Tissue-Class Segmentation

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Image segmentation

- We are often interested in subdividing or segmenting the brain into meaningful biological regions of interest (ROIs) for an analysis.
- Examples: tissue segmentation, segmentation of gray matter structures, segmentation of pathology (MS lesions, tumors, . . .)

Goals of this tutorial

- ▶ Perform tissue segmentation in R using FSL and ANTs.
- ▶ Discuss multi-atlas label fusion techniques for segmentation.
- ▶ Perform automatic MS lesion segmentation using OASIS.

Loading Data

▶ Let's read in the training T1s and brain mask for subject 05.

```r
library(ms.lesion)
library(neurobase)
library(fslr)
library(scales)
library(extrantsr)
all_files = get_image_filenames_list_by_subject(
  group = "training",
  type = "coregistered")
files = all_files$training05
t1 = readnii(files["MPRAGE"])
mask = readnii(files["Brain_Mask"])
# t1s = lapply(files, function(x) readnii(x["MPRAGE"]))
# t1 = t1s[[5]]
# masks = lapply(files, function(x) readnii(x["Brain_Mask"]
# mask = masks[[5]]
```

Tissue Segmentation: Large Outliers

- Many tissue class segmentations are based on k-means clustering.
- These methods can be skewed by large outliers.

```r
hist(t1, mask = mask, breaks = 2000)
```
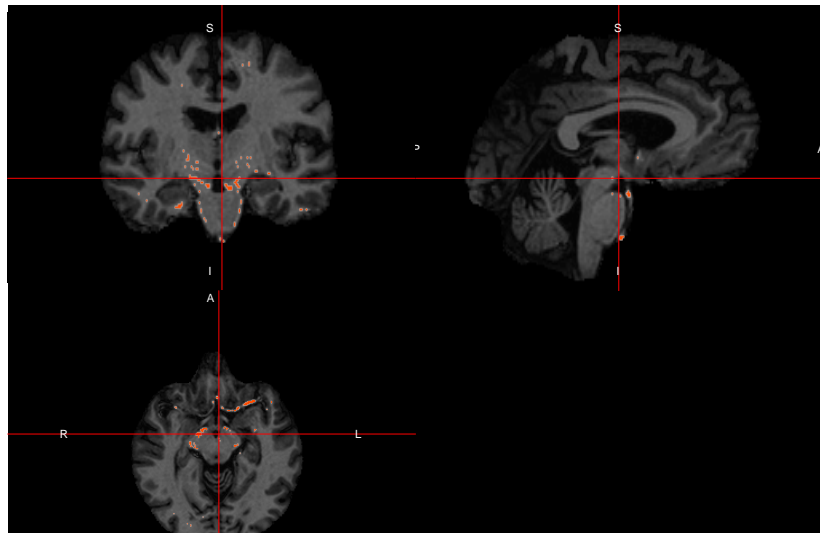
Histogram of x
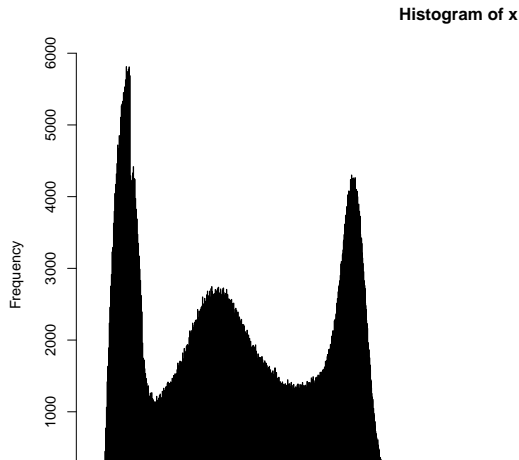
Where are the outliers

We see some values that may have been improperly segmentated:

```
ortho2(t1, t1 > 400, xyz = xyz(t1 > 400))
```

Tissue Segmentation using FSL FAST

- FAST is based on a hidden Markov random field model and an Expectation-Maximization algorithm.
- It jointly produces a bias field corrected image and a probabilistic tissue segmentation.
- More robust to noise and outliers than finite mixture model-based methods that do not incorporate spatial information.

