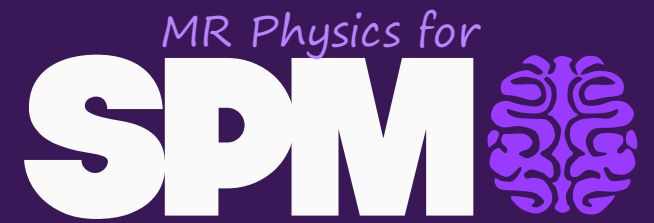


Anatomical Imaging for fMRI and VBM



Martina Callaghan

MR Physics for SPM – 26 May 2026

Functional Imaging Laboratory
Department of Imaging Neuroscience

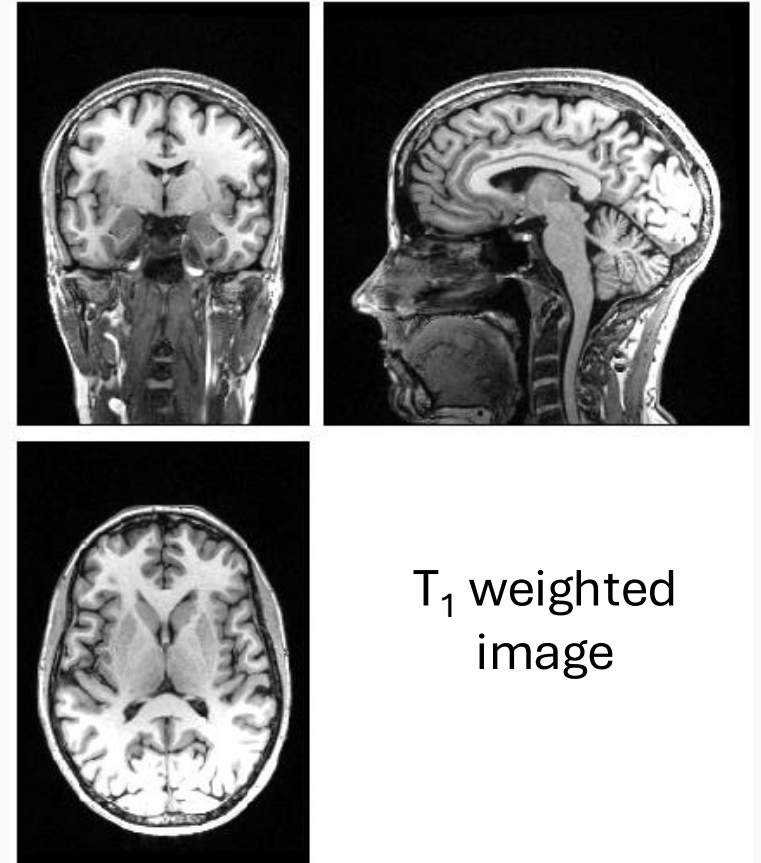


Learning Objectives

- ☐ Understand the uses of anatomical imaging in fMRI and VBM
- ☐ Understand how to generate T_1 weighting
- ☐ Identify key acquisition parameters, sequences and when to choose them
- ☐ Be mindful of the impact of image contrast on VBM conclusions

Requirements for anatomical imaging

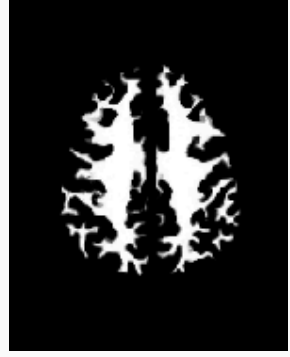
- ☐ High image contrast that accurately depicts morphology
- ☐ Whole brain coverage
- ☐ High spatial resolution
- ☐ Minimum acquisition time
- ☐ No signal losses or artefacts



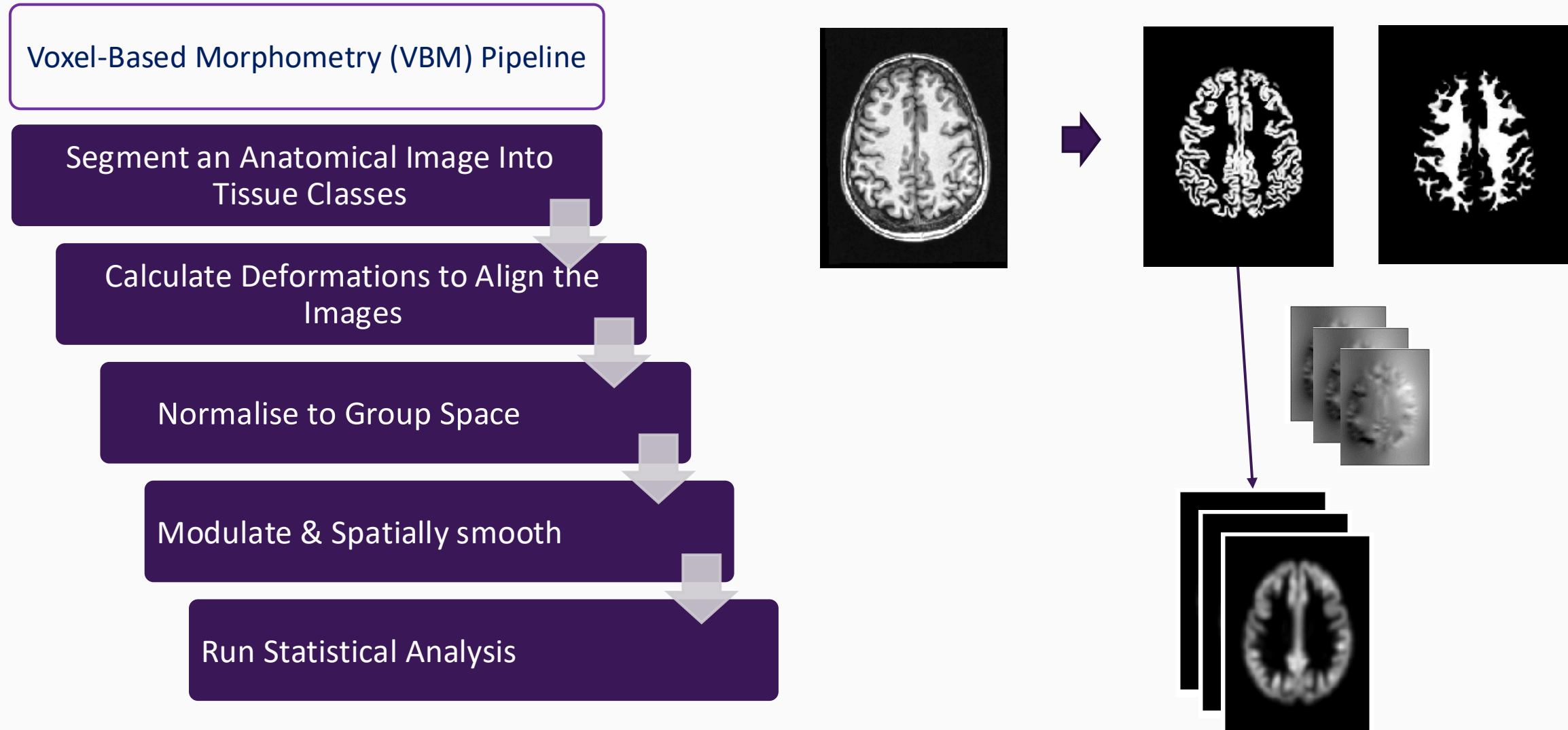
Purpose: Testing for volumetric differences

Voxel-Based Morphometry (VBM) Pipeline

Segment an Anatomical Image Into
Tissue Classes



Purpose: Testing for volumetric differences

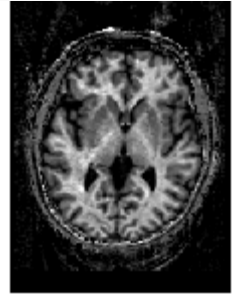


Purpose: Bringing brains into alignment

Participant 1

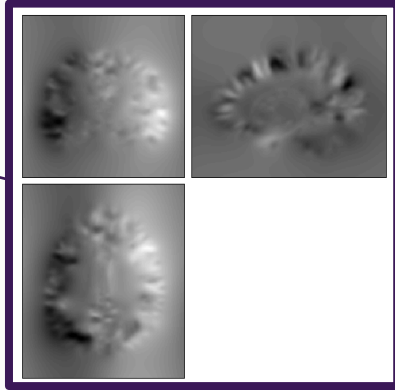
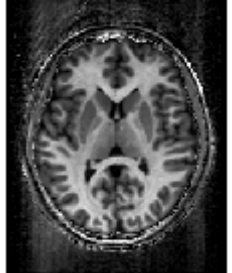


Participant 2

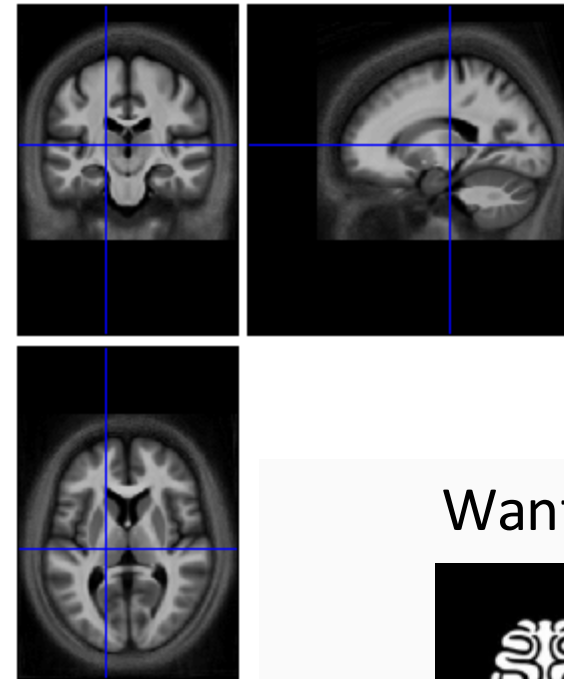


⋮

Participant N



Study-Specific Brain Template



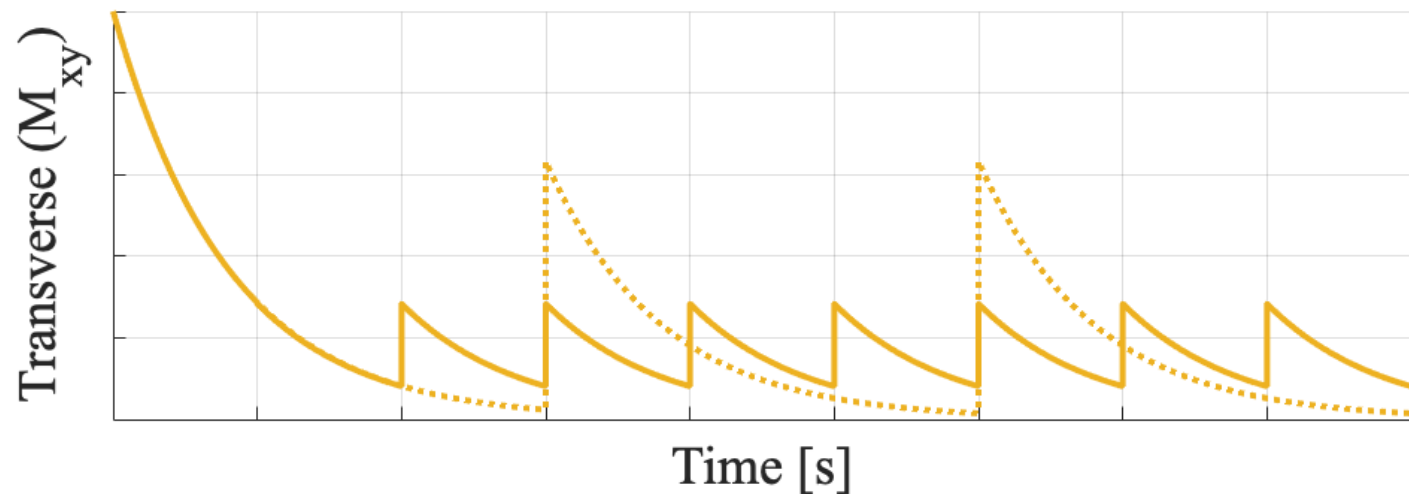
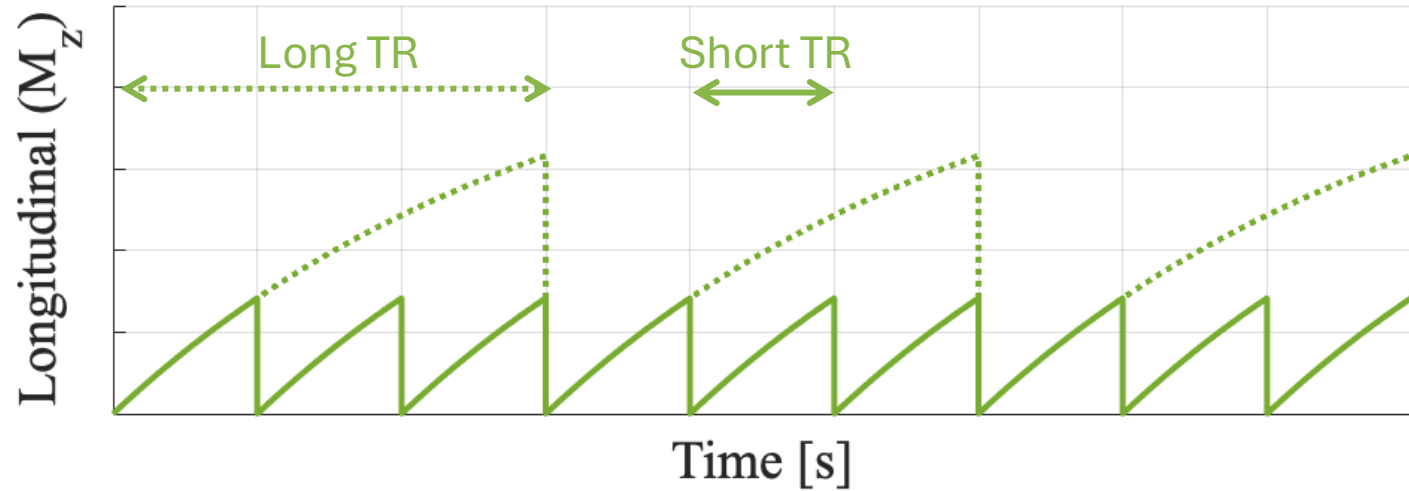
Want to learn more?



