



An automated tool for fast quality analysis of animal MRI

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Objectives

Which dataset is of sufficient quality for further processing?

- > Standardization of image quality in preclinical, animal MR imaging.
- > Categorization of good vs. bad quality datasets

Structure How is this pipeline structured? Input Data Stage (I): Parsing Stage (II): Feature Calculation Stage (III): Plots 1. SNR 1. Parse all MR files QCplot.py 2. tSNR 2. Extract T2w, DTI, fMRI data Excel: MR PDF: ParsingAllrawData.py **Excel: Results** Raw **Bruker** MR Data CheckingFeatures.py 3. Movement variability Histograms 3. Create an excel sheet of all **Data Path** 4. Spatial resolution QCtable.py the available MR data homogeneity

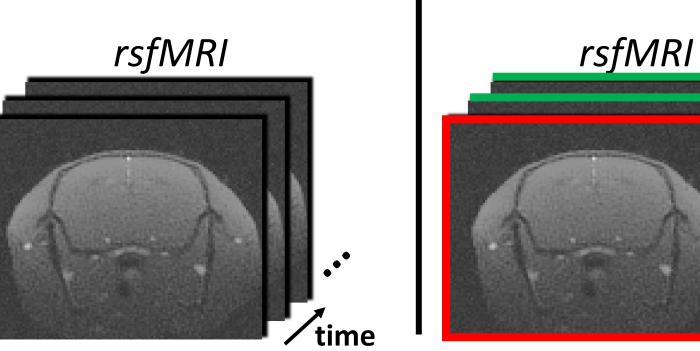
Python 3.8.12 (Libraries used: NumPy, Pandas, Matplotlib, Nibable, Alive_progress, Os, Glob, ...)

Methods

SNR: Method (Fully Chang automated approach without the need to define regions of interest)

$$tSNR = \frac{\mu_{SNR}}{\sigma_{SNR}} = \frac{\mu}{\sqrt{1/N\sum_{i=1}^{N} (I_i - \mu)^2}}$$

SNR & tSNR



Movement Severity

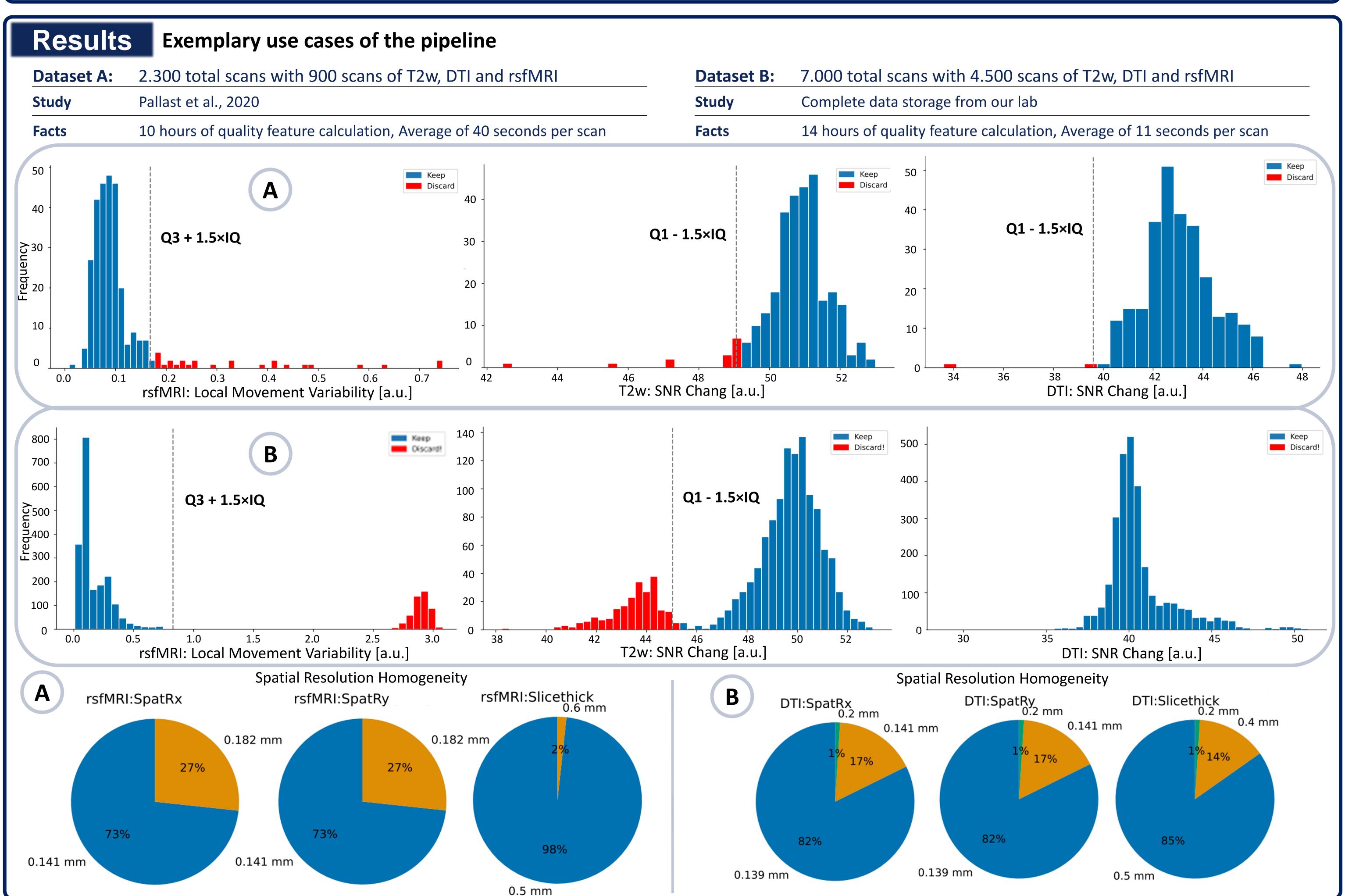
Mutual information between $I(t_0)$ and $I(t_0+t)$ for $\mathbf{t} = \mathbf{1}, ..., \mathbf{t}_{N}$ resulting in \mathbf{N} values or a mutual information vector.

Standard deviation of the mutual information series is the metric of movement severity

Categorization in good vs. bad quality

Based on the definition of **outliers** via **IQR rule**:

- Lower than Q1 1.5×IQ
- Higher than **Q3** + **1.5**×**IQ**



Outlook What are some ideas for the future?

- 1) Increasing sequence variety in addition to T2w, DTI, fMRI sequences 2) Further improving calculation speed via implementing parallel computing 3) Creating compatibility with Bruker's ParaVision software to directly compare quality features of new data with the main data storage
- The pipeline is available on GitHub under: https://github.com/aswendtlab/AIDAqc
- Necessary information on how to install the pipeline is available in addition to some examples and a test dataset