Tissue Segmentation using FSL FAST

- The `fslr` function fast calls fast from FSL (Zhang, Brady, and Smith 2001). The `--nobias` option tells FSL to not perform inhomogeneity correction (was already performed in `ANTsR`).

```
t1file = files["MPRAGE"]
t1fast = fast(t1,
              outfile = paste0(nii.stub(t1file), "_FAST"),
              opts = "--nobias")
```
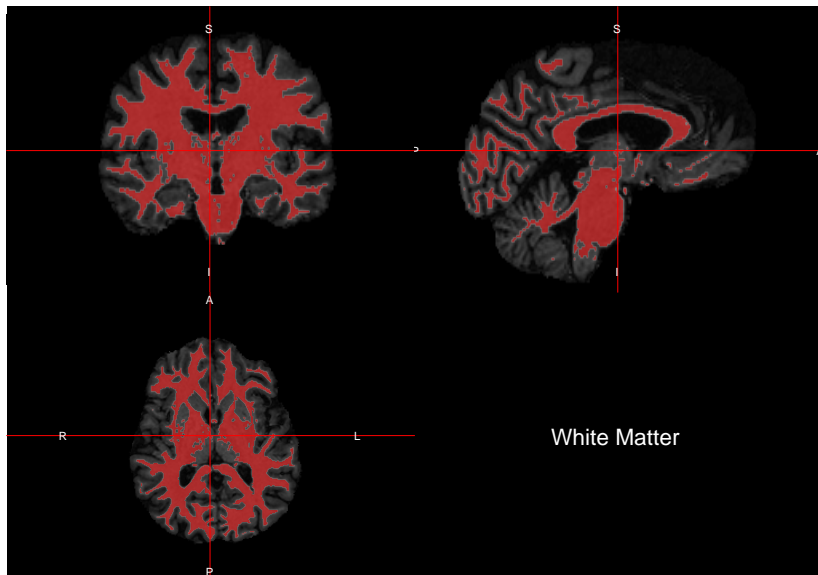
Results

FAST assumes three tissue classes and produces an image with the three labels, ordered by increasing within-class mean intensities. In a T1 image, this results in:

- ▶ Level 1: CSF
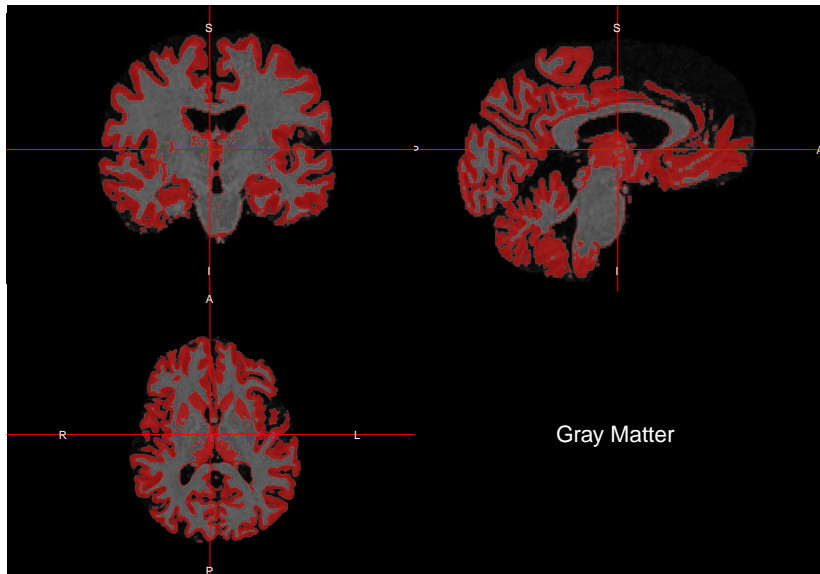- ▶ Level 2: Gray Matter
- ▶ Level 3: White Matter

White Matter

```
ortho2(t1, t1fast == 3, col.y = alpha("red", 0.5), text = '
```
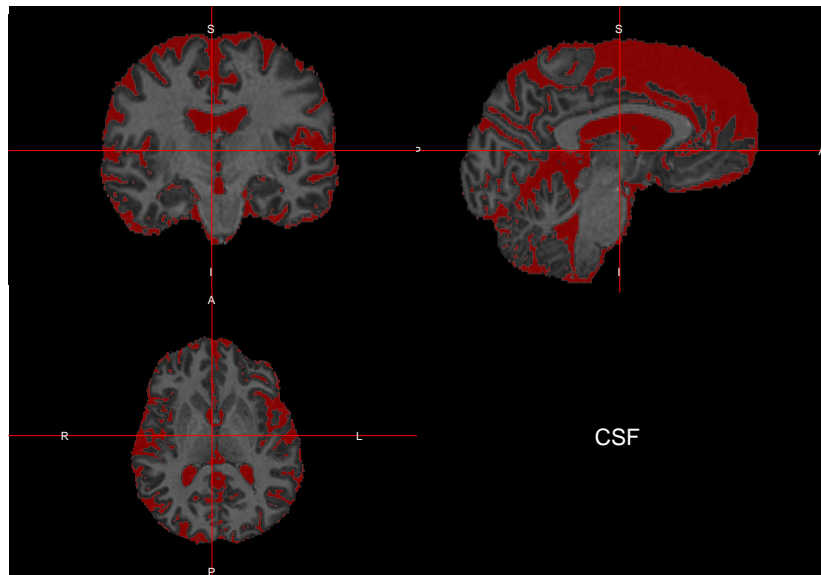