Covered in main SPM Course

Let's revisit: Relaxometry to Generate Contrast

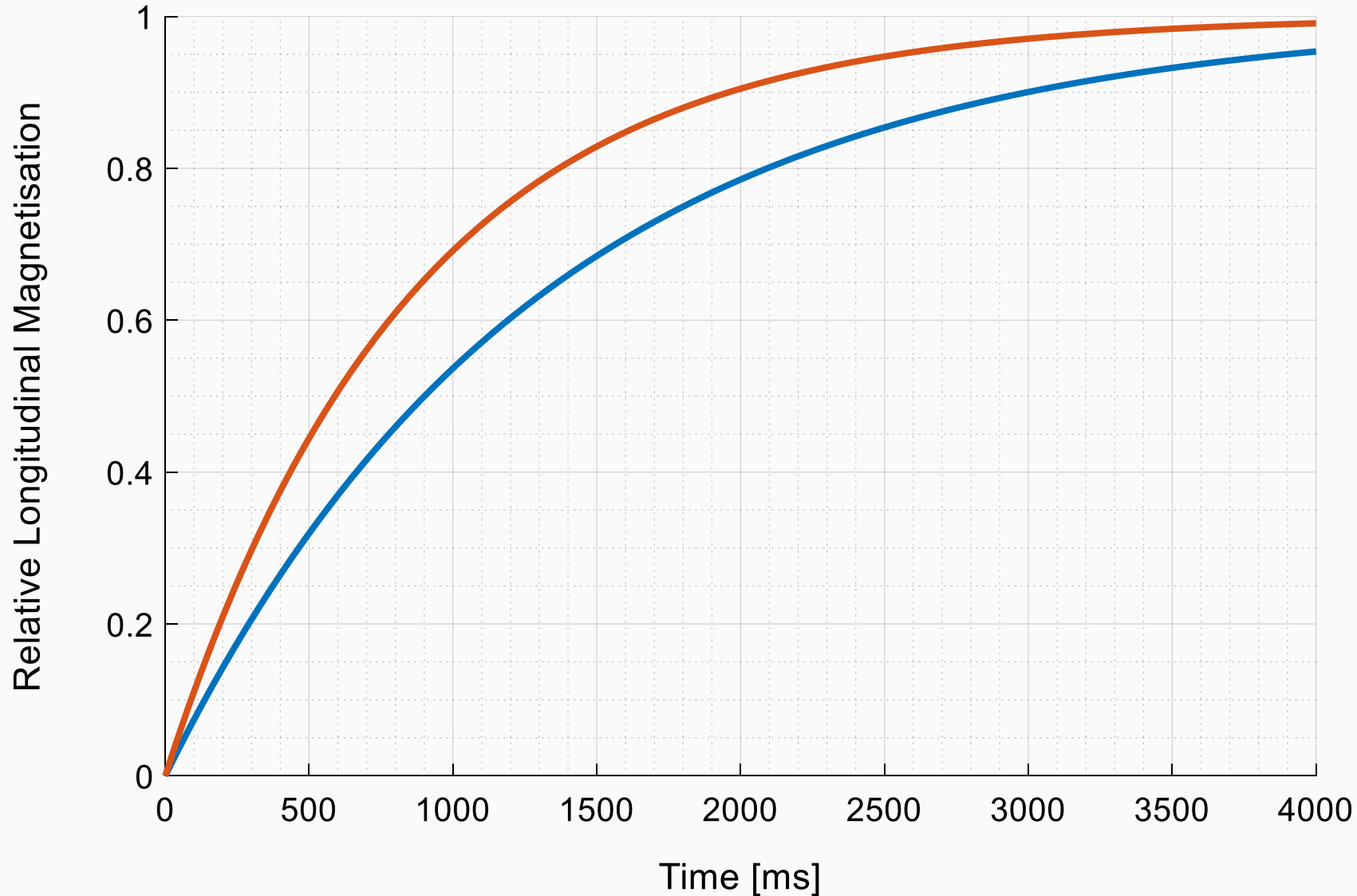
Recap: relaxometry



TR dictates how much time there is for longitudinal magnetisation to relax...

And therefore how much signal there is to sample...

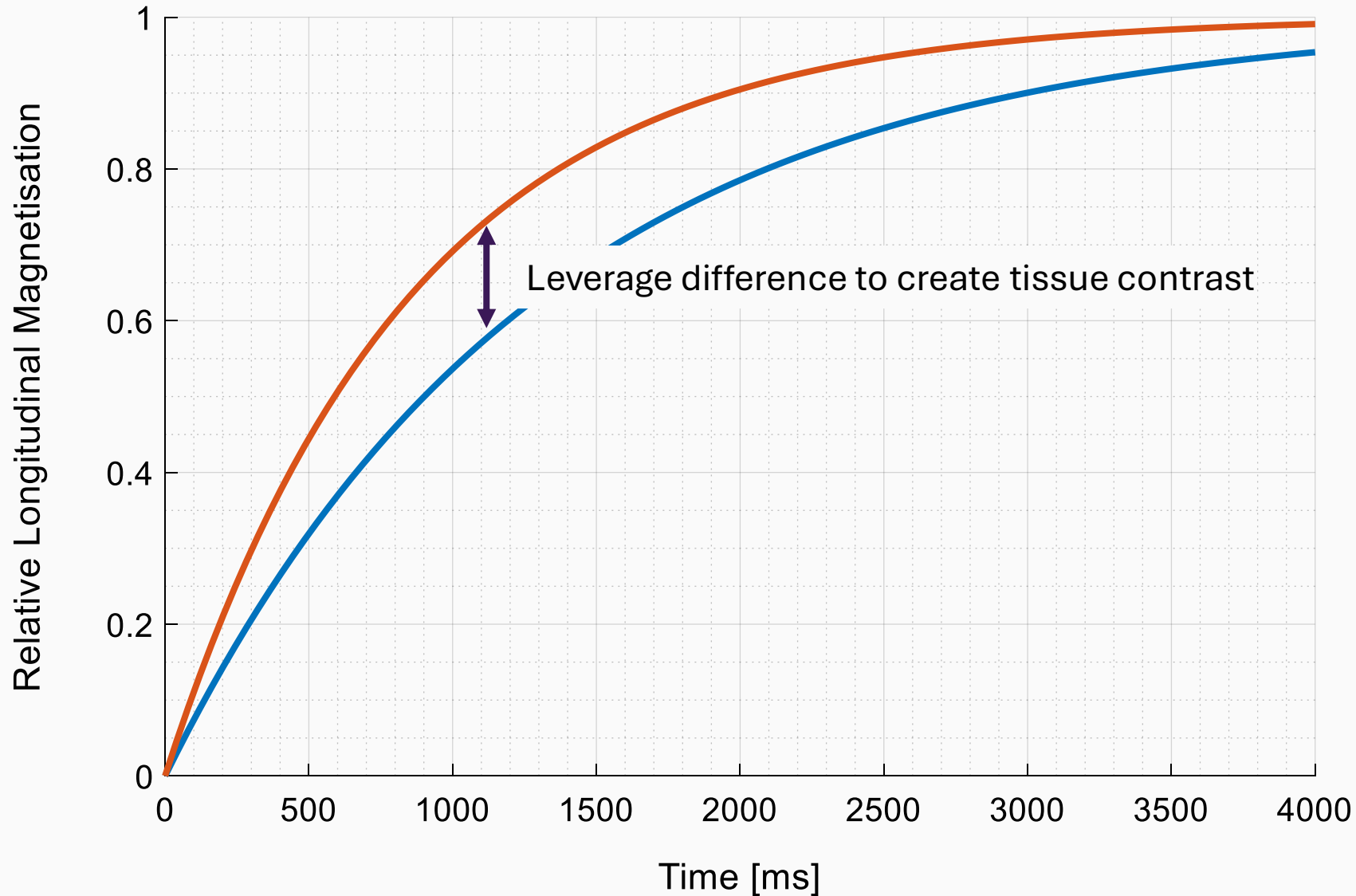
Contrast: T_1 weighting



T_1 describes the relaxation of longitudinal magnetisation

It depends on the local microstructure

Contrast: T_1 weighting



T_1 describes the relaxation of longitudinal magnetisation

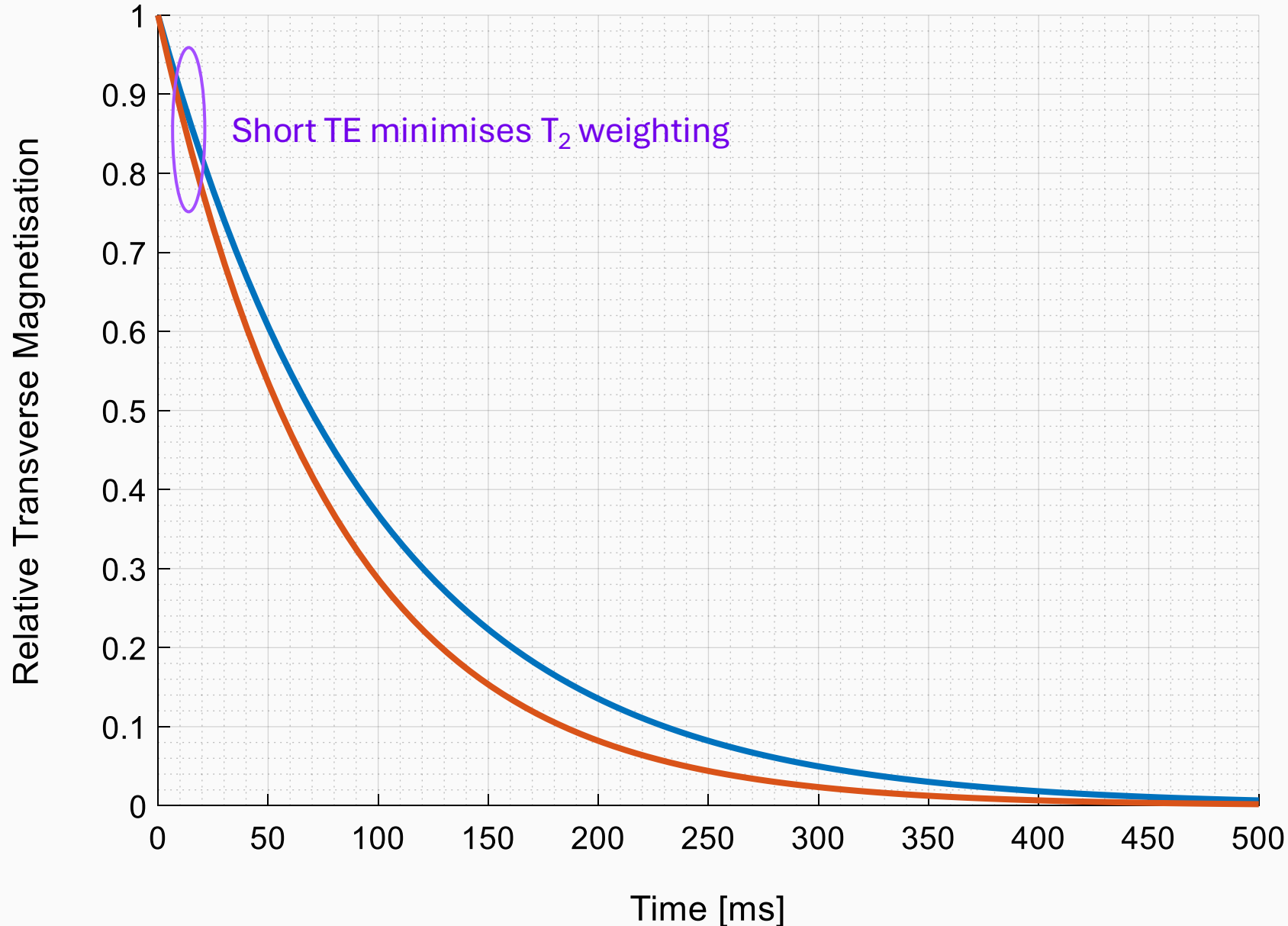
It depends on the local microstructure

At 3T:

