

# Susceptibility Weighted Imaging of Arteries and Veins with Ferumoxytol

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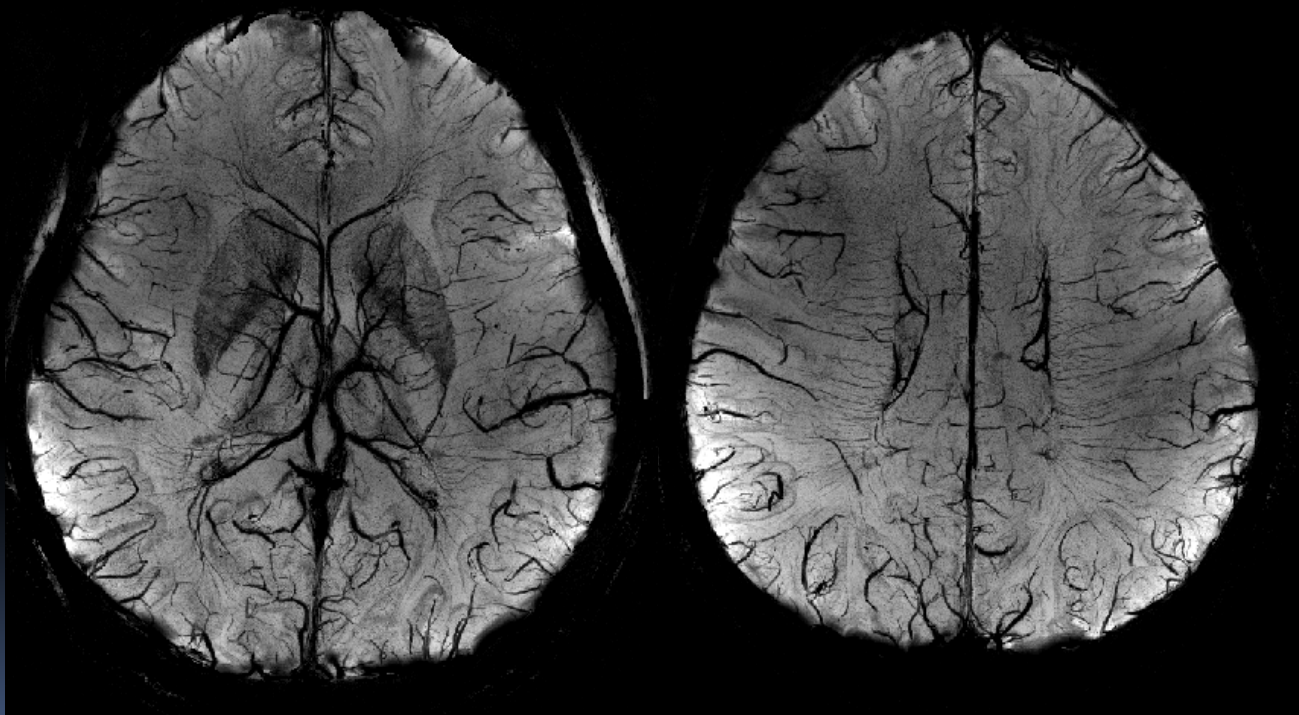
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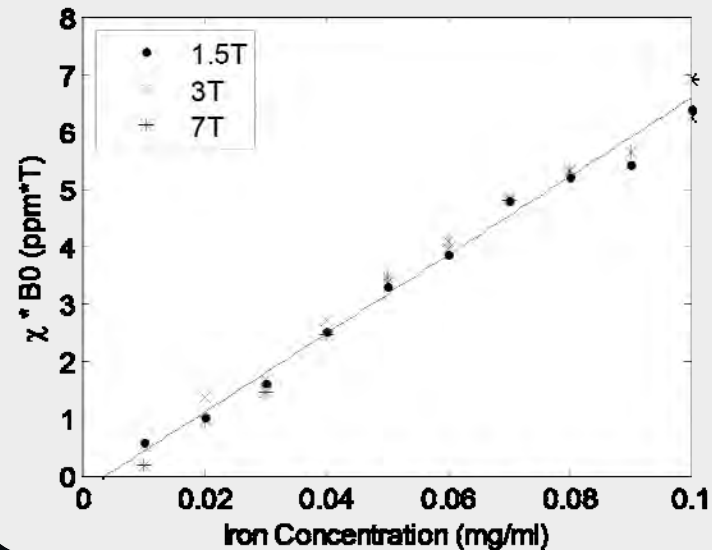
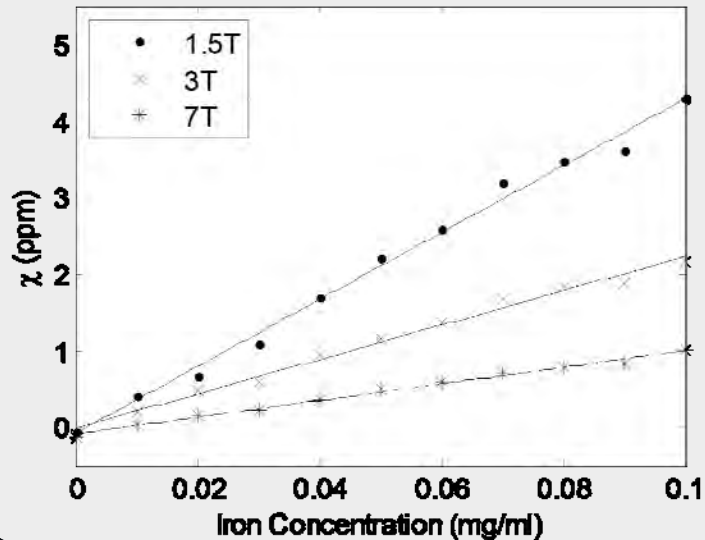
The 28<sup>th</sup> Annual International Conference, Society for Magnetic Resonance Angiography  
September 22<sup>nd</sup>, 2016, Chicago, USA

