

Susceptibility Weighted Imaging of Arteries and Veins with Ferumoxytol

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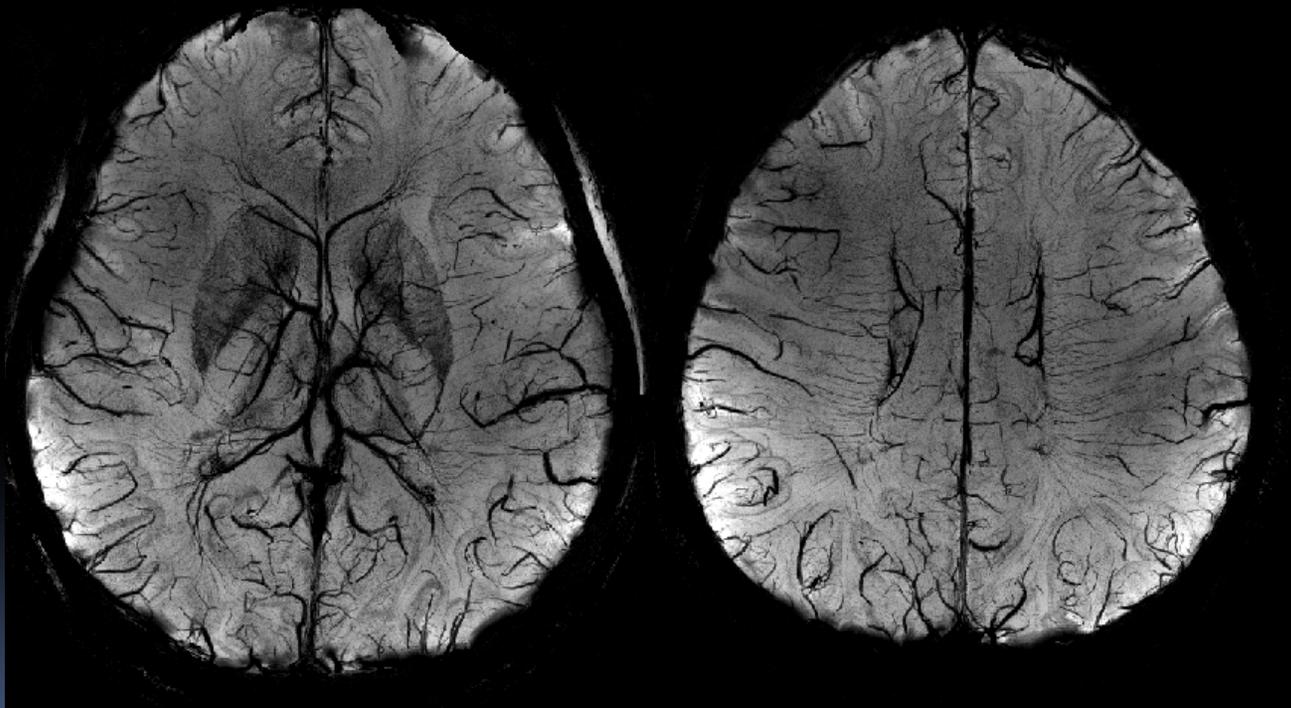
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