White Matter $T_1 \approx 0.8$ s

Grey Matter $T_1 \approx 1.3$ s

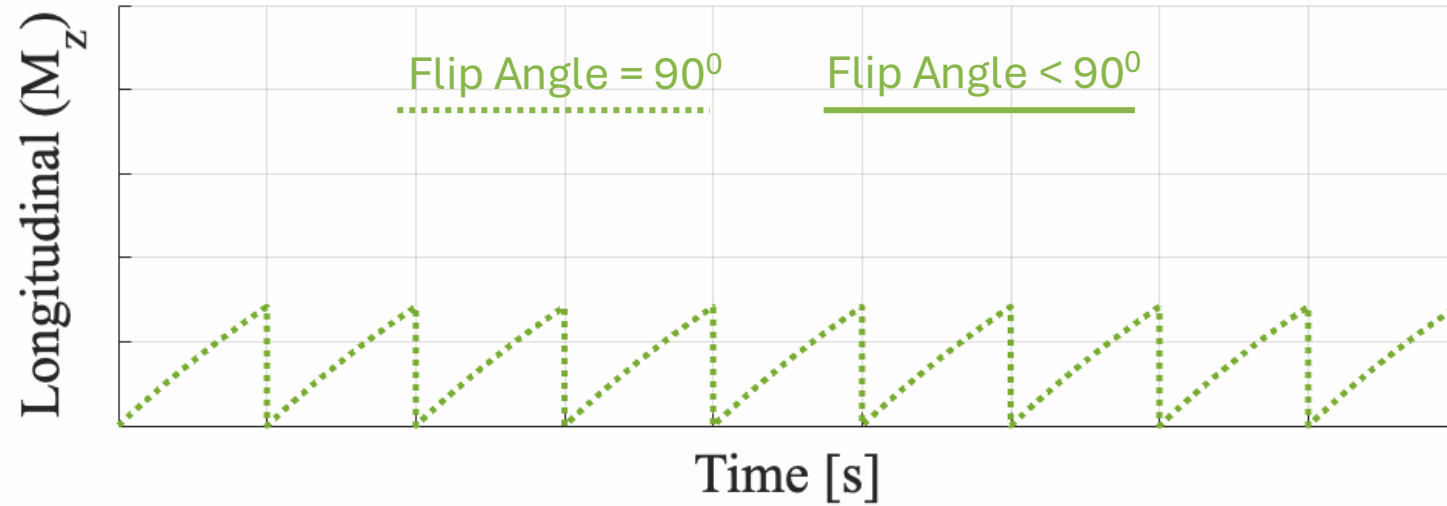
T_2^* weighting also occurs!



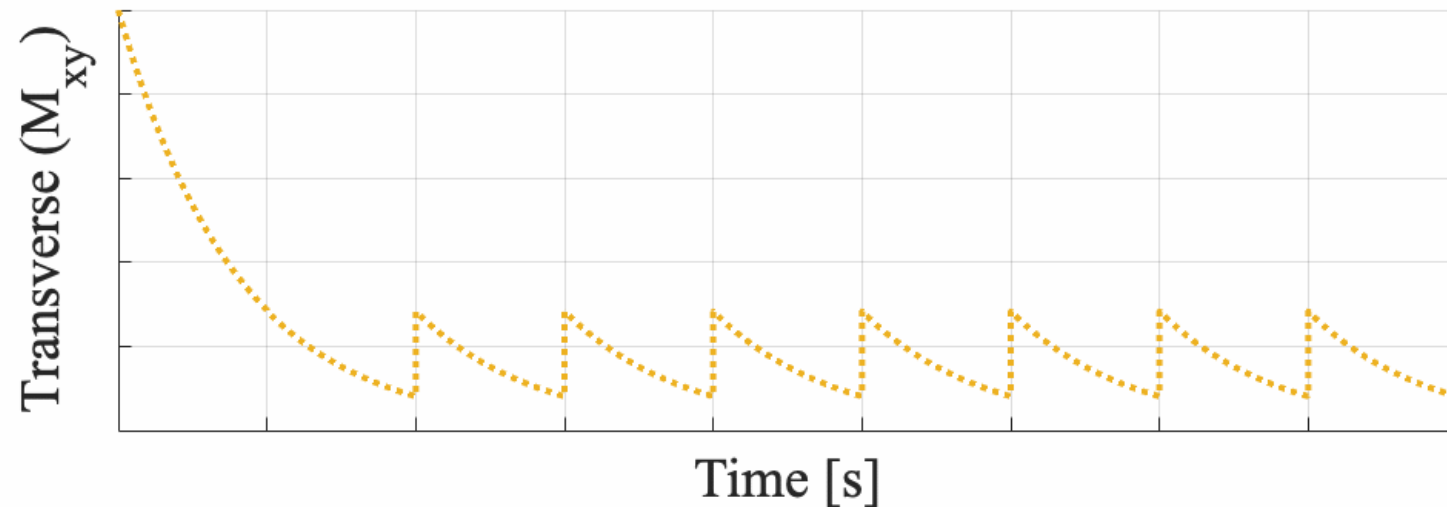
T_2^* describes how quickly transverse magnetisation decays

White Matter T_2^* < Grey Matter T_2^*

Recap: relaxometry

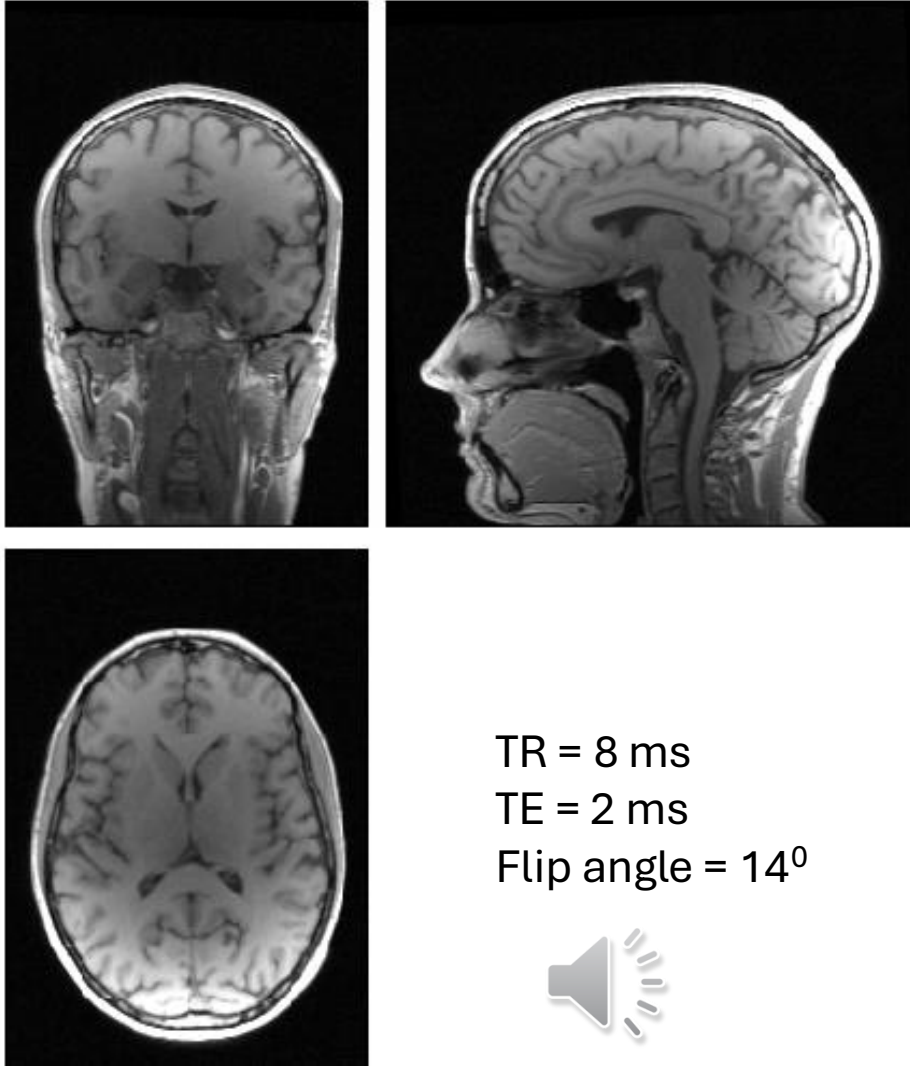


Flip Angle dictates how much longitudinal magnetisation we use each TR...



And therefore how much signal there is to sample...

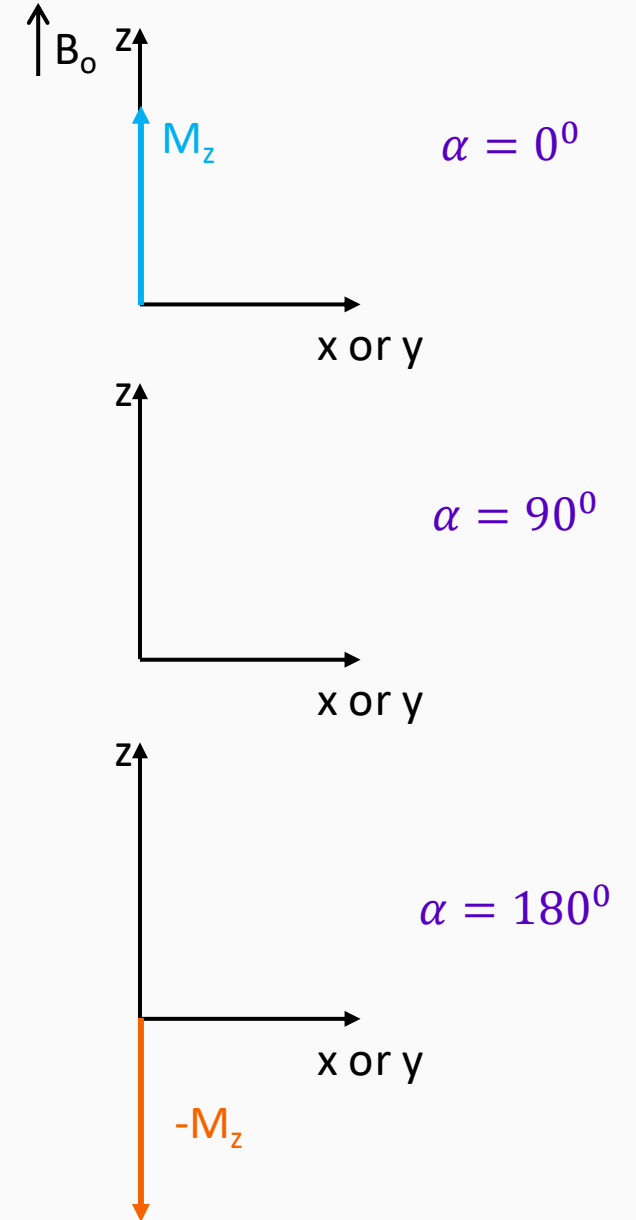
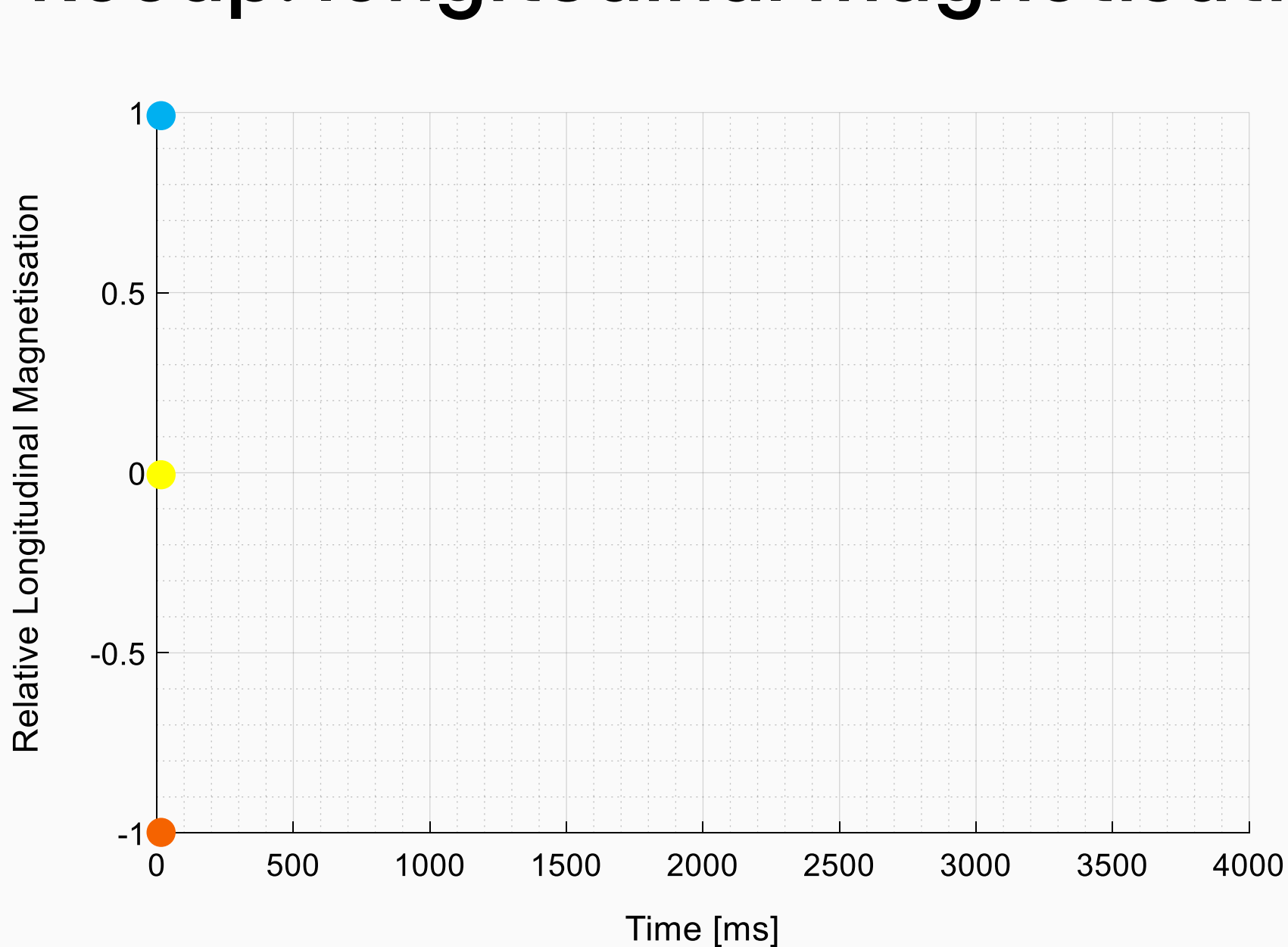
Fast Low Angle Shot = FLASH



