A generalizable approach to quality assessment of voxel-based morphometry data

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Introduction

We introduce and evaluate an automated quality assessment approach based on low-rank representations via Orthonormal Non-negative Matrix Factorization (OPNMF)[1] of preprocessed structural MRI data. OPNMF produces:

- W matrix - sparse spatial distribution of the components
- H matrix - loading coefficients for each image

Aim: to provide a method for detection of aberrant images, which is:

1. **Generalizable**
2. **Data-driven** (i.e. no manual labels required)

The performance of the proposed approach is assessed as:

1. "Outlier detection", i.e. within-sample factorization
2. "Quality control", i.e. out-of-sample factorization - generalizability

Methods

Datasets

I. Outlier detection paradigm
1000BRAINS[2] dataset (n=1,324; age-range 18-85)

II. Quality control paradigm
multi-site clinical cohort (334 schizophrenia patients, 372 healthy controls; age-range 18-65; 13 sites)

Preprocessing

- CAT12[3] with standard settings
- DARTEL-normalization and tissue segmentation
- Unsmooth, non-linear modulated only GM segments are used.
- Manual visual assessment by two raters of both datasets

Analysis outline

Flagging protocol
(1) Calculation of Quality metrics (z-scored):
- **Component loadings** (**CL**; from H matric)
- **Whole-brain reconstruction error** (**RE**; from W and H)
- **ROI reconstruction error** (**ROI-RE**; maximal taken).

(2) Any image with at least two flags of |z-score|>2 is suspicious of artifacts.

I. Outlier detection paradigm (within-sample factorization)
(0) OPNMF is performed on 1000BRAINS
(1):(2) Perform Flagging protocol

II. Quality control paradigm (out-of-sample factorization)
(0) Factorization from 1000BRAINS is transferred to clinical sample.
(1):(2) Perform Flagging protocol

III. Evaluate performance against manual visual inspection

Results

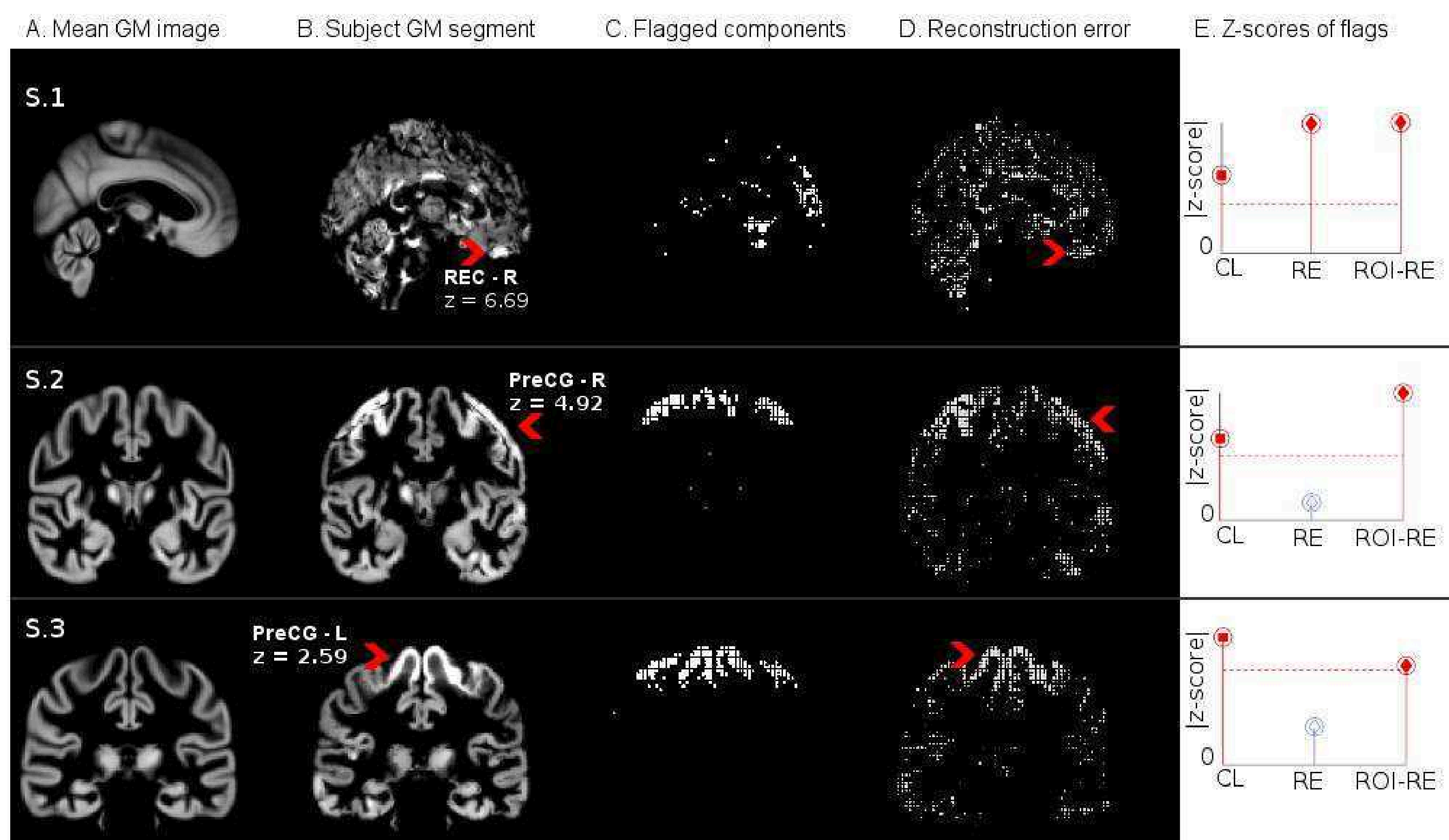### Flagged images in outlier detection paradigm (1000BRAINS)

