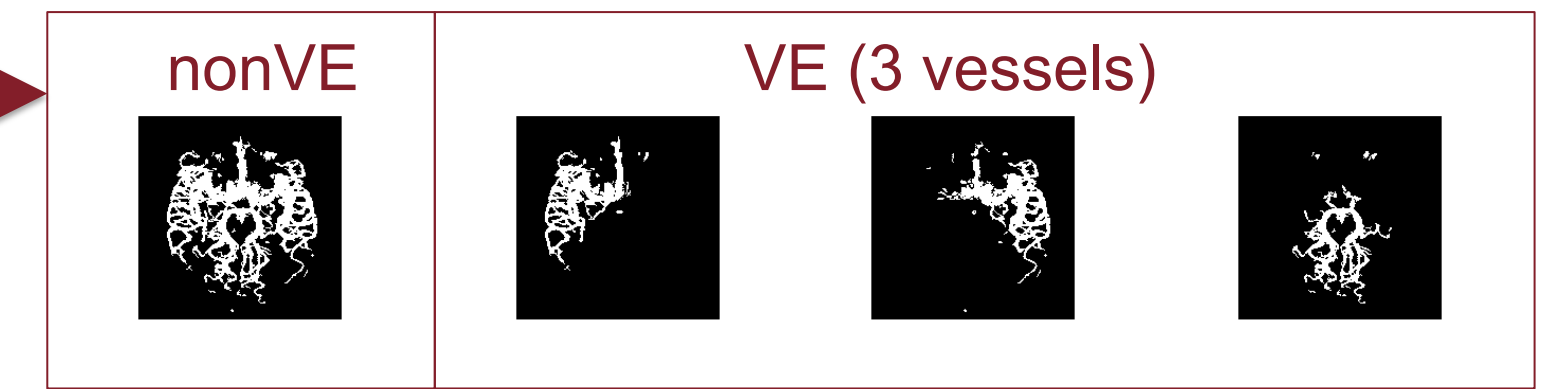
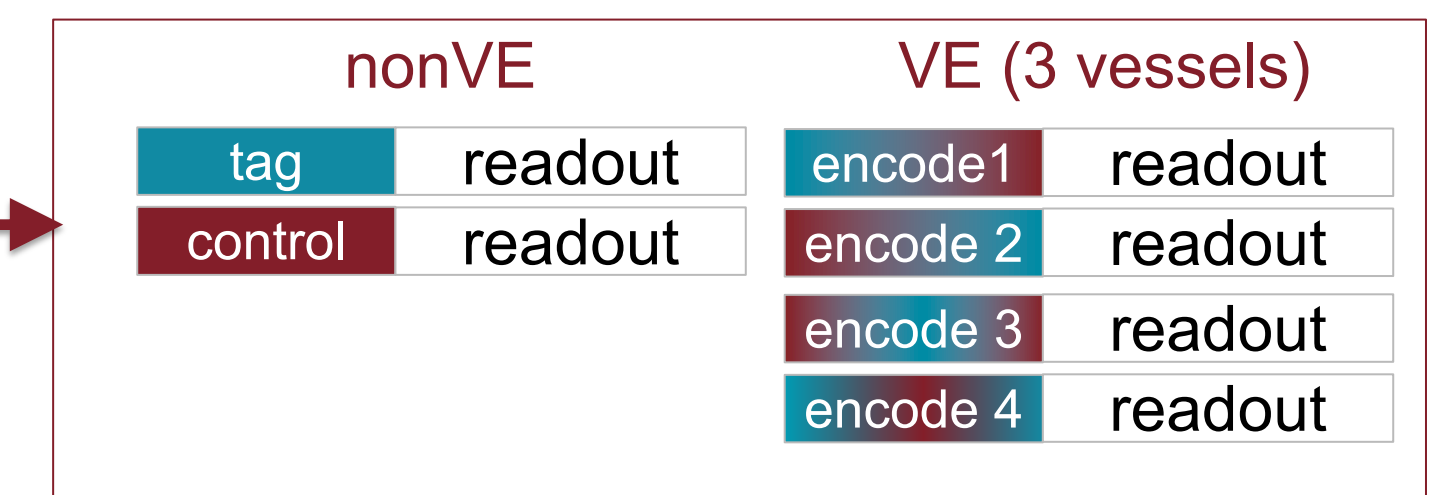