## Visualizing Veins using Susceptibility Weighted Imaging (SWI)



Minimum intensity projections of SWI. Effective slice thickness = 16mm.  
 $B_0=7T$ ,  $TE=16ms$ , voxel size =  $0.22 \times 0.22 \times 2 \text{ mm}^3$ .

# Phantom Studies



- $\chi_{1.5T} = 43.75Iron - 0.09$  [1]
- $\chi_{3T} = 22.47Iron - 0.02$  [2]
- $\chi_{7T} = 10.87Iron - 0.09$  [3]
- $\chi_{B_0} = 68.45Iron - 0.27$  [4]



Assuming blood volume 4700ml and weight 60kg, with 4mg/Kg Ferumoxytol at 7T (1.0mg/Kg, at 1.5T; 1.6mg/Kg, at 3T), the expected susceptibility of the arteries is ~450ppb.

# In vivo Data Acquisition and Processing

- Pre- and post- Ferumoxytol (1mg/Kg, 2mg/Kg, 3mg/Kg, 4mg/Kg) data were collected on 3 healthy volunteers at 7T, using a high resolution asymmetric gradient echo sequence. The highest in-plane resolution was 110um.
- Phase images were reconstructed with Echo Center Correction<sup>1</sup>, and were unwrapped using Laplacian unwrapping<sup>2</sup>.
- For generating SWI, 64x64 homodyne high-pass filter was used to remove the artifacts in phase images<sup>3</sup>.
- Susceptibility maps were generated using iterative SWIM algorithm<sup>4</sup>. The background field was removed using SHARP<sup>5</sup>.