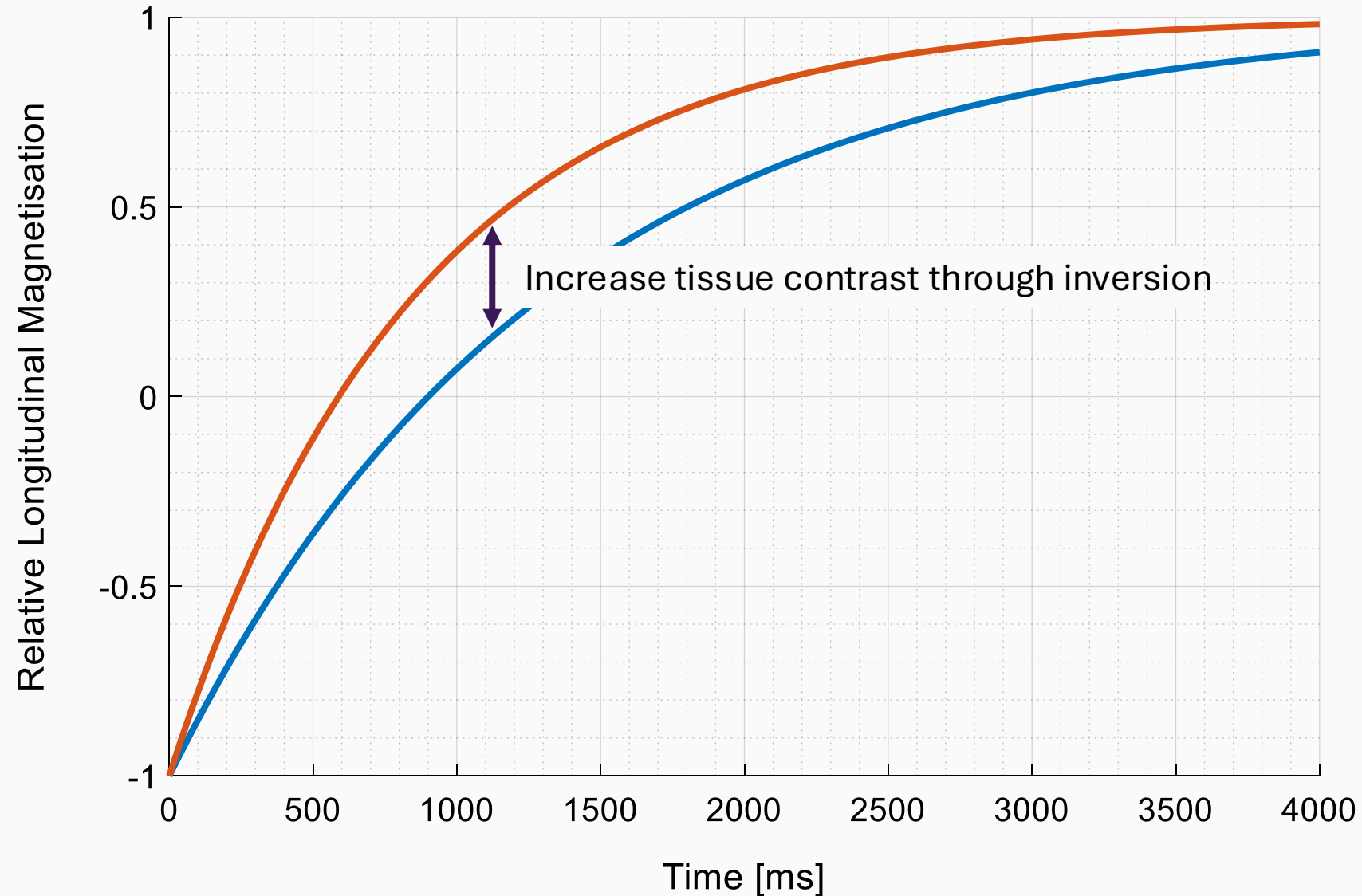
An efficient gradient echo
that combines short TR
with small flip angle

What if we want *more*
contrast?

