1,2}, JX Wang⁵, A Chen⁶,
A Ouriadov¹, M Fox^{1,3}, B Rutt^{1,7}, T Scholl^{1,2},
G Santyr^{1,4}, C McKenzie^{1,2}

¹Robarts Research Institute, ²Medical Biophysics, ³Physics and
Astronomy, ⁴Medical Imaging, University of Western Ontario,
⁵Global Applied Science Laboratory, GE Healthcare,
⁶GE Healthcare, Toronto, Ontario, ⁷Diagnostic Radiology
and Richard M Lucas Center for Imaging,
Stanford University, California



Introduction

- Due to limited life times of pre-polarised spins ($T_1 \approx 1\text{min}$ for $[1-^{13}\text{C}]$ -pyruvate at 3T) imaging of hyperpolarized ^{13}C -enriched substrates requires fast imaging
- Traditional use of field gradients for spatial encoding is very slow
- Parallel MRI can acquire images faster with little to no loss in SNR
(Lee, et. Al. MRM 2006 **55**:1132)

Purpose

To demonstrate the feasibility and advantages of parallel MRI acceleration at high spatial and temporal resolutions after injection of a hyperpolarized ^{13}C -enriched substrate

Methods

Hyperpolarized Substrate Preparation

- [1- ^{13}C]-pyruvic acid (CIL, Cambridge MA) with 15mM OX63 trityl radical (Oxford Instruments, Abingdon UK) and 1mM Dotarem (Guerbet)
- Pyruvic acid mixture was polarized using a HyperSense DNP polarizer (Oxford Instruments, Abingdon UK)



Methods

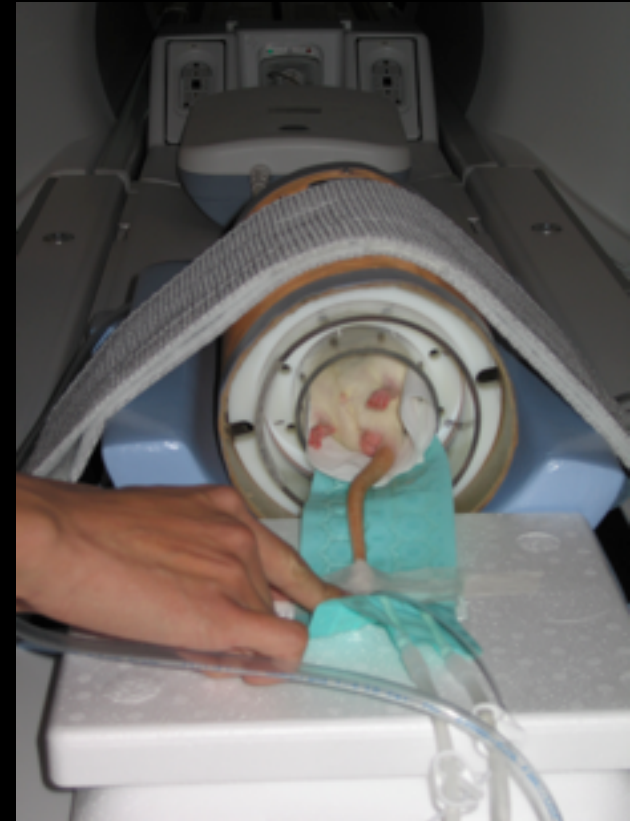
Animal Preparation

- Approval was obtained from the institutional Animal Use Subcommittee
- Normal Sprague-Dawley rats were anesthetized with isoflurane
- 2.5mL of hyperpolarized $[1-^{13}\text{C}]$ -pyruvate (80mM, pH=7.8) was injected into the tail vein over 13 seconds

Methods

RF Coils

- Custom dual tuned RF coil (^1H transmit/receive, ^{13}C transmit)
- Custom 8 coil array designed for ^{13}C at 3.0T (MR750, GE Healthcare, Waukesha, WI, see poster #1489)