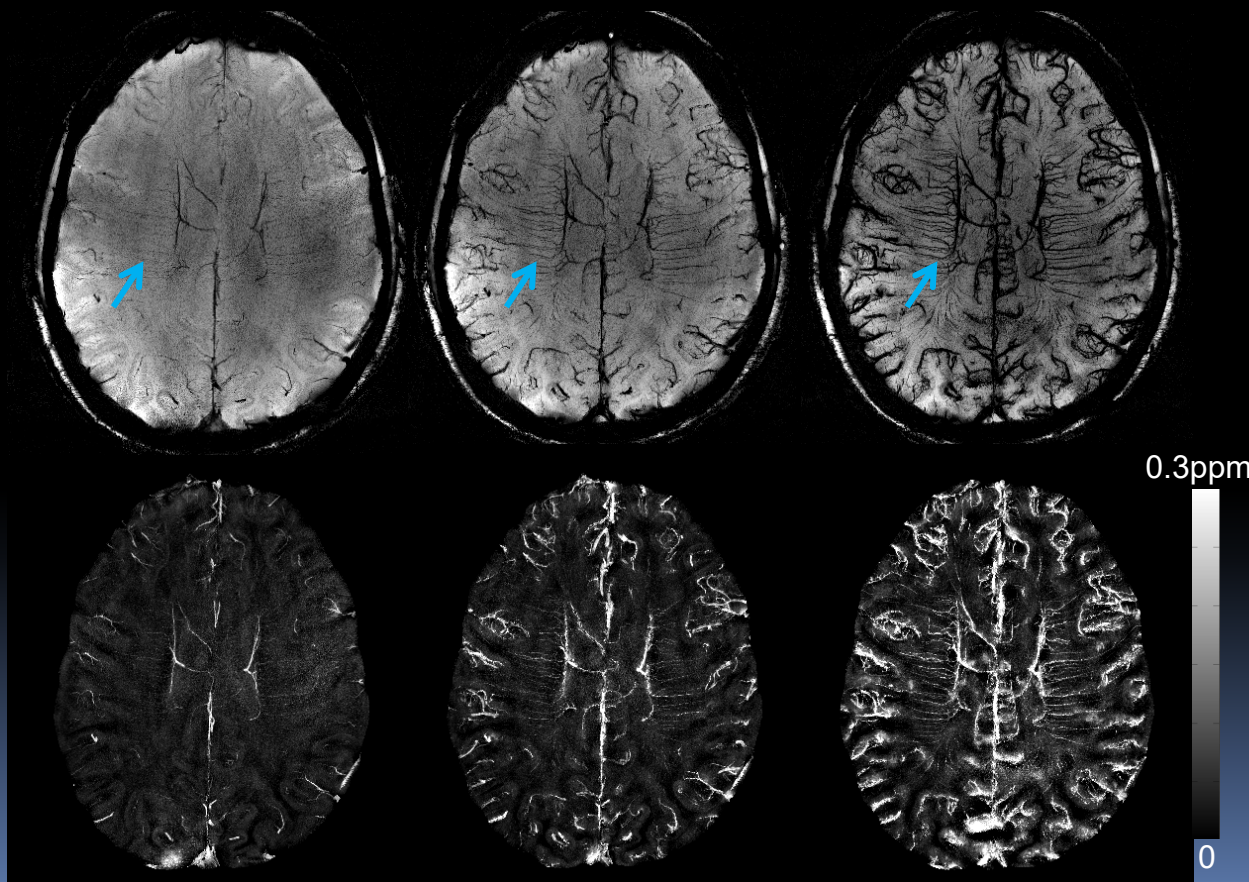
1. Liu, S. et al. 2016, NMR Biomed. doi: [10.1002/nbm.3552](https://doi.org/10.1002/nbm.3552). 2. Haacke et al. 2015, MRI 33:1–25  
3. Haacke et al. 2004, MRM 52:612–618. 4. Tang et al. 2013, MRM, 69: 1396–1407  
5. Schweser et al. 2011, NeuroImage 54:2789–2807

# In vivo Data Results

Pre

Post 1mg/Kg

Post 3mg/Kg



mIP of SWI, Effective  
Slice Thickness = 6.4mm

$B_0=7T$ ,  $TE=10ms$ ,  $TR=24ms$ ,  
 $FA=10^\circ$ ,  $BW/px = 244 \text{ Hz/px}$ ,  
Voxel size =  $0.13 \times 0.13 \times 0.8 \text{ mm}^3$

MIP of QSM, Effective  
Slice Thickness = 6.4mm

## Differentiate Arteries from Veins

- Strategies to differentiate arteries from veins in the post-contrast data:
  - Pre-contrast magnitude => arteries
  - Pre-contrast SWI and QSM => veins
  - Susceptibility difference between arteries and veins in the post-contrast QSM
  - Connectivity between the pixels