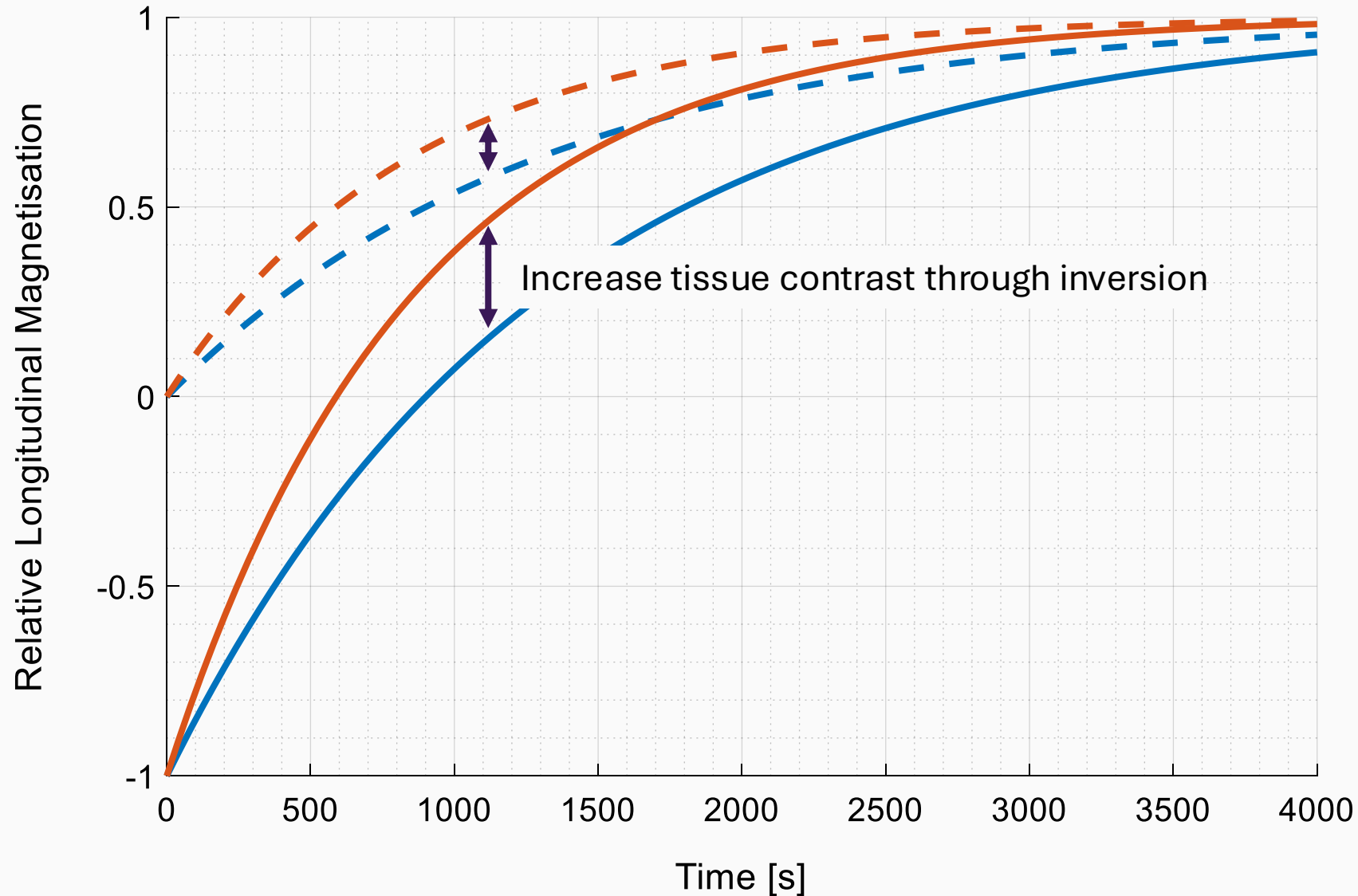
Recap: longitudinal magnetisation



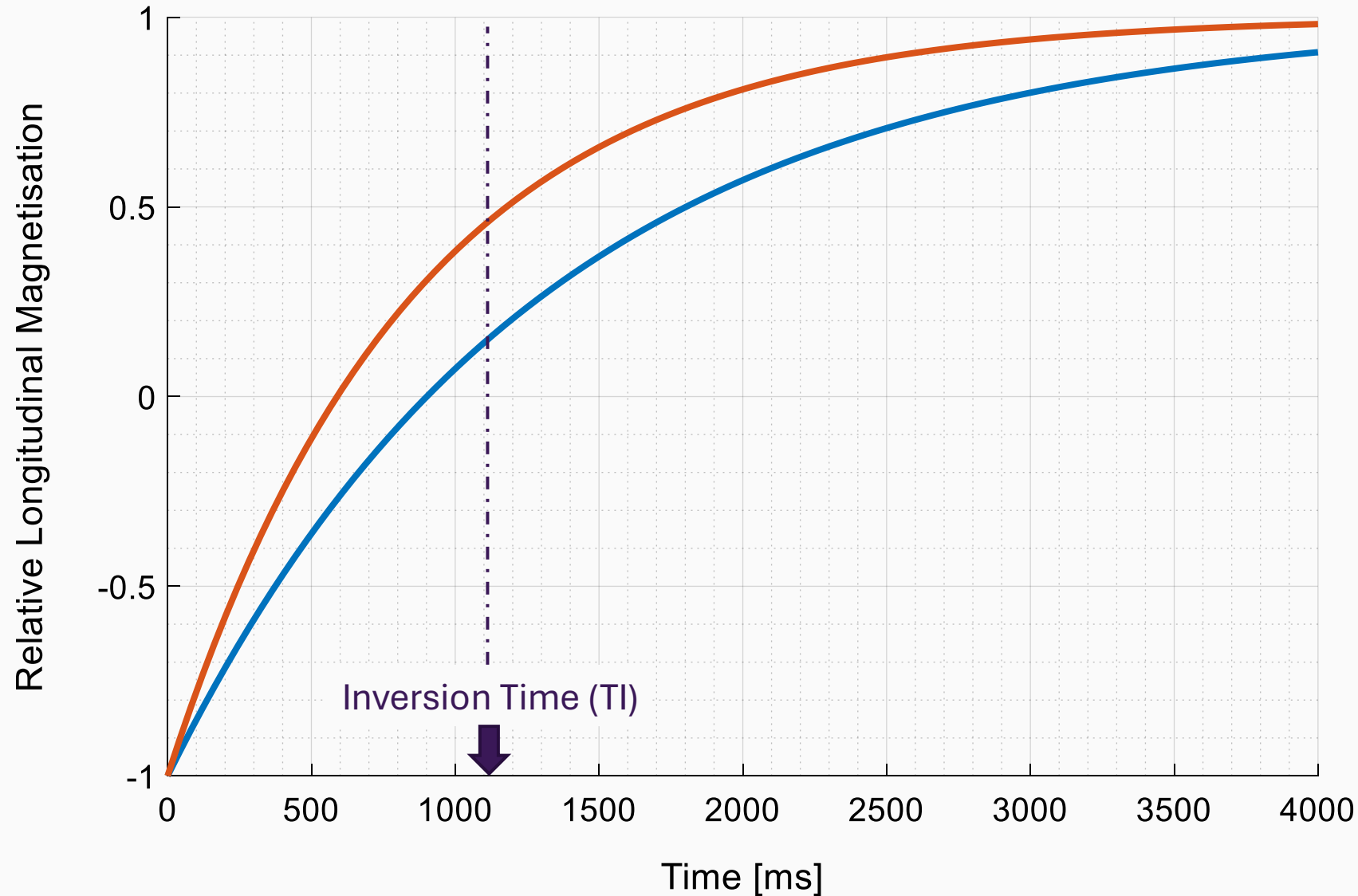
Magnetisation Preparation: Inversion



Magnetisation Preparation: Inversion

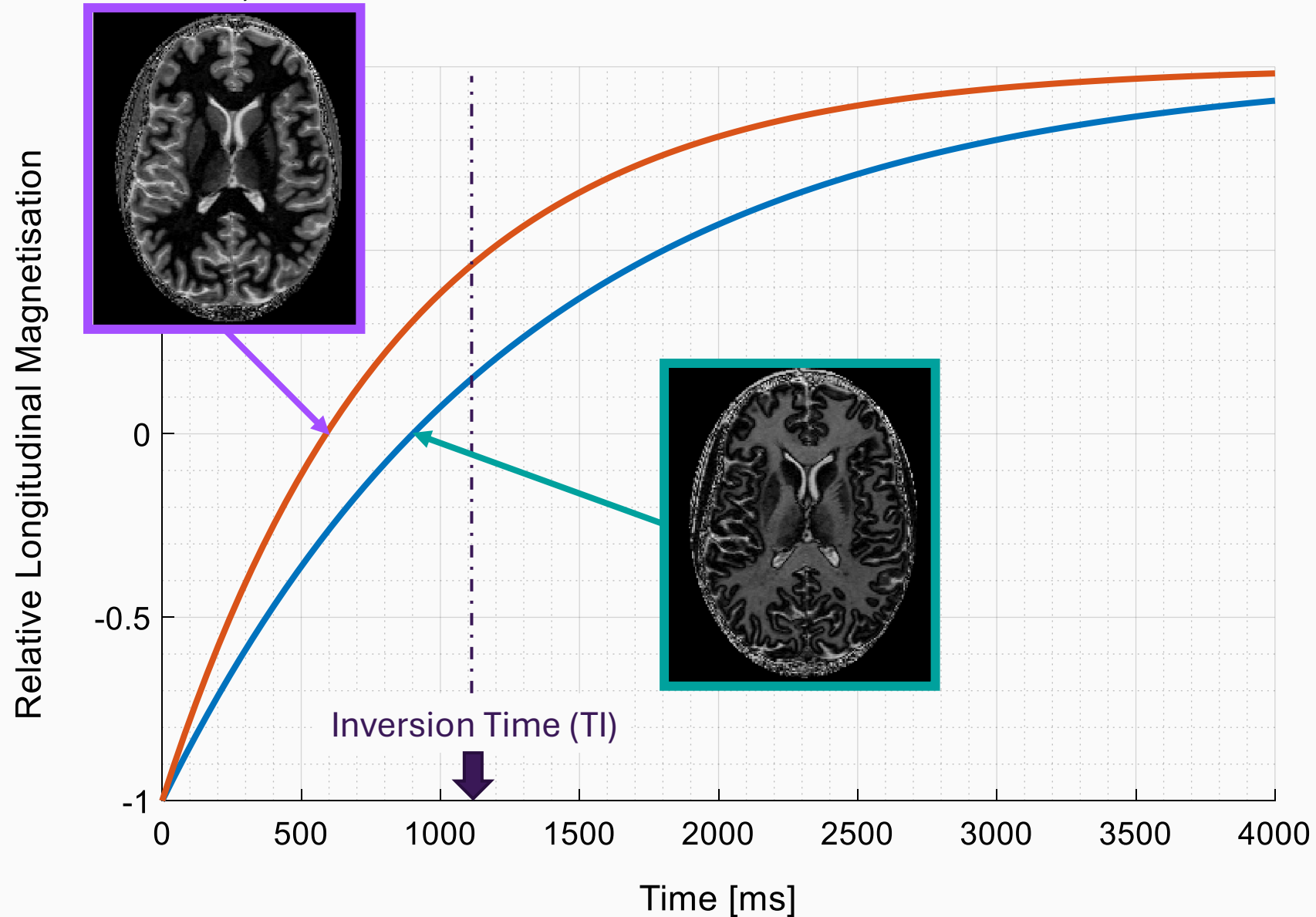


Magnetisation Preparation: Inversion



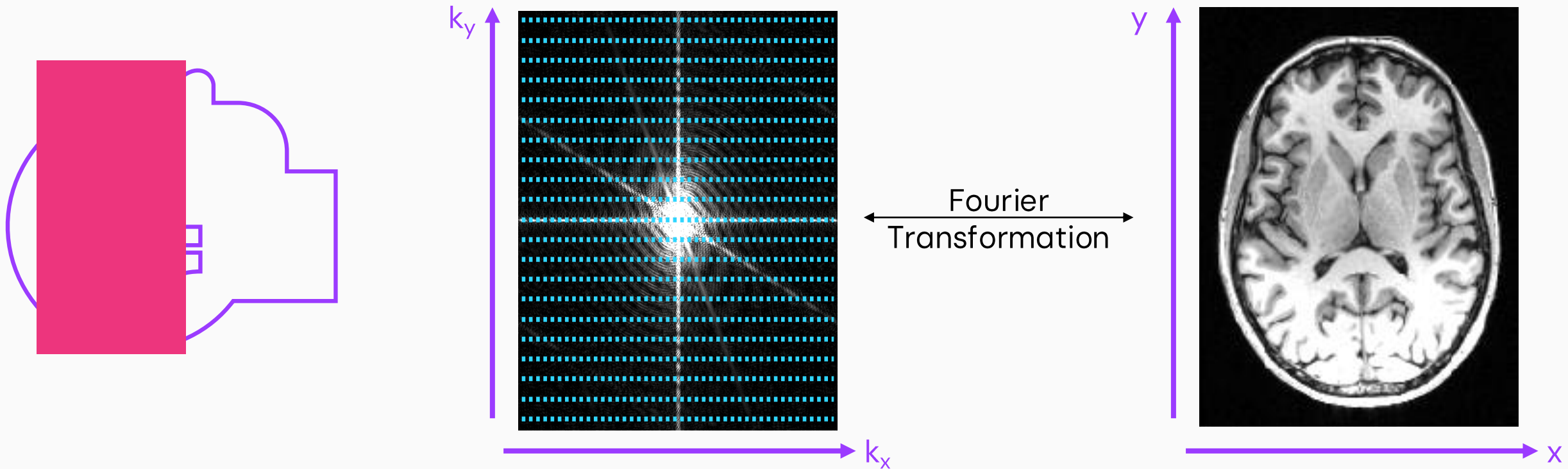
The inversion time is the time we allow contrast to develop between tissues

Magnetisation Preparation: Inversion



We can choose a TI to null a given tissue type!

Recap: filling k-space



Example: axial Field of View (FoV) = 180 mm (left-right) x 220 mm (anterior-posterior) with 1 mm resolution => 39,600 TRs!
TR ~ 10 ms => scan time of 6.6 minutes

Magnetisation Preparation

3D Magnetisation Prepared RAPid Gradient Echo:

1. Invert the magnetisation with a non-selective 180° pulse to amplify T1 based contrast
2. Wait some inversion time
3. Excite and acquire a series of gradient echoes for the inner phase-encoded direction
4. Allow further magnetisation recovery
5. Repeat for the outer phase-encoded direction

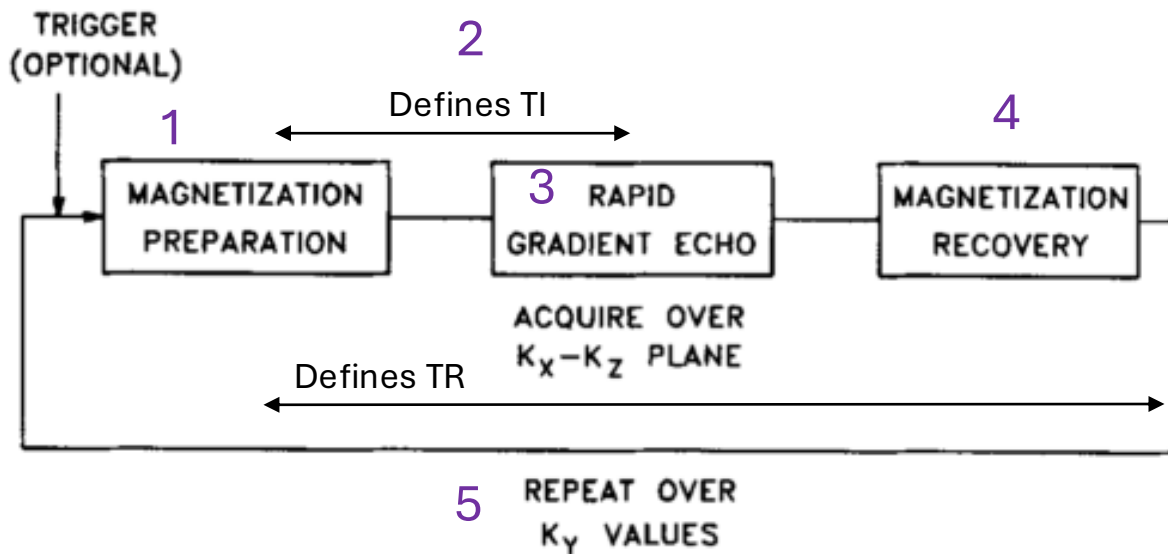


FIG. 1. Schematic diagram of the 3D MP RAGE sequence.

Magnetisation Preparation

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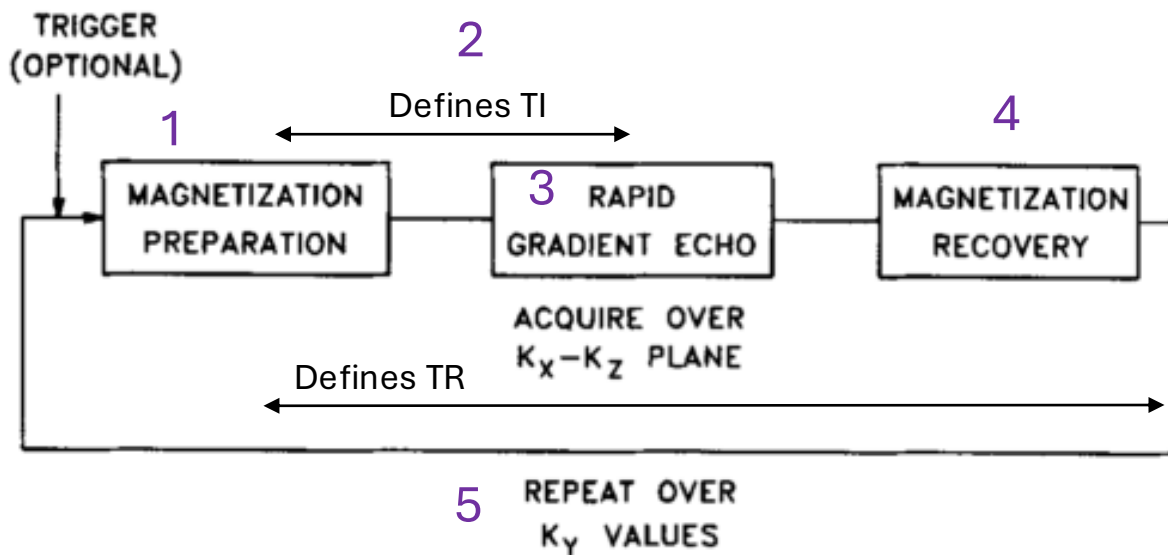
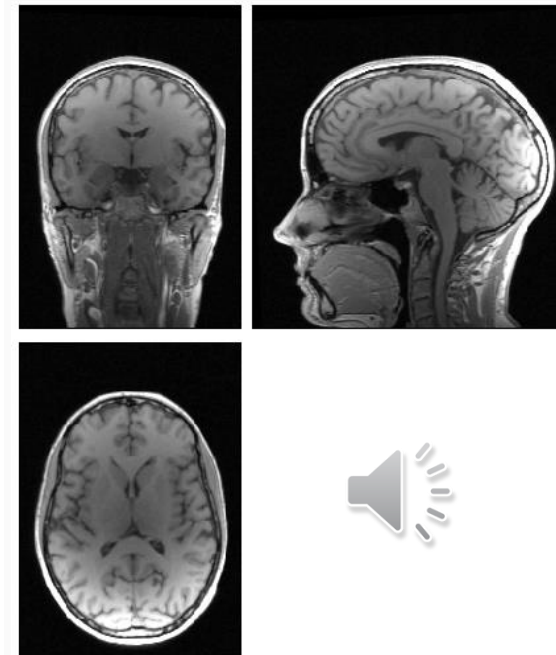


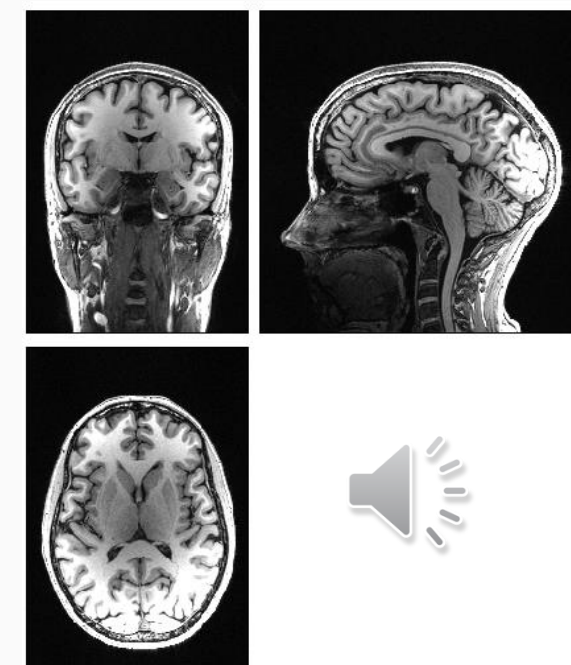
FIG. 1. Schematic diagram of the 3D MP RAGE sequence.