White Matter

Gray Matter

```
ortho2(t1, t1fast == 2, col.y = alpha("red", 0.5), text = '
```


Gray Matter

CSF

```
ortho2(t1, t1fast == 1, col.y = alpha("red", 0.5), text = '
```
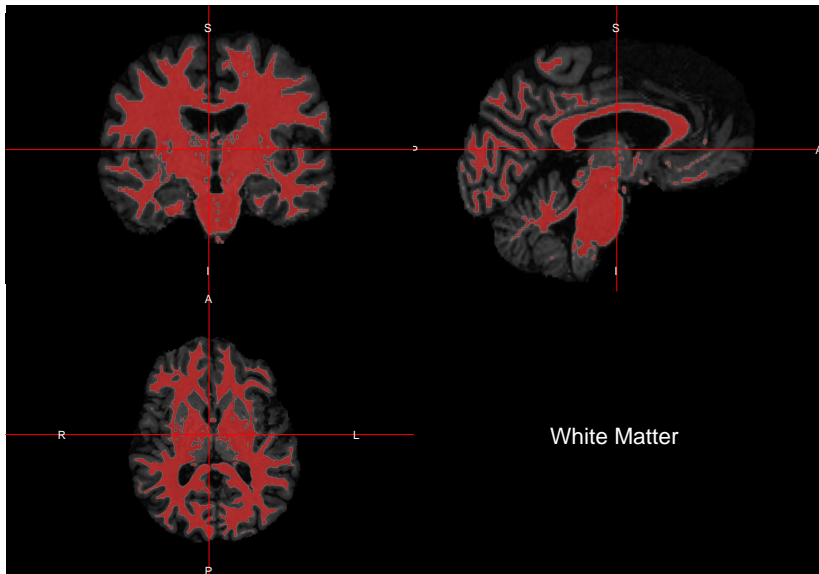

CSF

Windowing

```
rb = robust_window(t1)
robust_fast = fast(rb,
                outfile = paste0(nii.stub(t1file), "_FAST"),
                opts = "--nobias")
```
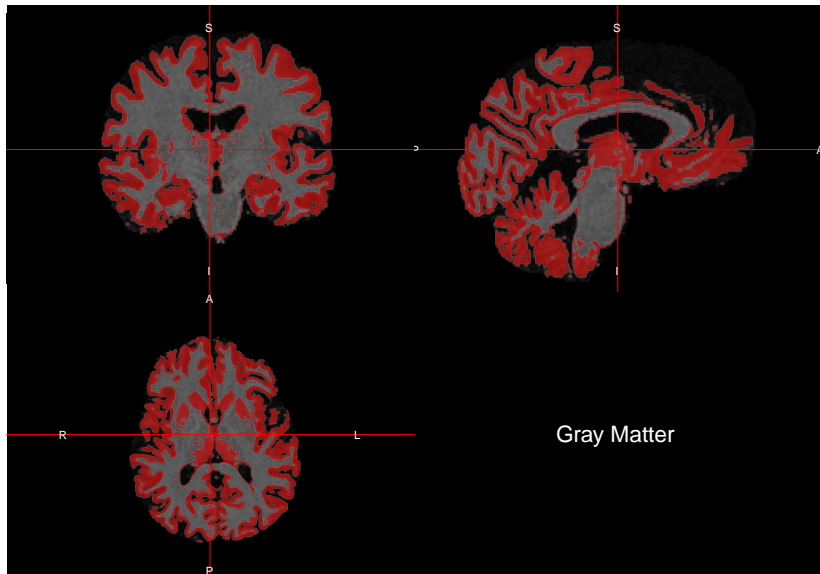
White Matter

```
ortho2(t1, robust_fast == 3, col.y = alpha("red", 0.5), tex
```
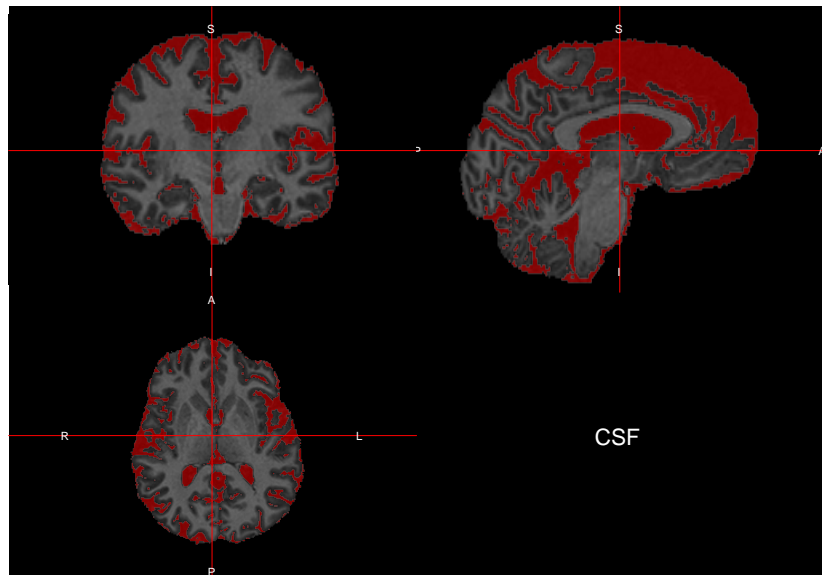