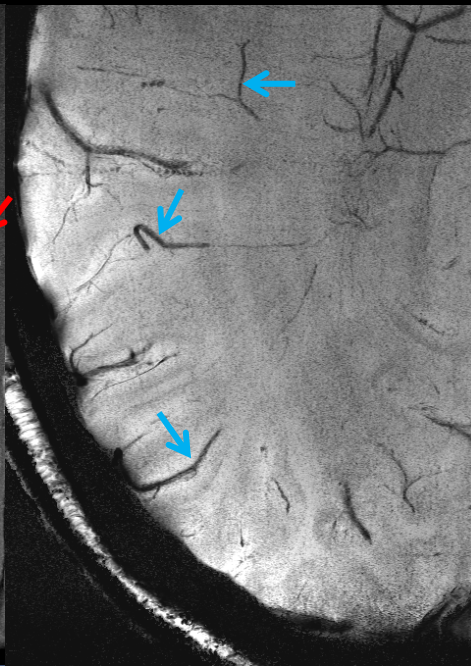


# Differentiate Arteries from Veins

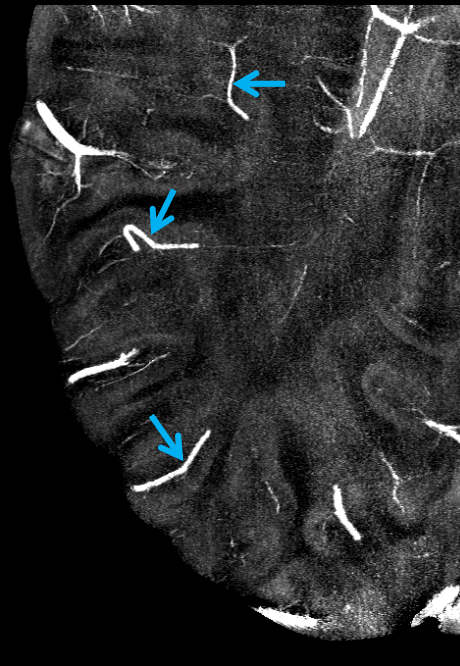
MIP of pre-contrast  
magnitude images



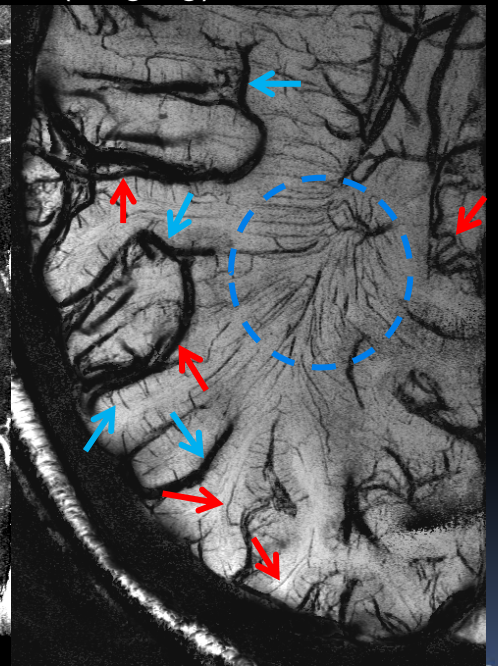
mIP of pre-contrast  
SWI data



MIP of pre-contrast  
QSM data



mIP of post-Ferumoxytol  
(3mg/Kg) SWI data

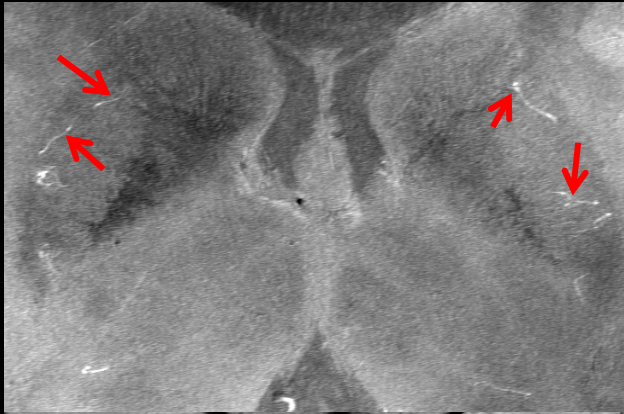


↑ : Artery

↑ : Vein

Effective Slice Thickness = 6.4mm

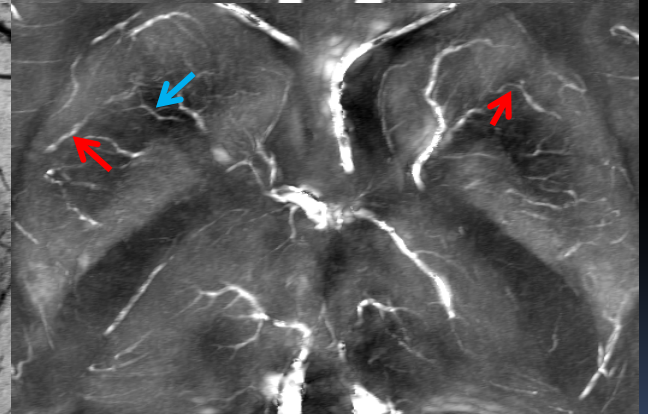
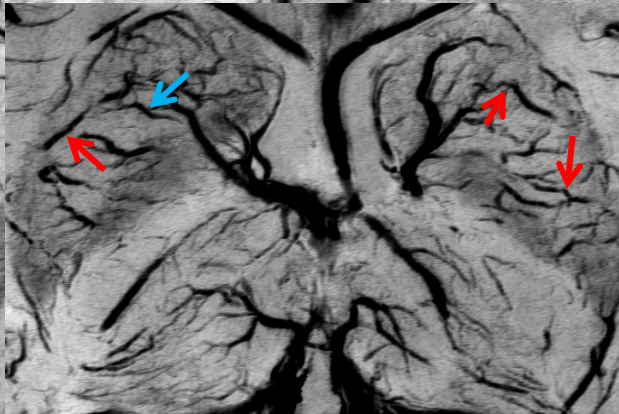
MIP of pre magnitude