4D Vessel-Encoded pCASL Angiography in a Five-Minute Scan

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Introduction

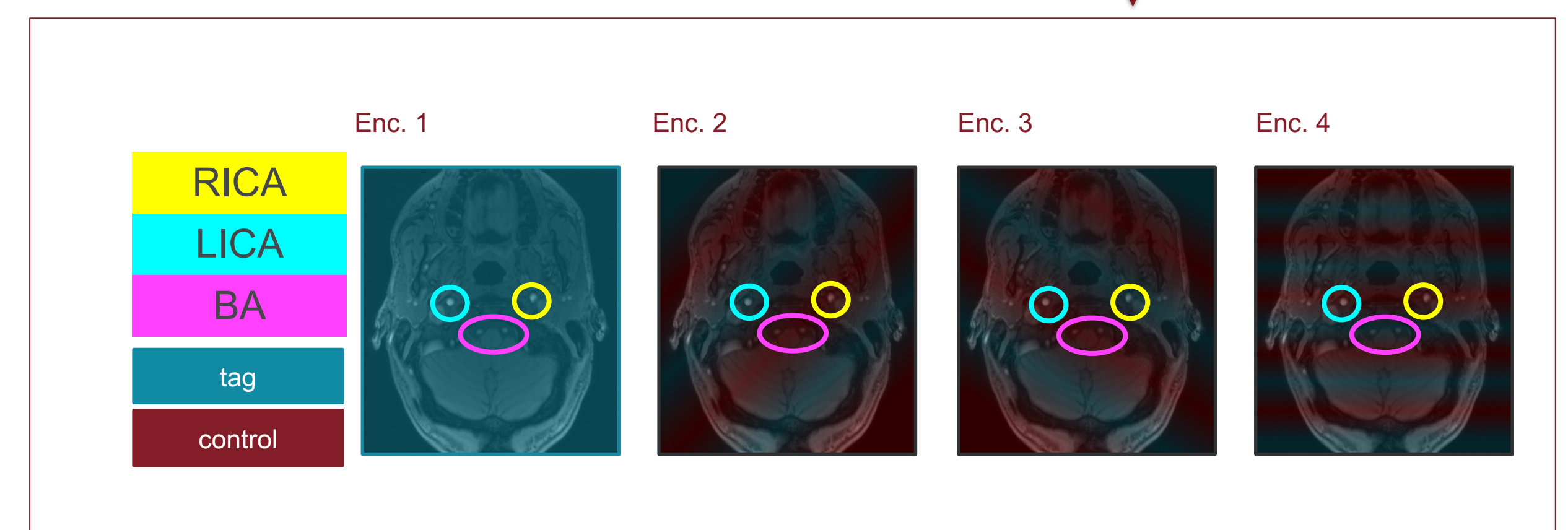
- Vessel encoded VE-pCASL¹ angiography² provides potentially crucial information for a range of cerebrovascular diseases in a non-invasive way
- VE-pCASL is as SNR efficient as conventional pCASL, which is an advantage over single-artery selective methods
- Long scan times because of:
 - additional encoding steps (N+1 encodings for N vessels) compared to conventional (nonVE) angiography
 - many repeats needed to fill k-space at sufficient spatial (~1mm isotropic) and temporal (~200ms) resolution in 3D
- Can be accelerated by:
 - acquiring with trajectory that produces more benign undersampling artefacts (e.g. radial)³
 - reconstructing in a compressed sensing framework⁴, exploiting the images' spatial sparsity. VE is especially well suited for this due to higher relative sparsity than nonVE angiography



Methods

Acquisition:

- Proof-of-principle study on one healthy volunteer, acquired on 3T Siemens Verio
- Three vessels (right and left internal carotid arteries (RICA, LICA) + basilar artery (BA)) were labelled with a Hadamard encoding scheme
- Scan parameters:
 - Trajectory: 3D golden angle radial spokes⁴
 - TE/TR = 5.9/11.6ms - Flip angle = 7°
 - Labelling duration: 1.0s
 - Readout time: 1.3s (108 readout spokes acquired per ASL preparation)
 - Total scan time: 5:16min (label + readout 33 times for each encoding)
 - Acceleration factor, R = 97



Reconstruction:

- Compressed sensing with spatial sparsity enforced + coil sensitivity estimation
 - Combining vessel decoding with image reconstruction
- Cost function:

$$cost = \frac{1}{2} \|E\mathbf{x} - \mathbf{d}\|_2^2 + \lambda \|\mathbf{x}\|_1$$

$E = \text{acquisition operator}$, $\mathbf{x} = \text{image}$, $\mathbf{d} = \text{data}$, $\lambda = \text{regularisation weighting}$
- Reconstruction algorithm: FISTA⁵
- Reconstruction parameters:
 - Spatial resolution: 1.1mm isotropic
 - Temporal resolution: 210ms (6 frames)