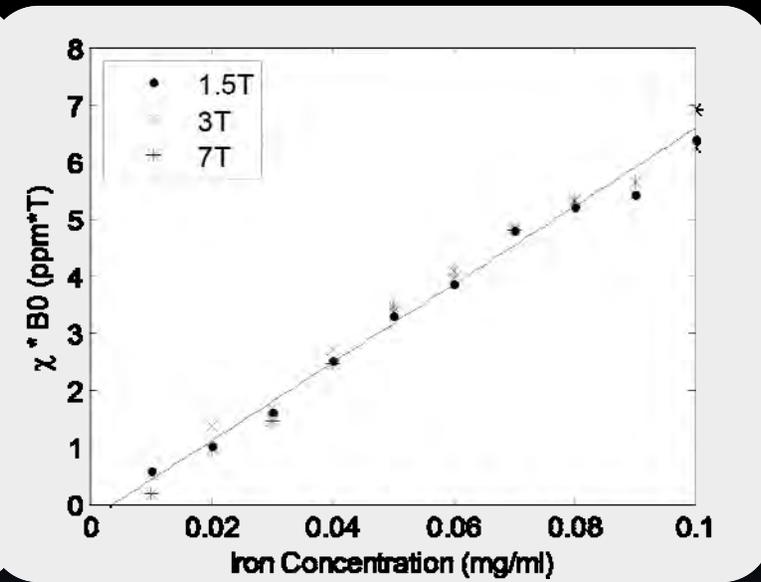
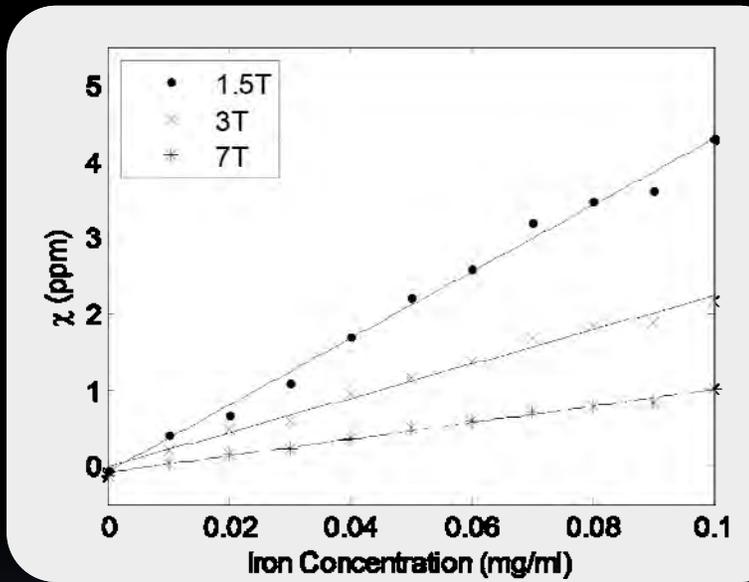
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Visualizing Veins using Susceptibility Weighted Imaging (SWI)