Figure 1. Examples of images correctly flagged as "Bad". The individual rows illustrate examples of images flagged as suspicious by at least two flags. Column A displays the mean GM image for the whole sample, for use as a visual reference. In column B there are the actual GM segments of interest. The red arrows indicate the ROIs with the largest |z-score|. Column C illustrates the voxels belonging to flagged components (i.e. components with |z-score| of at least 2.5); in column D is the reconstruction error for the voxels; and in column E the z-scores of each flag are plotted. Abbreviations: REC-R – Gyrus rectus (right hemisphere); PreCG-R/L – precentral gyrus (right/left)/

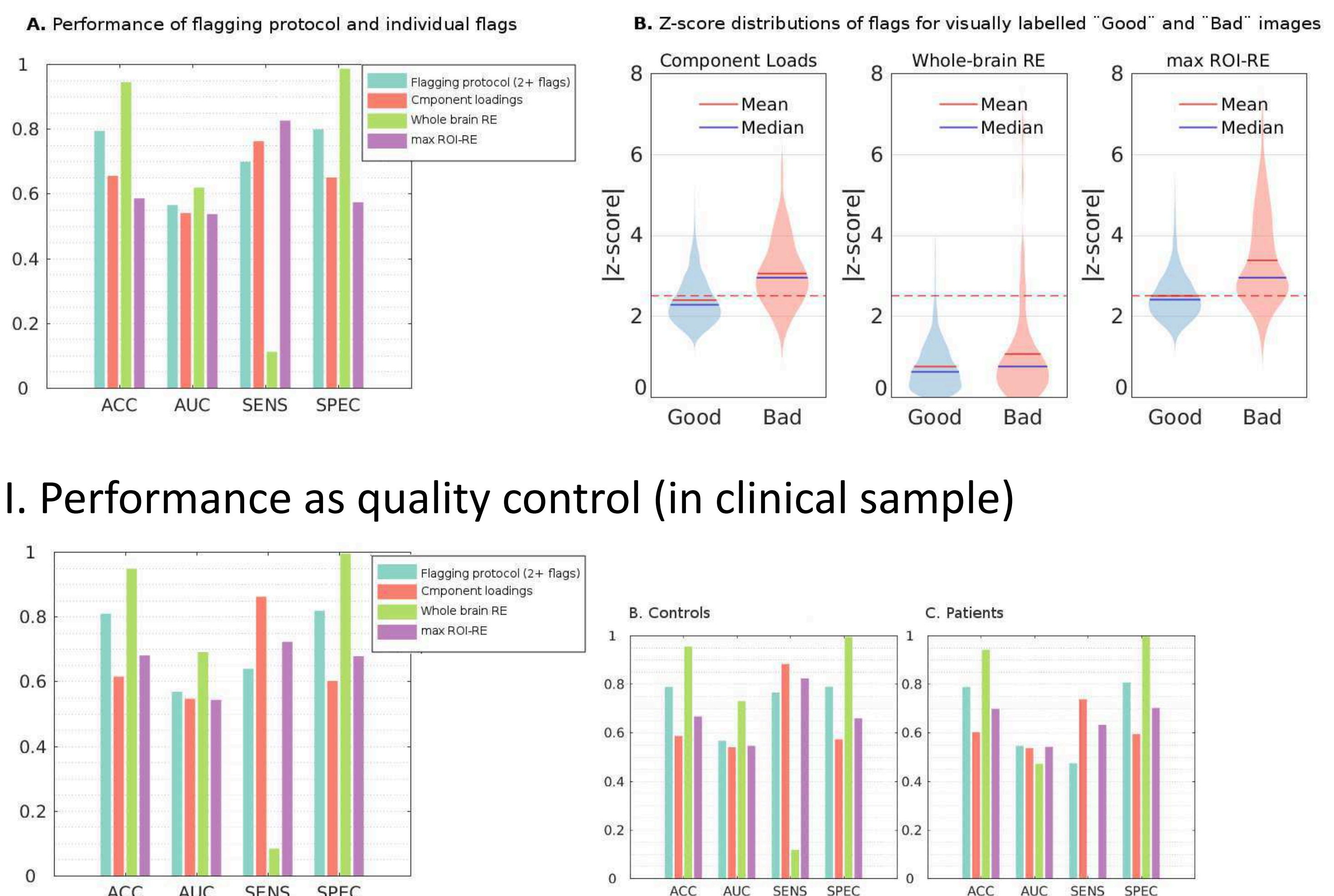### I. Performance as outlier detection in healthy sample (1000BRAINS)