Methods

^1H MR Imaging

- ^1H images were collected using a spoiled gradient sequence (SPGR)
TR=4.3ms, TE=1.5ms, FOV=200x200mm, 5mm slice thickness, 5mm slice spacing, 128x128 matrix

Methods

Dynamic ^{13}C MR Imaging

- Fully gradient-encoded datasets were collected beginning 8s after the start of $[1-^{13}\text{C}]$ -pyruvate injection.
- 10 coronal images collected at 2s intervals using 2D fast gradient recalled echo (FGRE) sequence
TR=10.7ms, TE=4.8ms, flip angle=5°, bandwidth= $\pm 4\text{KHz}$, FOV=200x200mm, 200mm slice thickness, 64x64 matrix, acquisition time=0.58s/image

Methods

Dynamic ^{13}C Image Reconstruction

- Twelve, fully sampled lines at centre of k-space were used to determine coil sensitivities
- Generalized encoding matrix reconstruction with:
Sub-sampling to simulate outer reduction factors (ORF) of 2, 3, and 4 and net reduction factors (R) of 1.7, 2.2 and 2.6 respectively

Methods

High Resolution ^{13}C MR Imaging

- Fully gradient encoded images collected at 8 and 15s post-injection of $[1-^{13}\text{C}]$ -pyruvate using 2D FGRE sequence

TR=11.2ms, TE=5.0ms, flip angle=10°,
bandwidth= $\pm 8\text{KHz}$, FOV=180x72mm, 200mm slice
thickness, 128x52 matrix, acquisition time=0.58s/image

Methods

High Resolution ^{13}C Image Reconstruction

- Sixteen, fully sampled lines at centre of k-space were used to determine coil sensitivities
- Generalized encoding matrix reconstruction with:
Sub-sampling to simulate outer reduction factors (ORF) of 2, 3, and 4 and net reduction factors (R) of 1.5, 1.9 and 2.1 respectively