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 Correction¹, and were unwrapped using Laplacian unwrapping².
- For generating SWI, 64x64 homodyne high-pass filter was used to remove the artifacts in phase images³.
- Susceptibility maps were generated using iterative SWIM algorithm⁴. The background field was removed using SHARP⁵.

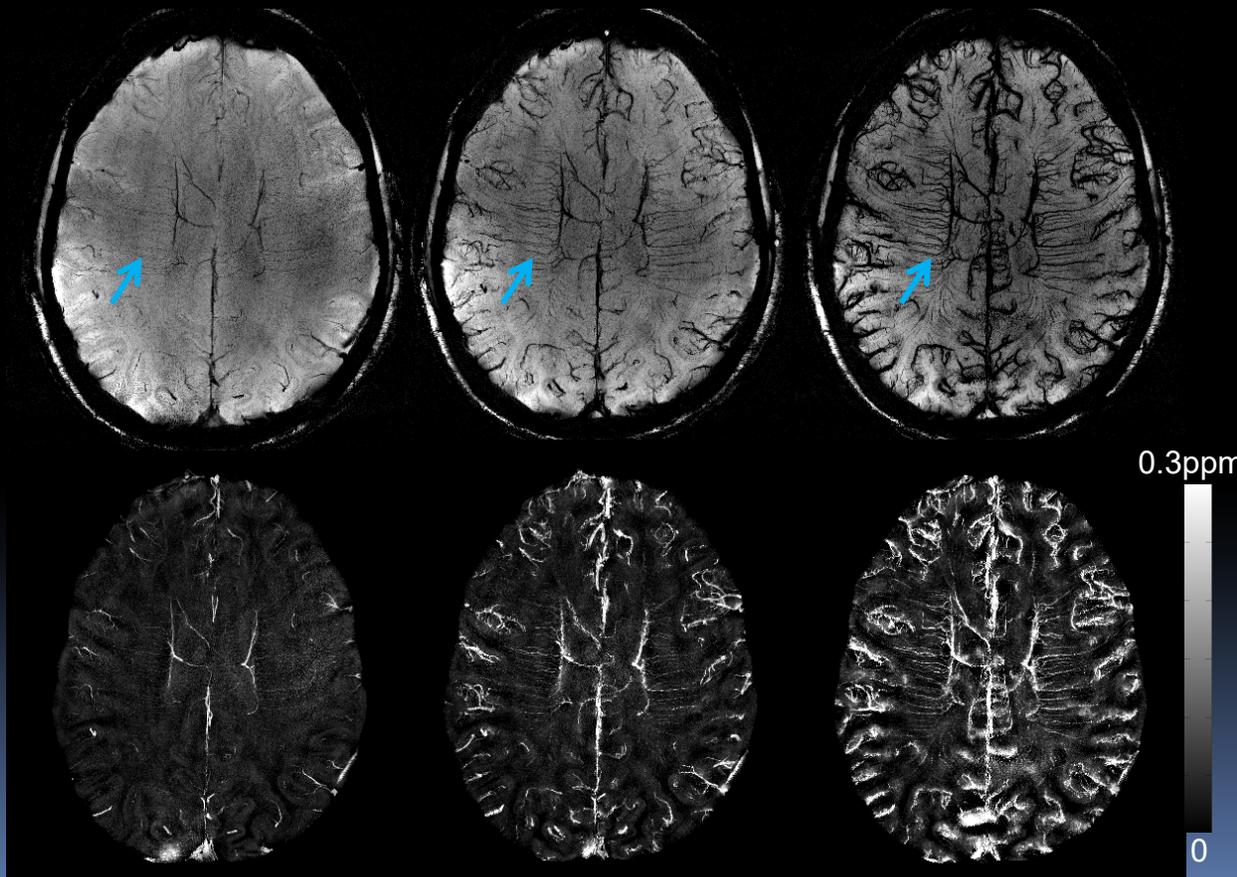
1. Liu, S. et al. 2016, NMR Biomed. doi: [10.1002/nbm.3552](https://doi.org/10.1002/nbm.3552). 2. Haacke et al. 2015, MRI 33:1–25
3. Haacke et al. 2004, MRM 52:612–618. 4. Tang et al. 2013, MRM, 69: 1396–1407
5. Schweser et al. 2011, NeuroImage 54:2789–2807

In vivo Data Results

Pre

Post 1mg/Kg

Post 3mg/Kg



mIP of SWI, Effective
Slice Thickness = 6.4mm

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

MIP of QSM, Effective
Slice Thickness = 6.4mm

Differentiate Arteries from Veins

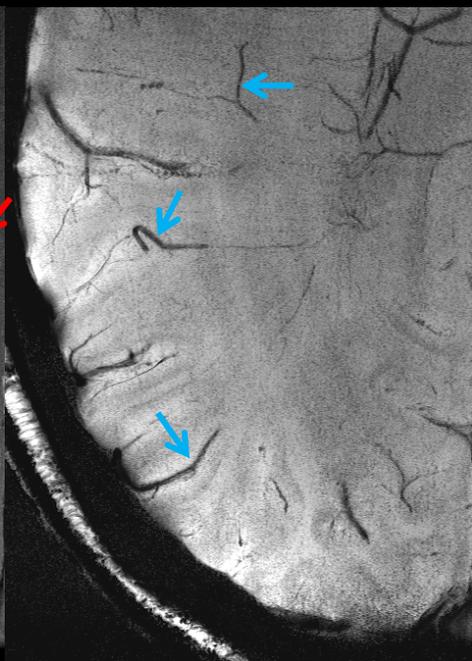
- Strategies to differentiate arteries from veins in the post-contrast data:
 - Pre-contrast magnitude => arteries
 - Pre-contrast SWI and QSM => veins
 - Susceptibility difference between arteries and veins in the post-contrast QSM
 - Connectivity between the pixels