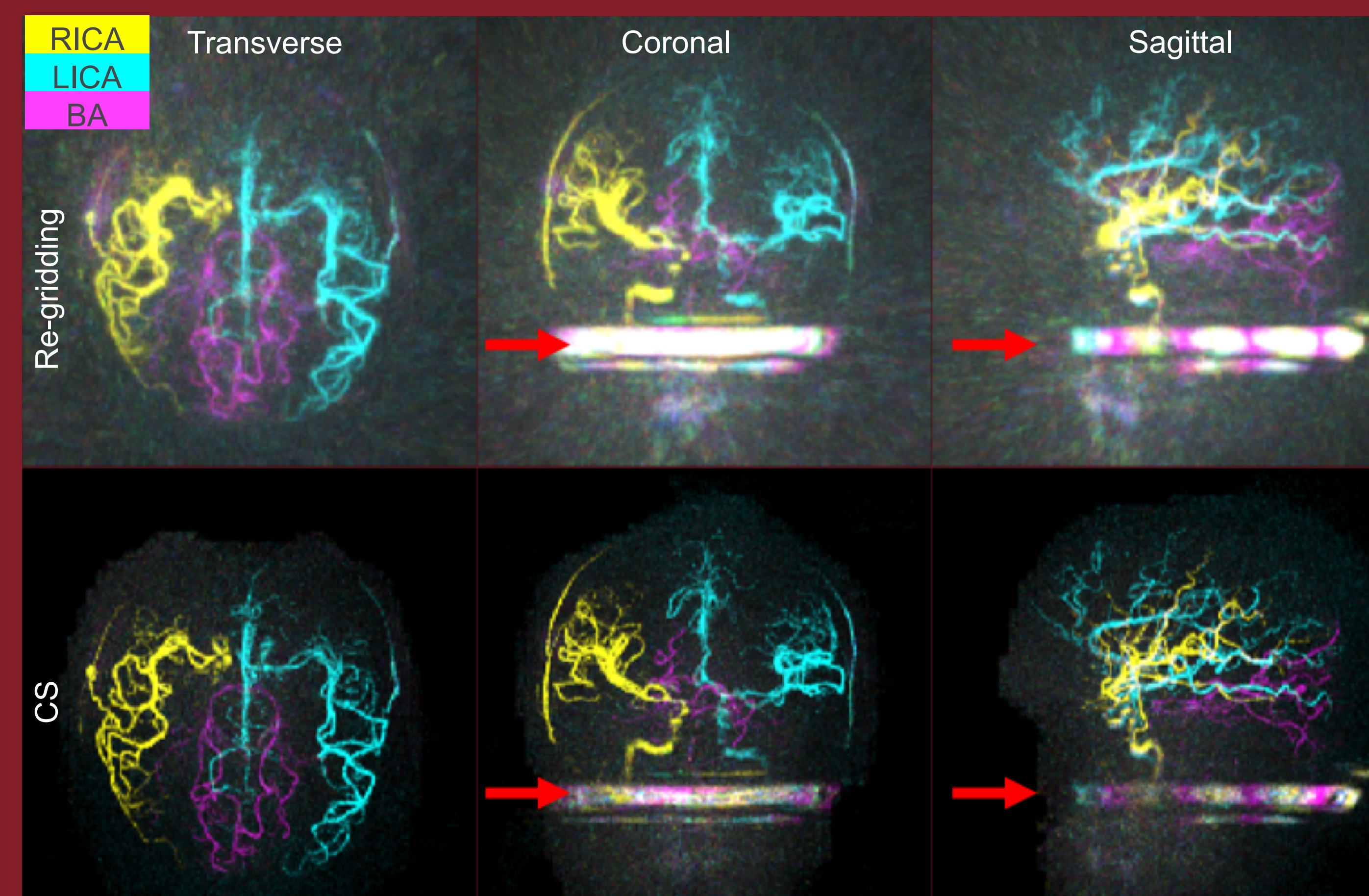
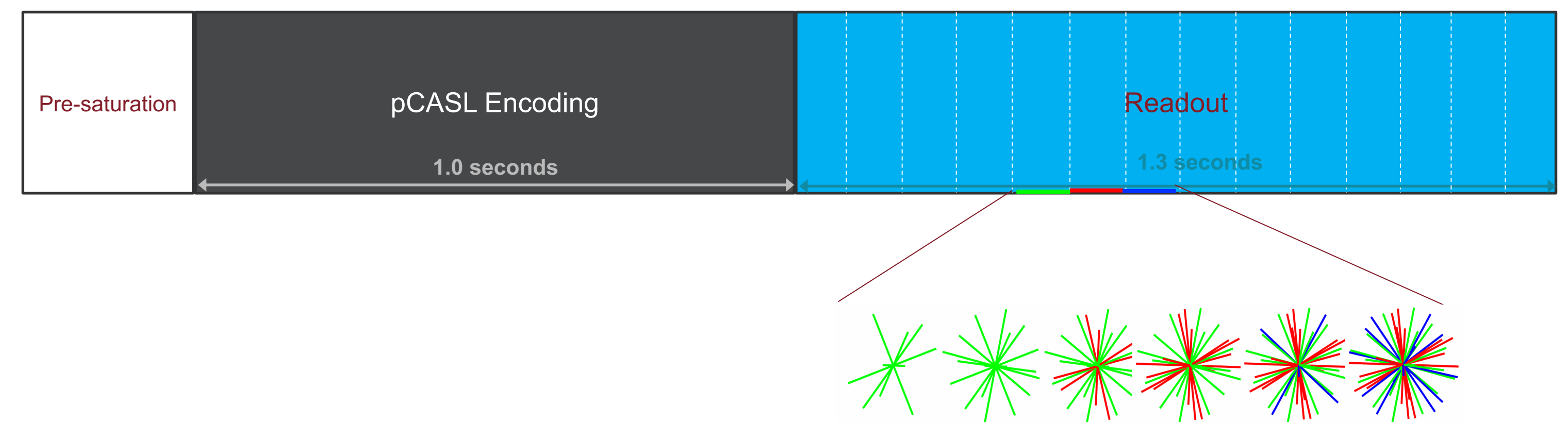