II. Performance as quality control (in clinical sample)

Figure 2. Performance of the flagging protocol when applied in the outlier detection paradigm (**I**) and when applied as quality control (**II**). The violin plots on **I.B.** illustrate the separation between the images manually labelled as "Good" and "Bad" by the human raters, as measured by each of the individual flags – component loadings, whole-brain reconstruction error, and maximal ROI reconstruction error.
Figure **II.A.** shows the performance of the flagging protocol on the whole clinical sample (i.e. when applied to controls and patients pooled together). The performance of the protocol when applied to the two groups separately is shown on Figure **II.B.** and **C.**
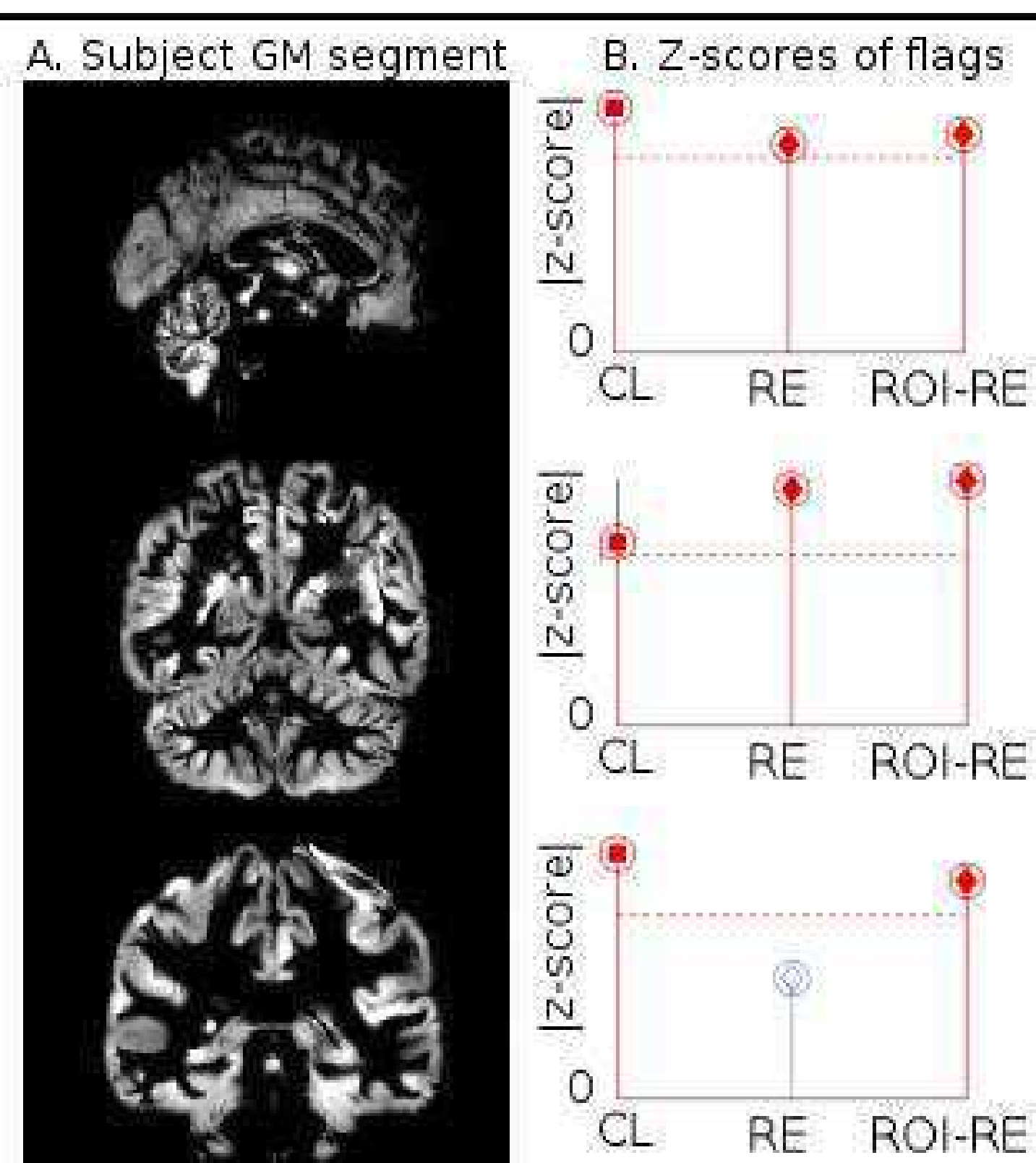
Images missed by the human raters

Figure 3. "False positive" images containing artifacts, i.e. real true positives. These are examples of three individual images detected by our automatic flagging protocol but missed by the human raters. The left column A shows GM slices at the coordinates of the voxel with the highest reconstruction error for each respective image. The scores of the individual flags are plotted in column B.

Discussion

Our method is applicable to any large, already preprocessed VBM data at a low computational cost. The evaluation showed that:

- The component loadings and the ROI reconstruction errors can be used to flag aberrant images with good performance compared to manual ratings.
- Our method allows generalization to new datasets to identify aberrant images with good performance, especially on healthy subjects.
- Our method detects artifacts human raters can miss.

Data-driven techniques eliminate the dangers of human error and subjectivity. Unsupervised methods such as OPNMF have the advantage of not requiring training, yet are generalizable to new, unseen data.

References:
1. Sotiras et al.(2014): Finding imaging patterns of structural covariance via Non-Negative Matrix Factorization. Neuroimage 108:1–16. http://dx.doi.org/10.1016/j.neuroimage.2014.11.045.
2. Caspers S et al.(2014): Studying variability in human brain aging in a population-based German cohort-rationale and design of 1000BRAINS. Front Aging Neurosci 6:1–14.
3. Gaser & Dahnke (2012): CAT - A Computational Anatomy Toolbox for the Analysis of Structural MRI Data. HBM Conf 2012 32:7743.