Differentiate Arteries from Veins

MIP of pre-contrast magnitude images



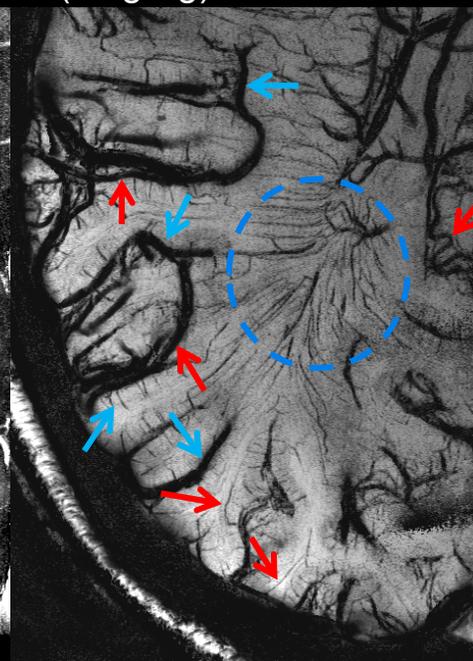
mIP of pre-contrast SWI data



MIP of pre-contrast QSM data



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



↑ : Artery

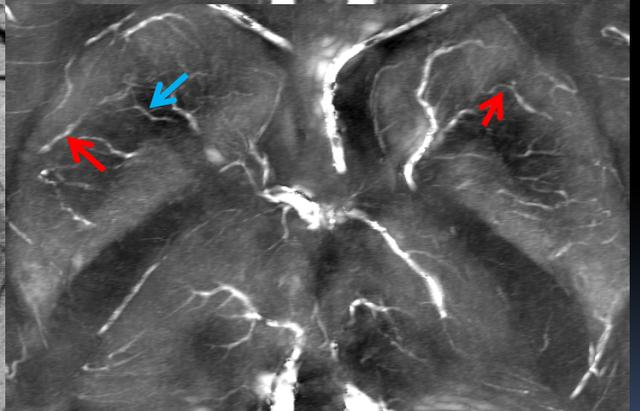
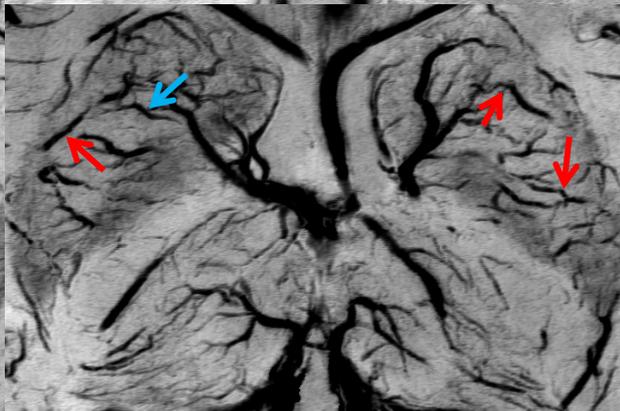
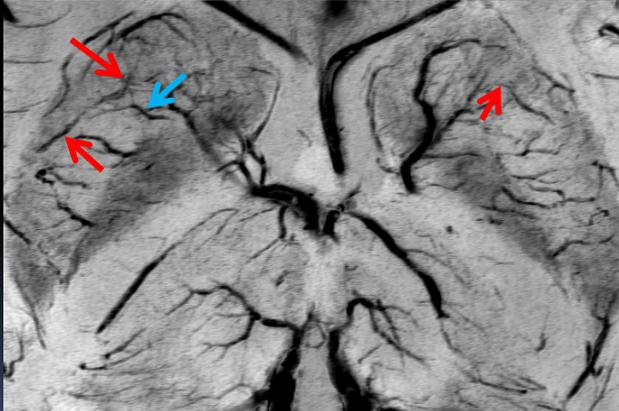
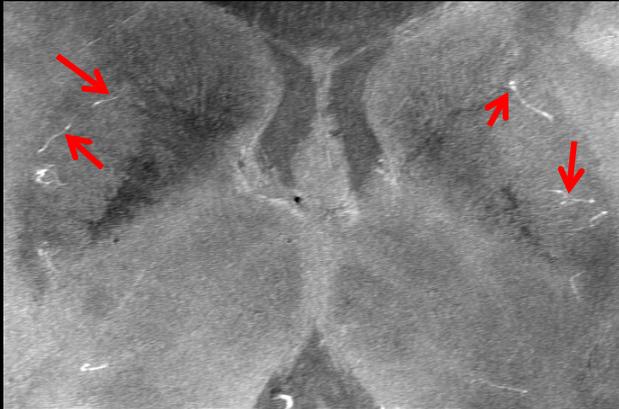
↑ : Vein