Figure 1: Above: Simple re-gridding reconstruction (temporal average MIP). Below: CS Reconstruction. The red arrows indicate the labelling plane artifact.

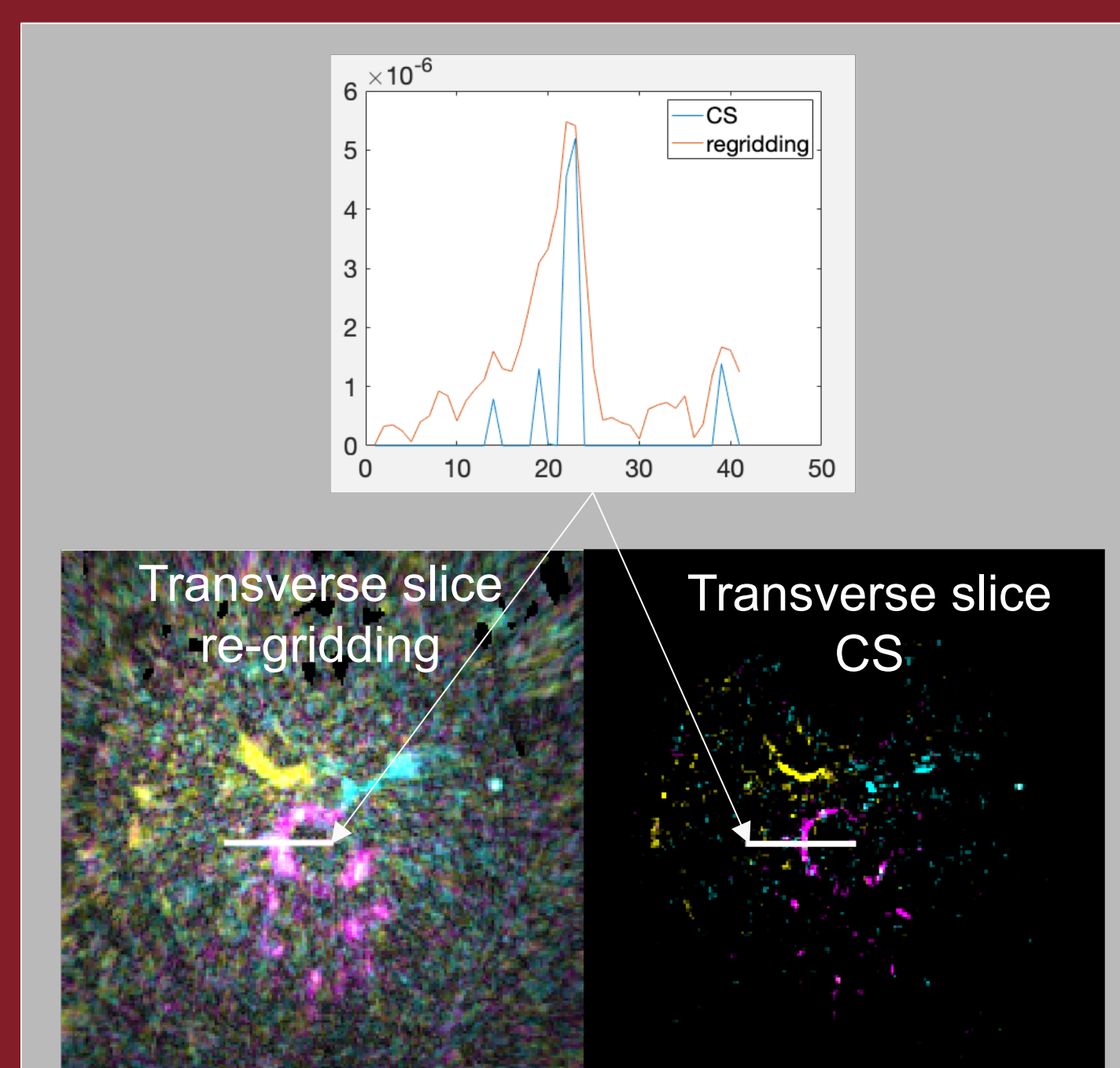


Figure 2: Example of increased SNR and lowered blurring for CS compared with re-gridding

Results

- Less blurring and a clear increase in SNR for the CS reconstructed images compared with reconstruction using simple re-gridding and coil sensitivities only (Figure 1)
 - FWHM of an example vessel (Figure 2) was 1.93 for CS and 5.77 for re-gridding
 - 19% increase in the apparent SNR (although rigorous assessment of SNR with this type of CS reconstruction requires further work)
- Small and faint vessels visible even at later frames (Figure 3)

Discussion and Conclusions

- Promising as proof-of-principle but not yet optimized
- Will need to verify these results in a more quantitative way across a larger number of subjects