3D MPRAGE: Mugler & Brookeman, MRM 1990

FLASH



MPRAGE

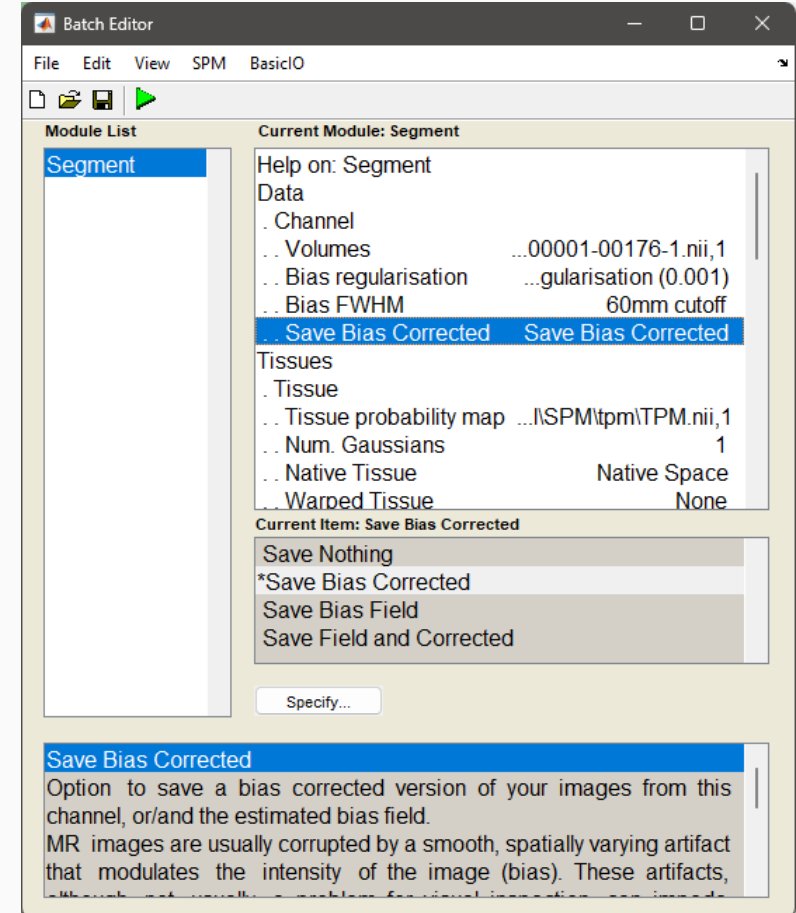
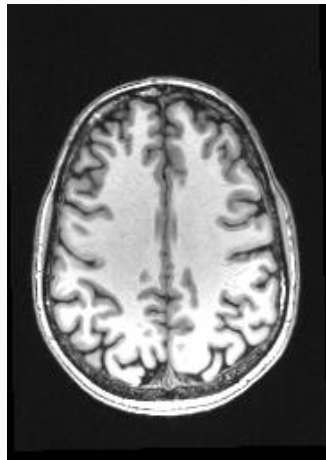


Impact of receiver coil

64
channels



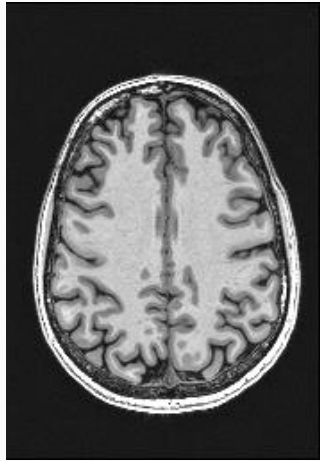
20
channels



Impact of receiver coil

Bias Field or Non-Uniform Intensity (NUI)

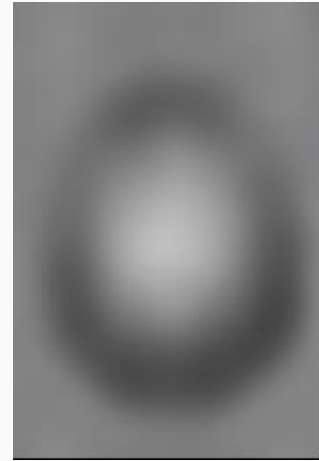
64
channels



=



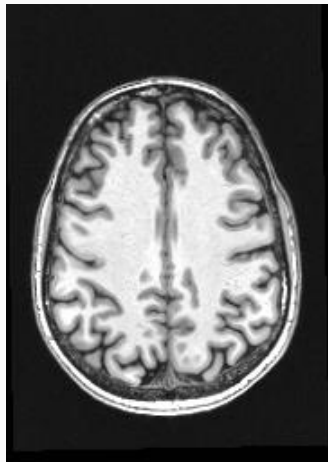
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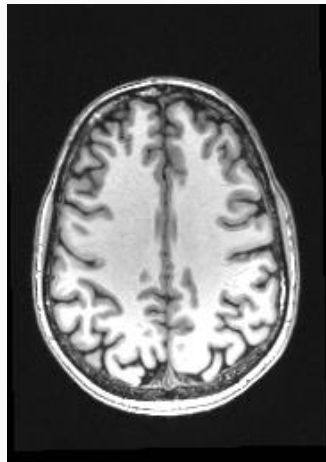
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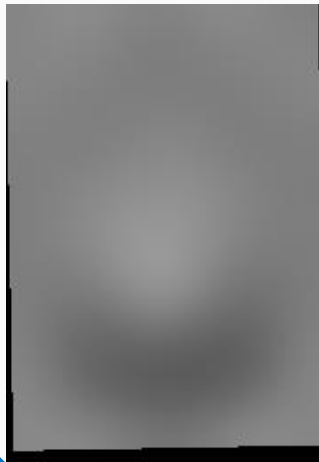
20
channels



=

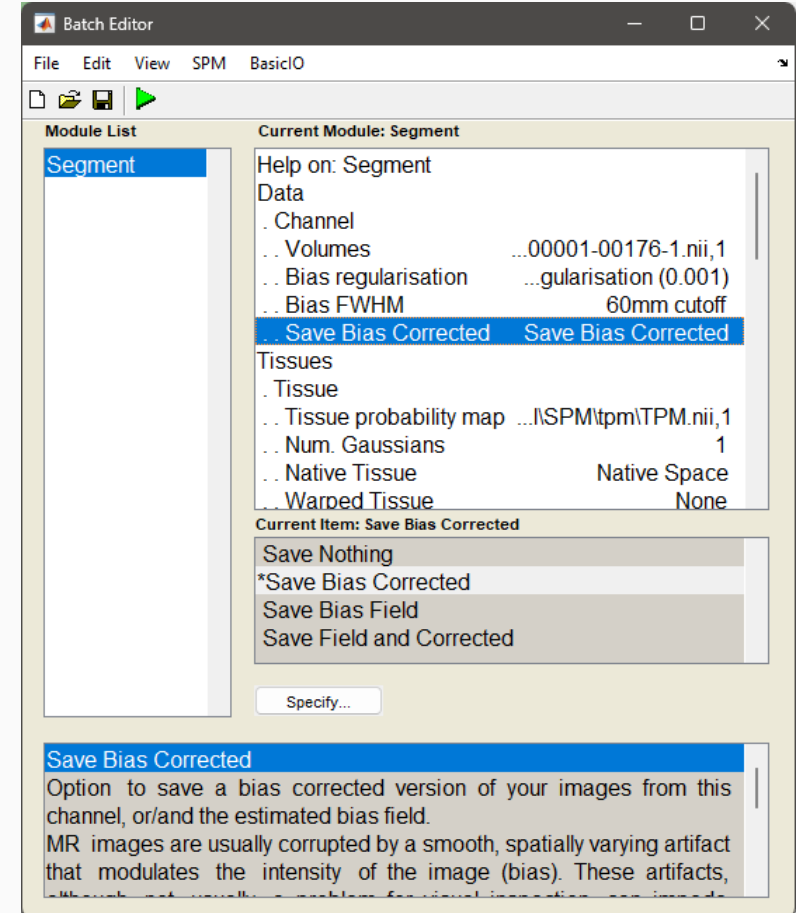


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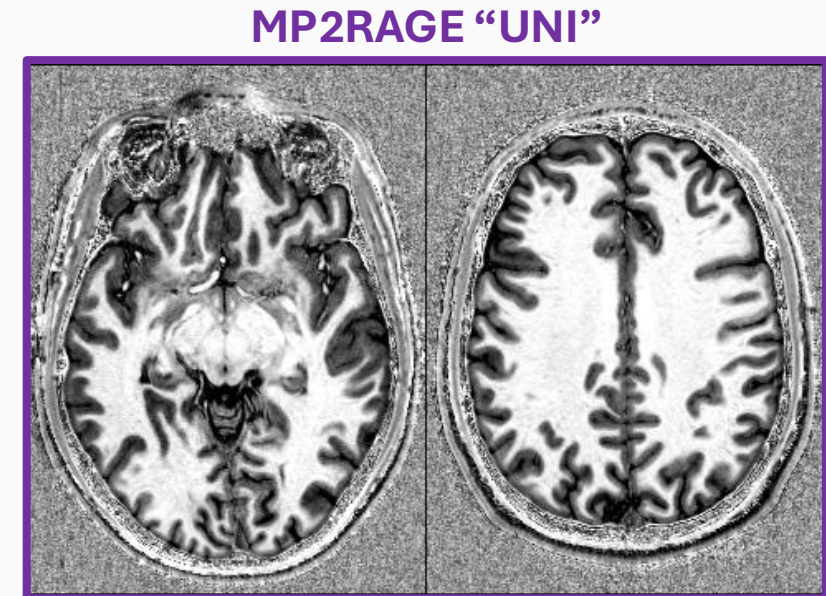
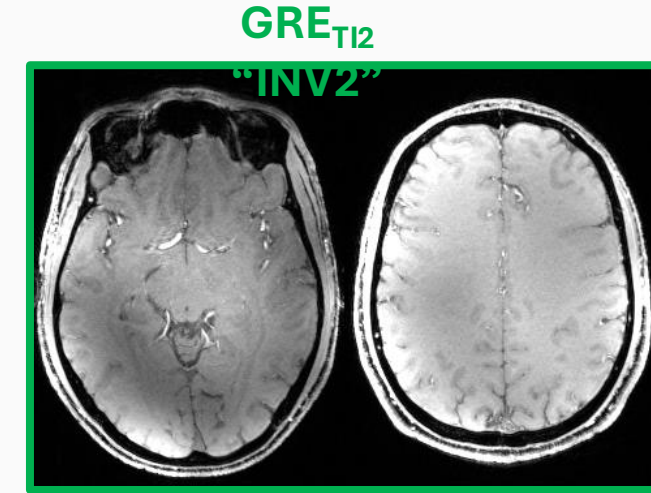
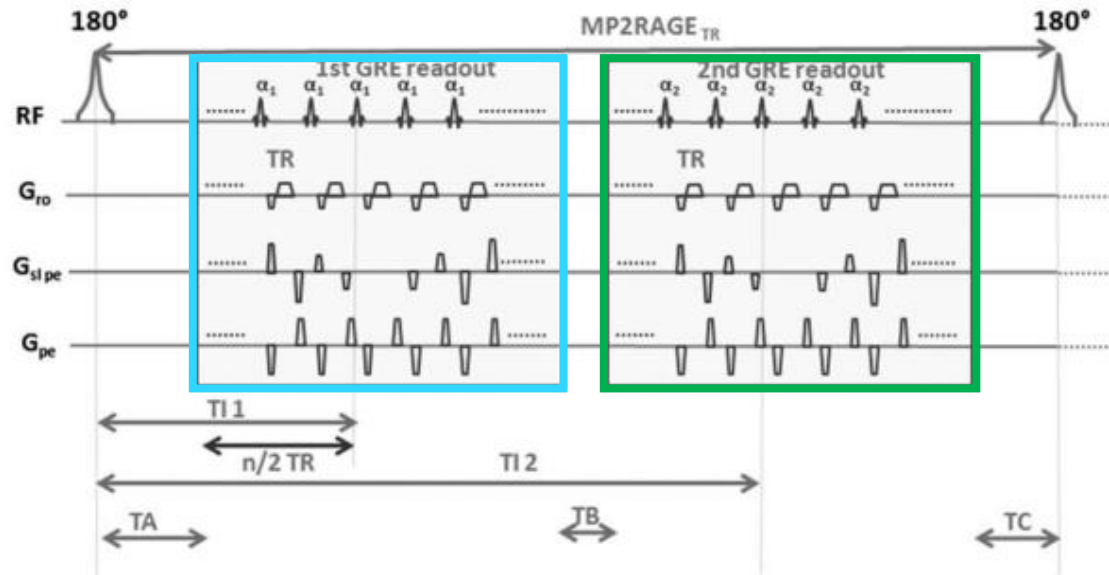