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

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

↑ : Artery ↑ : Vein

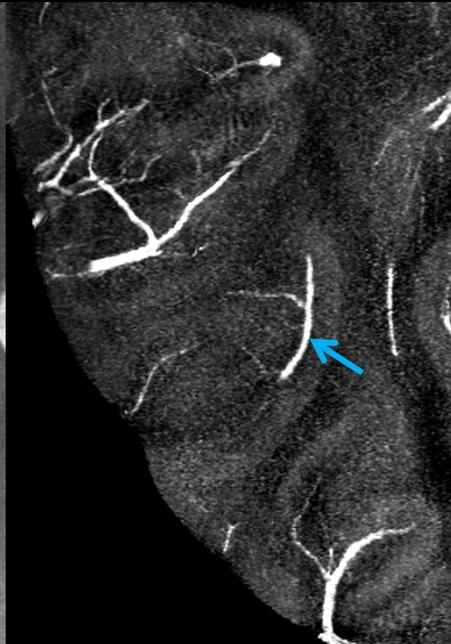
Differentiate Arteries from Veins

MIP of pre-contrast magnitude images



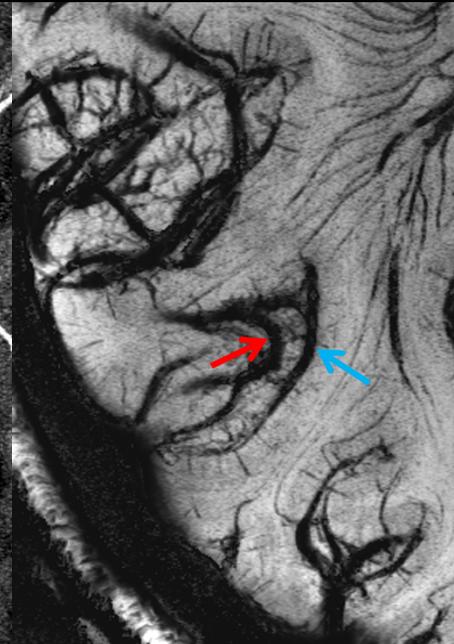
↑: Artery

MIP of pre-contrast QSM data



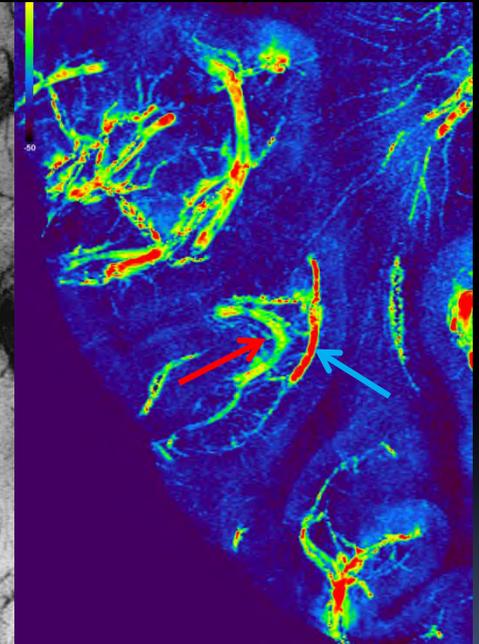
↑: Vein

mIP of pre-contrast SWI data



Effective Slice Thickness = 6.4mm

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



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.