White Matter

Gray Matter

```
ortho2(t1, robust_fast == 2, col.y = alpha("red", 0.5), tex
```


Gray Matter

CSF

```
ortho2(t1, robust_fast == 1, col.y = alpha("red", 0.5), tex
```

FAST Results

- overall the results look good
- not much difference after dampening outliers using `robust_window`
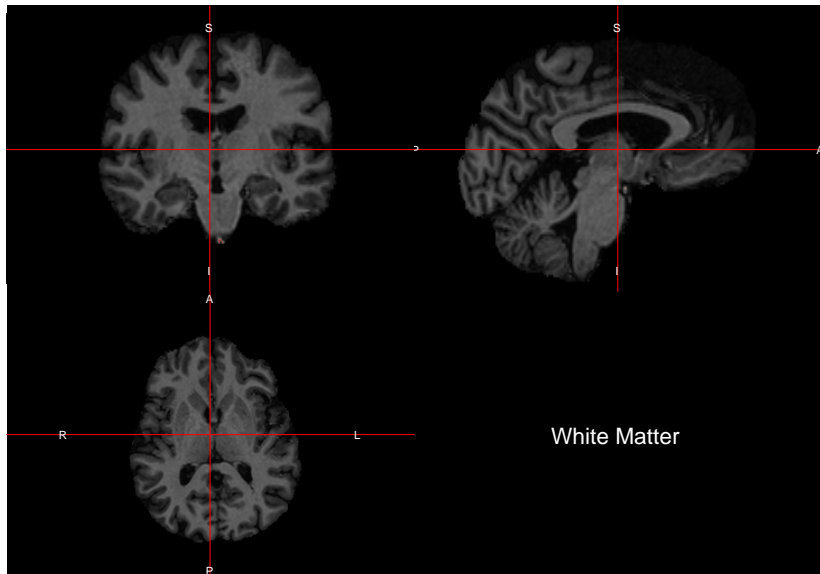- FAST is robust to noise

Tissue Segmentation using ANTsR, extrantsr

- Uses Atropos (Avants et al. 2011)

```
t1_otropos = otropos(a = t1, x = mask) # using original da
t1seg = t1_otropos$segmentation
```