2

0

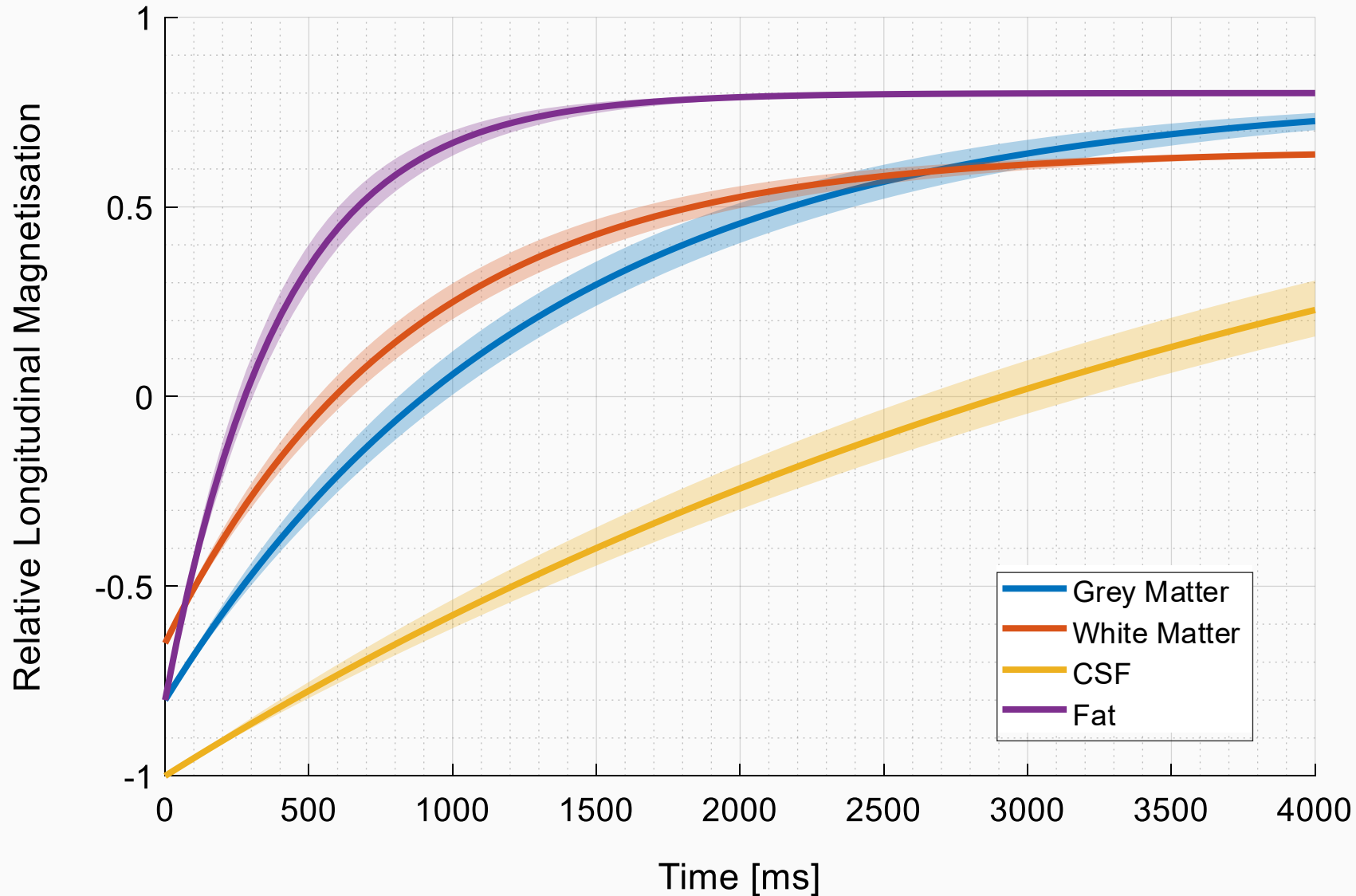


7T: MP2RAGE



$$MP2RAGE = \frac{GRE_{TI1} GRE_{TI2}}{GRE_{TI1}^2 + GRE_{TI2}^2} \in [-0.5, 0.5]$$

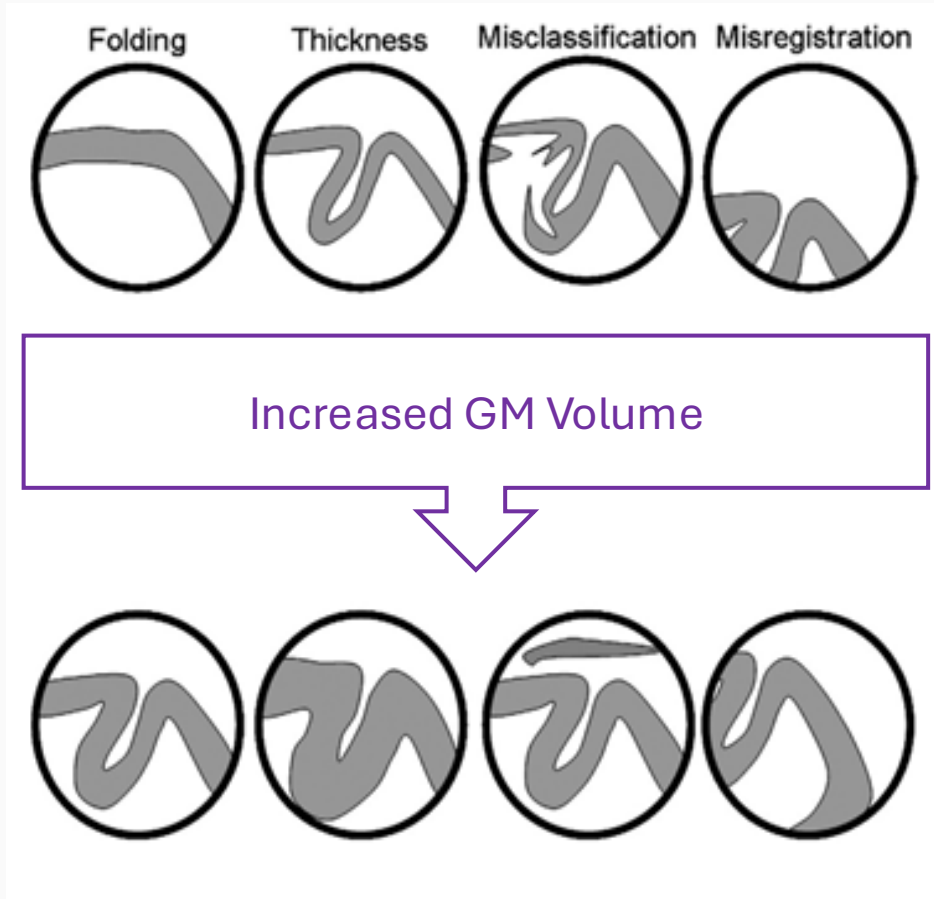
Magnetisation Preparation: Inversion



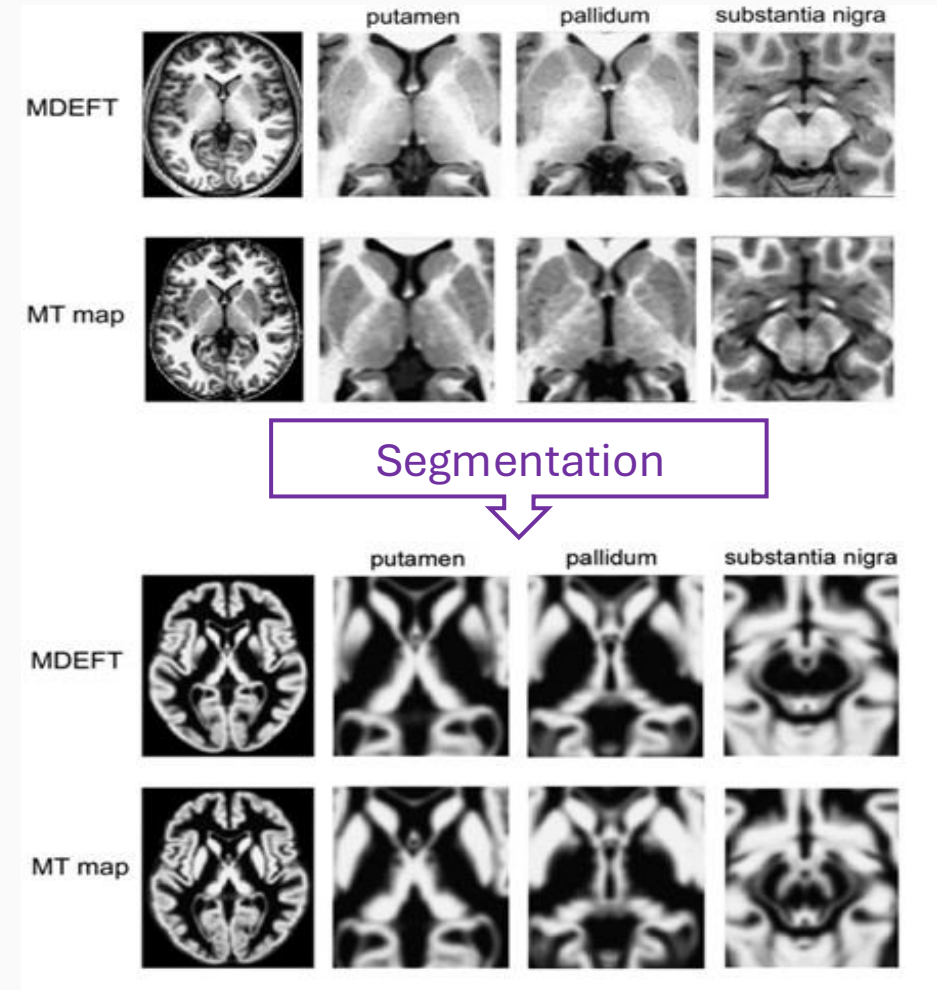
Optimisation must account for multiple tissue types, each with a range of T_1 times...

Contrast and Interpretation

Interpreting Morphological Differences



Ashburner, MRI, 2009

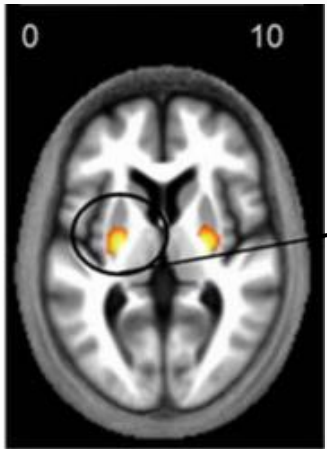


Helms *et al.*, NeuroImage, 2009

Interpreting Morphological Differences

Neurobiological Origin of Spurious Brain Morphological Changes: A Quantitative MRI Study

Lorio et al., HBM, 2016

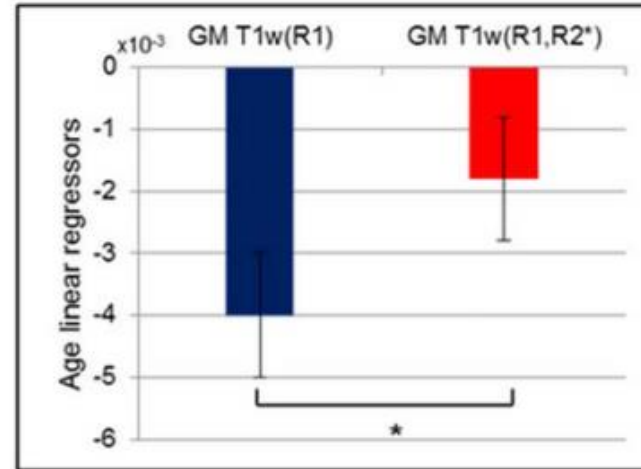
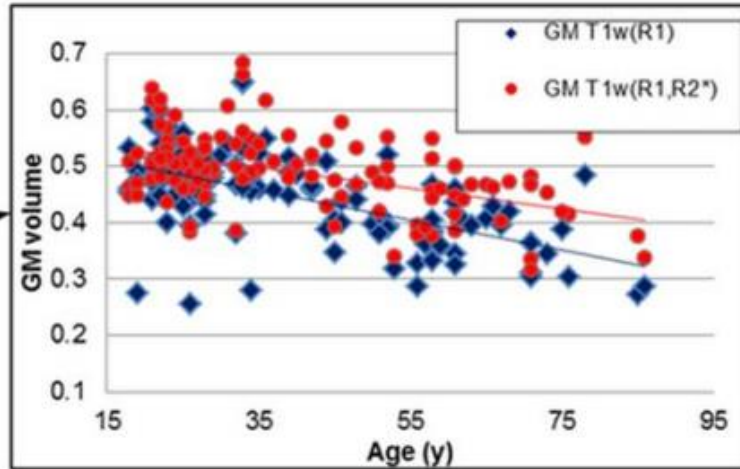
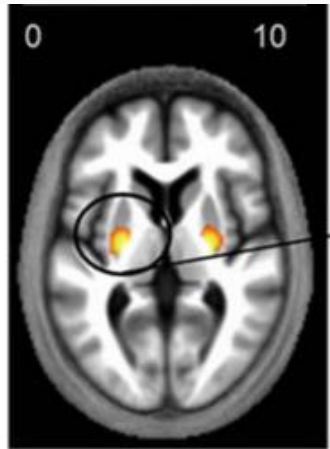