Methods

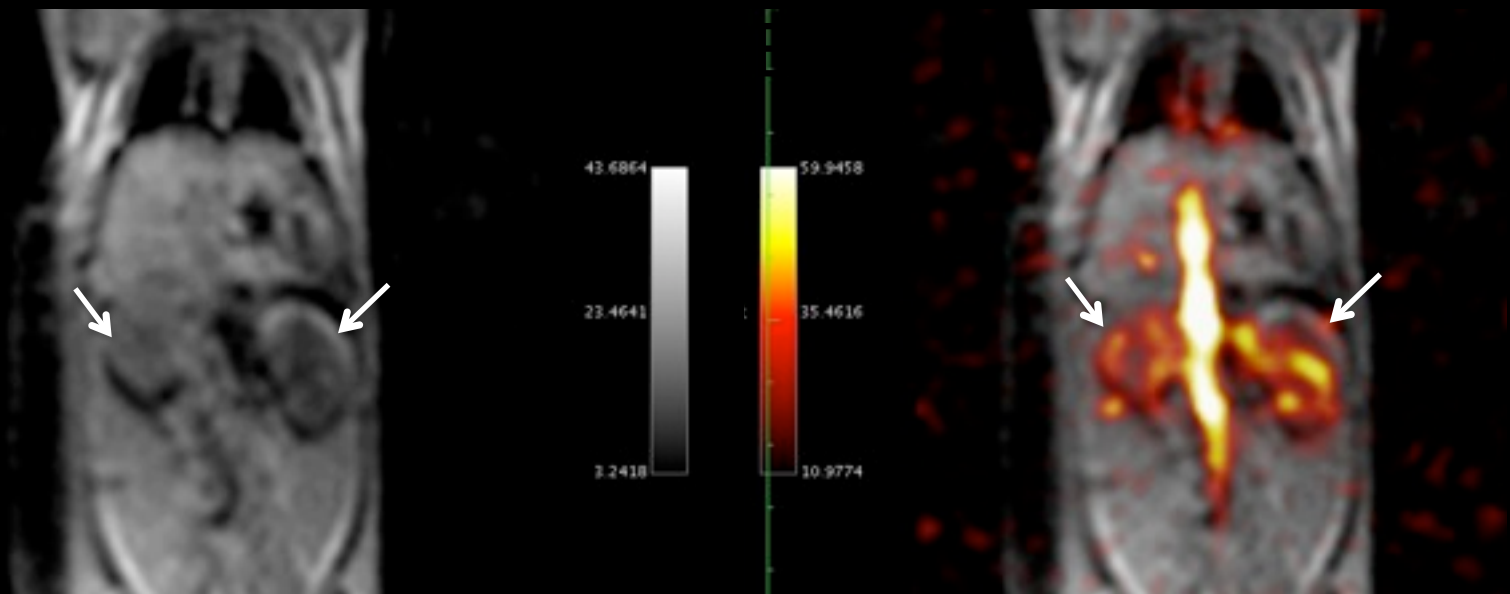
3-dimensional ^{13}C Imaging

- Fully gradient encoded images collected at 14 seconds post-injection of $[1-^{13}\text{C}]$ -pyruvate to capture $[1-^{13}\text{C}]$ -pyruvic acid in the vasculature and major organs (heart and kidneys)
- 3D FGRE sequence
TR=12.3ms, TE=5.0ms, flip angle=4°, bandwidth=±4KHz, FOV=200x100mm, 5mm slice thickness, 64x64 matrix

Results

Fig 1:

A) ^1H SPGR image. B) Overlay of ^{13}C image on the proton image shown in A. Arrows point to kidneys.



Results

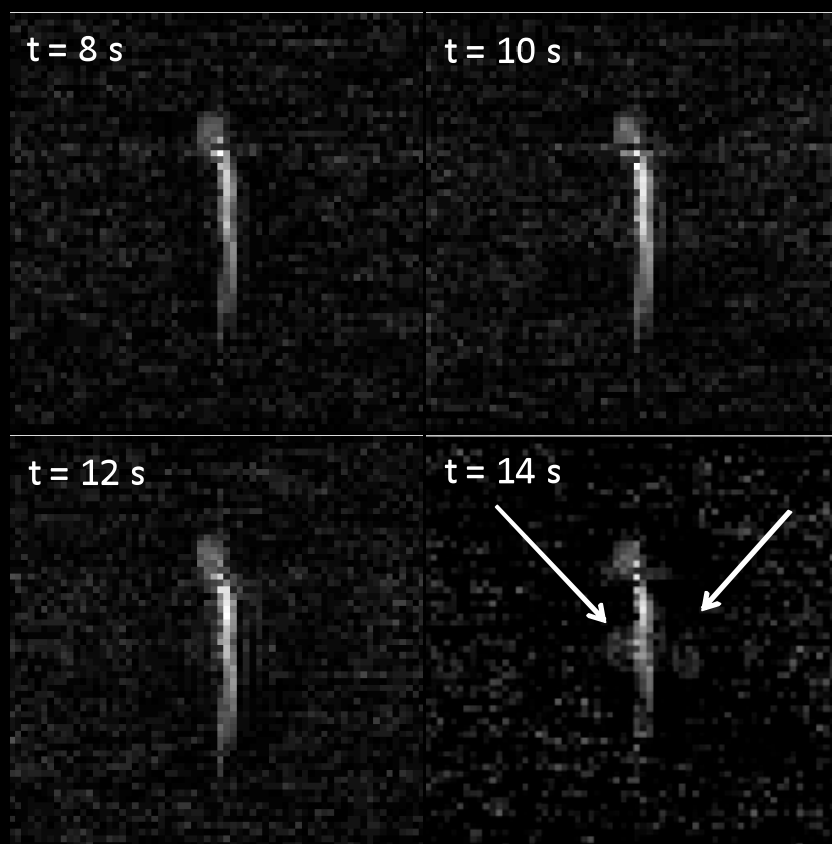


Fig 2:

- PMRI accelerated ($R=1.7$) projection images starting 8s following injection of 80mM $[1-^{13}\text{C}]$ -pyruvate.
- In-plane resolution=2.8mm, effective acquisition time=0.41ms, acquisition interval=2.0s.
- Flow of $[1-^{13}\text{C}]$ -pyruvate into the kidneys is apparent at 14s post-injection (arrows).

Results

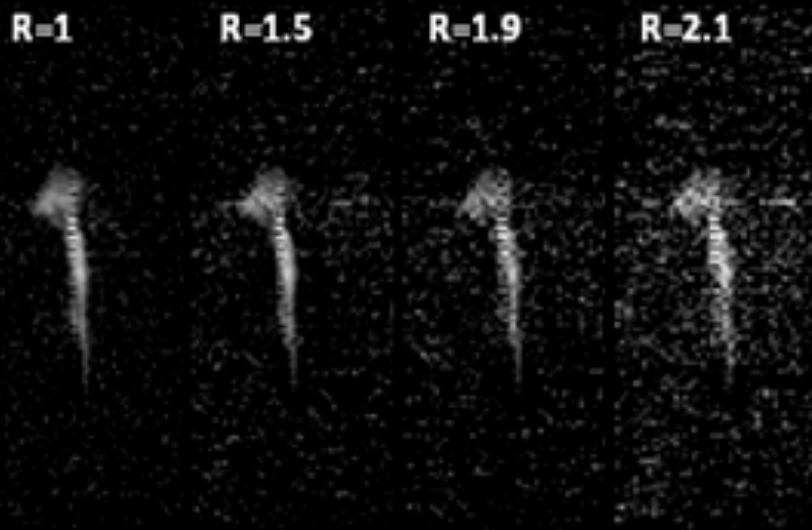


Fig 3:

- PMRI accelerated projection images acquired 8s following injection of 80mM $[1-^{13}\text{C}]$ -pyruvate.

- k-space data were decimated to obtain net reduction factors as shown (R=1=no acceleration).
- In plane resolution = 1.4mm, effective acquisition times = 0.58ms, 0.39ms, 0.31ms, and 0.28ms

Results

Fig 4:

Volume rendering of 3-dimensional ^{13}C images

Results

Fig 4:

Volume rendering of 3-dimensional ^{13}C images

WL: 2881 WW: 4907

S



RA

LP

I



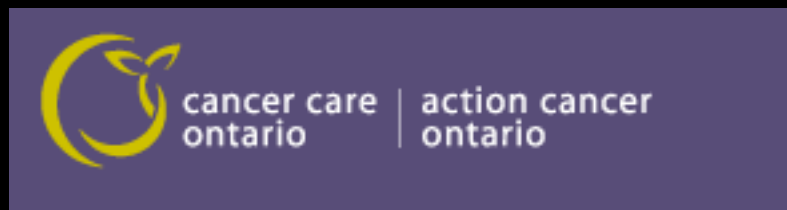
Discussion

- 1D reduction factors up to 2 were achieved before increasing g-factor caused excessive SNR loss.
- Artificial Decimation:
 - Led to significant under estimation of achievable SNR.
 - In truly accelerated images, fewer phase encodes are required allowing larger flip angles, and increased SNR.
- Variable flip angle schemes may significantly increase SNR.

Conclusion

- Self-calibrated parallel MRI was used to obtain dynamic and high resolution images of a small animal injected with hyperpolarized $[1-^{13}\text{C}]$ -pyruvate.
- ^{13}C images were obtained using a novel 8-channel receive array.

Acknowledgements



Canada Research
Chairs

Chaires de recherche
du Canada