mIP of pre SWI



MIP of pre SWIM



mIP of post-2mg/kg SWI

mIP of post-4mg/kg SWI

MIP of post-2mg/kg SWIM

↑ : Artery    ↑ : Vein

$B_0=7T$ ,  $TE=8ms$ , Voxel size:  $0.1094 \times 0.1094 \times 1.25mm^3$ .  
Effective slice thickness = 5mm.

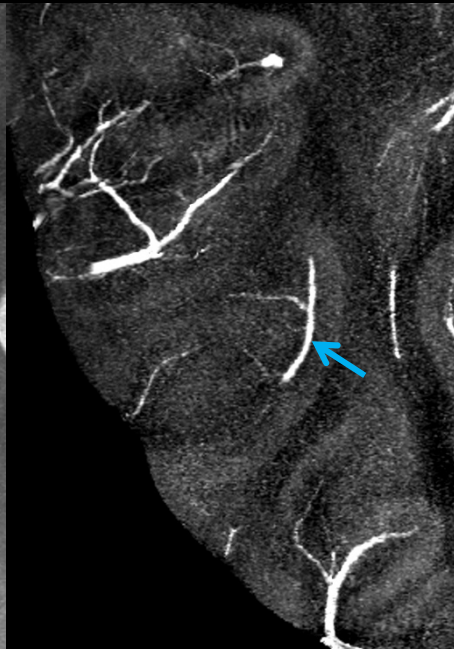


# Differentiate Arteries from Veins

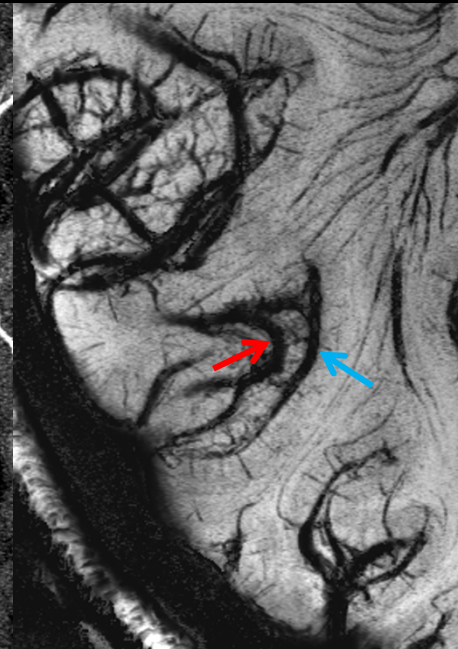
MIP of pre-contrast  
magnitude images



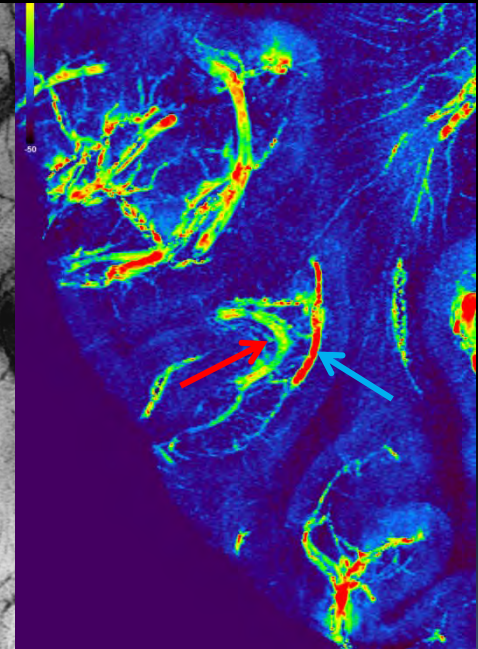
MIP of pre-contrast  
QSM data



mIP of pre-contrast  
SWI data



mIP of post-Ferumoxytol  
(3mg/Kg) SWI data



↑: Artery

↑: Vein

Effective Slice Thickness = 6.4mm

## Conclusions

- We have demonstrated the feasibility to generate ultra-high-resolution MRAV using USPIO-enhanced SWI.
- Using a combination of pre-contrast magnitude, SWI, QSM and the difference between the susceptibilities of arteries and veins, we are able to separate arteries and veins.
- This could represent a powerful new tool in detecting microvascular abnormalities not visible on conventional MRA.