Age-related atrophy

Interpreting Morphological Differences

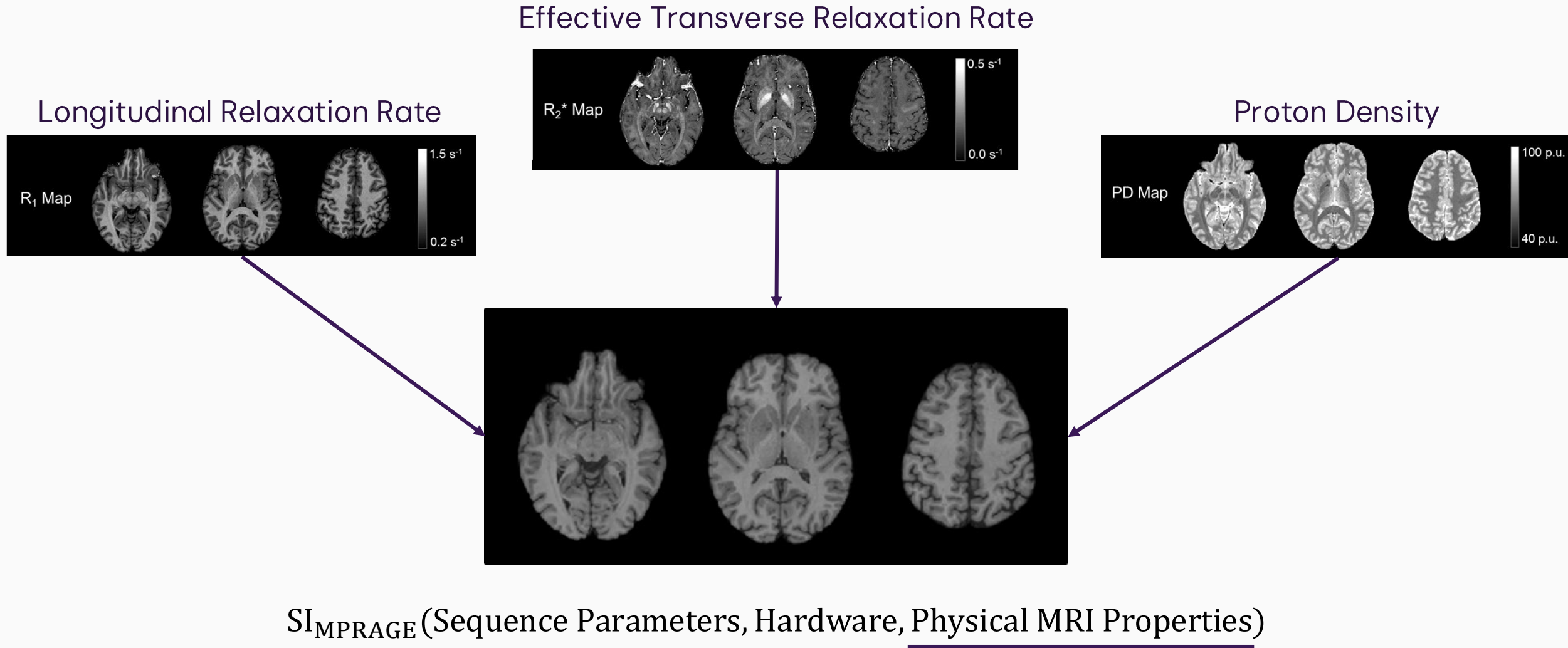
Neurobiological Origin of Spurious Brain Morphological Changes: A Quantitative MRI Study

Lorio et al., HBM, 2016

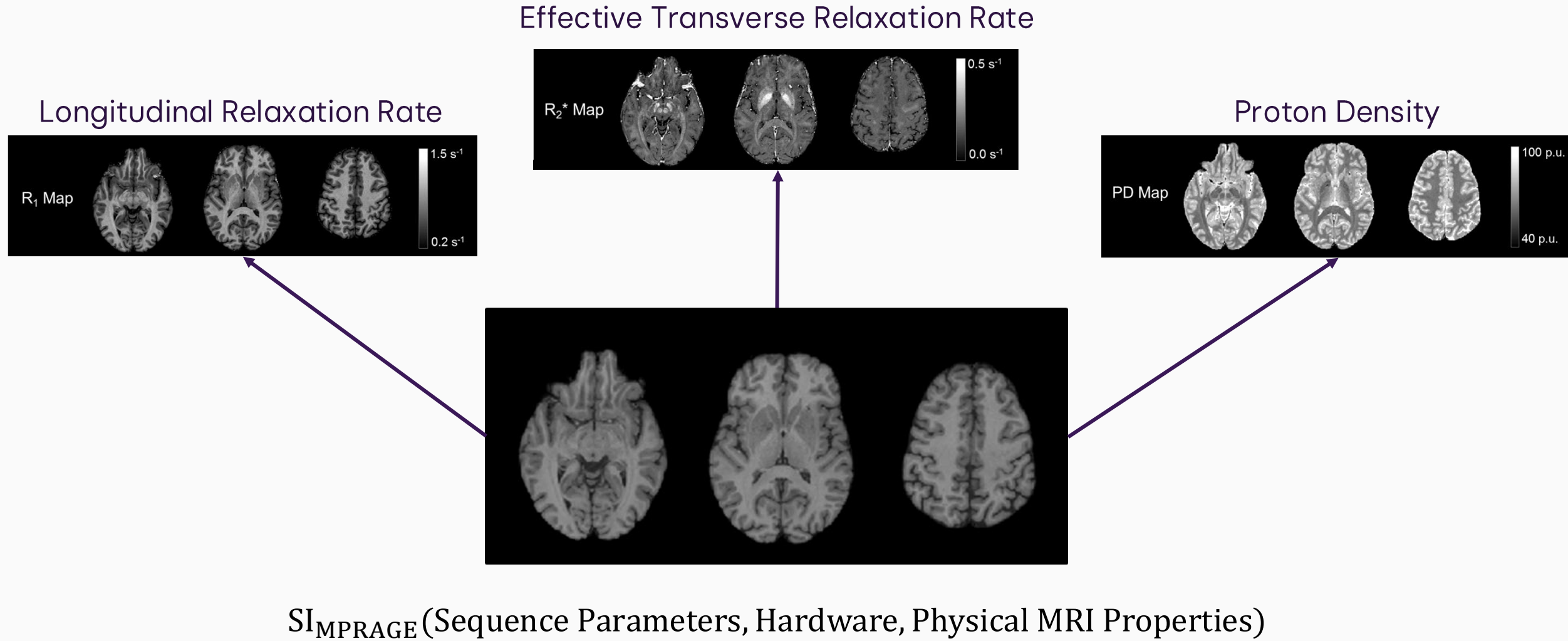


Rate of apparent age-related atrophy depends on image weighting!

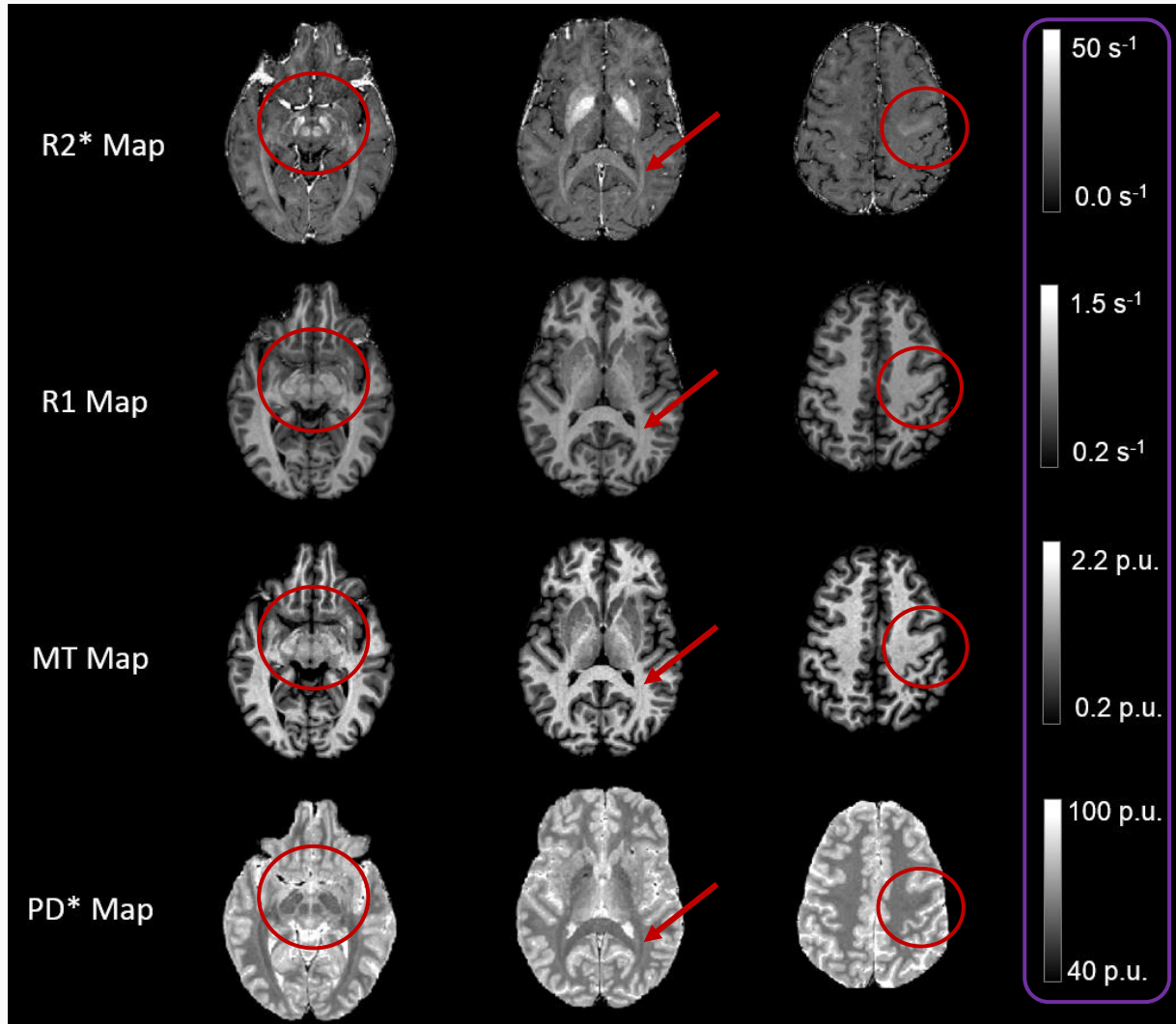
Interpreting Morphological Differences



Interpreting Morphological Differences



Quantitative MRI



Want to learn more?



Special Interest Workshop on
“Quantitative MRI”

Methods Section

MRI data acquisition

Experiments were performed on a 3 T whole-body MRI scanner (Magnetom Vida, Siemens Healthineers, Erlangen, Germany) with a 64-channel Head/Neck coil (Siemens Healthineers, Erlangen, Germany). The acquisition parameters for each of the protocols are listed in Table 1.



Fast and functionally specific cortical thickness changes induced by visual stimulation

Natalia Zaretskaya^{1,2,*}, Erik Fink¹, Ana Arsenovic^{1,2}, Anja Ischebeck^{1,2}

¹Department of Cognitive Psychology and Neuroscience, Institute of Psychology, University of Graz, Universitaetsplatz 2, 8010 Graz, Austria,
²BioTechMed-Graz, Mozartgasse 12, 8010 Graz, Austria

Table 1. Protocol overview.

Protocol type	MPRAGE (fMRI analysis)	2D GE-EPI
Isotropic voxel size (mm)	1	3
TR (ms)	2,530	3,220
TE (ms)	3.88	32
TI (ms)	1,200	n/a
flip angle	7°	82°
Matrix	256 × 256	76 × 76
Slices	176	46
PE	A>>P	A>>P
Acceleration	GRAPPA 2	None
Duration (min)	6:03	8:22
Partial Fourier	Off	Off
Bandwidth (Hz/pixel)	190	1,828

TR, repetition time; TE, time to echo; TI, inversion time.

Thank you! Any Questions?