- Currently, offline reconstruction times are slow due to the large memory burden (>30 GB), but optimisations for processing speed and memory efficiency are being investigated
- Temporal smoothness can also be leveraged as additional regularizer
- Variable flip angle acquisition⁵ or time-encoded ASL⁷ might preserve signal in later frames more
- Potential clinical use in evaluating the relative blood supply from different arteries in steno-occlusive disease and in planning embolization therapy for arteriovenous malformations

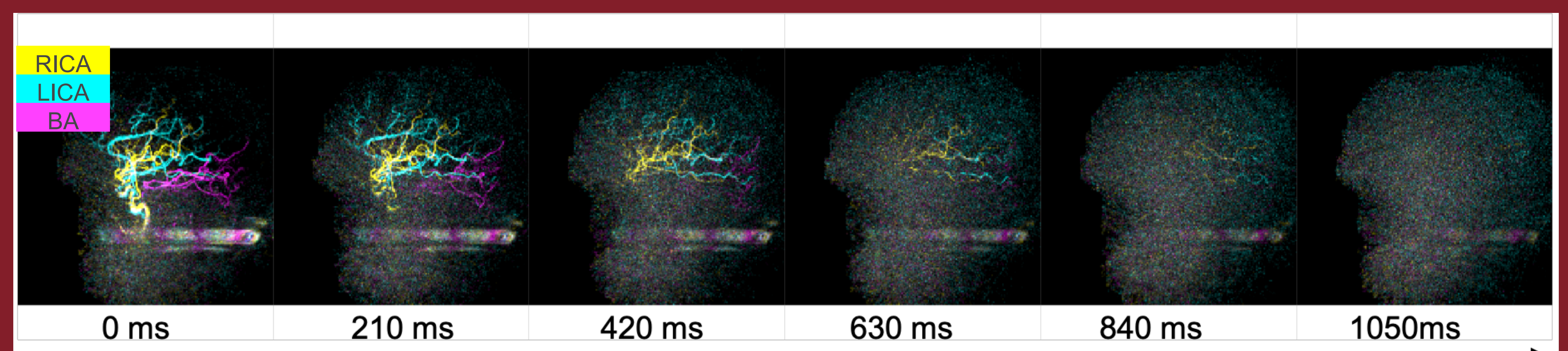


Figure 3: Temporal dynamics of the outflow of the labeled bolus. Sharp delineation of faint vessels is apparent even in later frames, despite the lower SNR. Effective post-labelling delays are shown beneath each frame.

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