White Matter

```
ortho2(t1, t1seg == 3, col.y = alpha("red", 0.5), text = "W
```
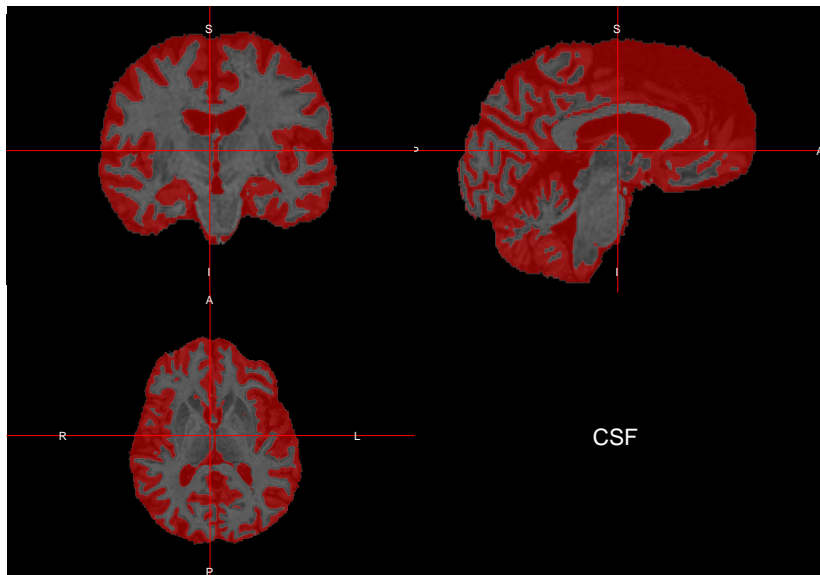
Gray Matter

```
ortho2(t1, t1seg == 2, col.y = alpha("red", 0.5), text = "(
```


Gray Matter

CSF

```
ortho2(t1, t1seg == 1, col.y = alpha("red", 0.5), text = "
```
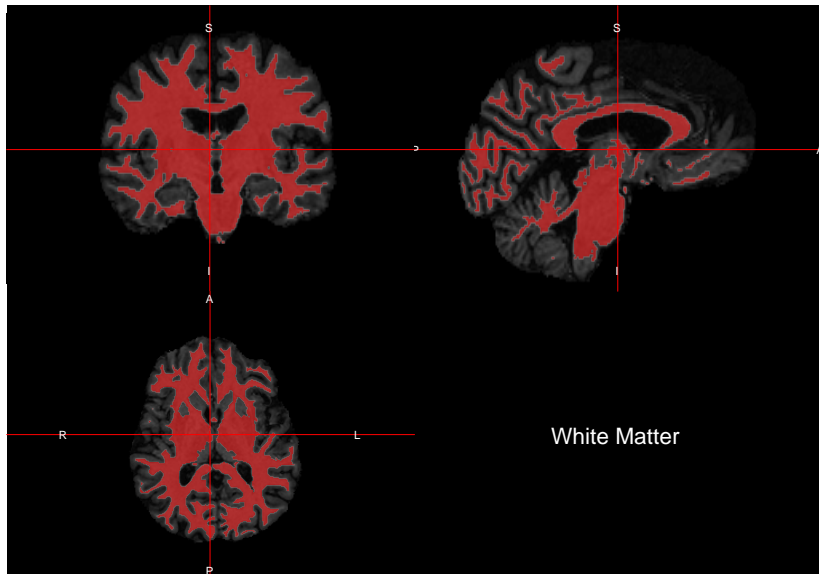

CSF

Tissue Segmentation using ANTsR, extrantsr

```
robust_t1_otropos = otropos(a = rb, x = mask) # using robus
robust_t1seg = robust_t1_otropos$segmentation
```
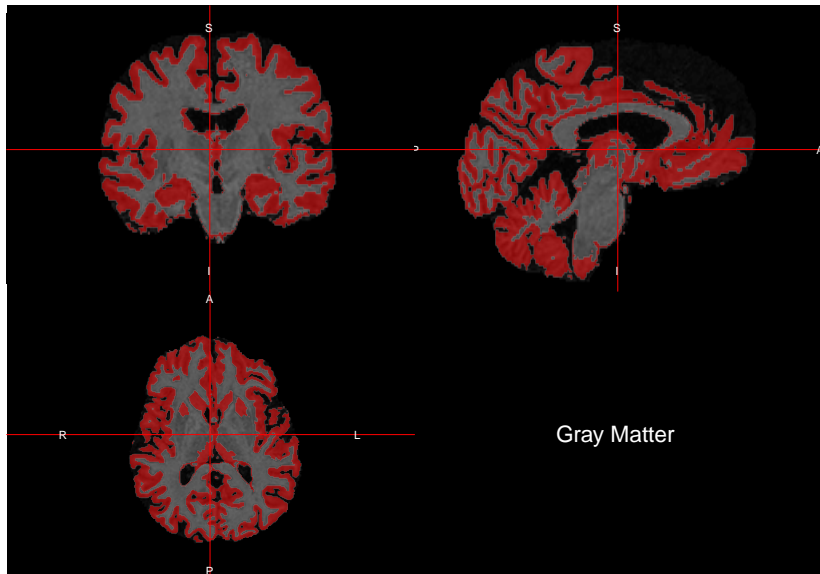
White Matter

```
ortho2(t1, robust_t1seg == 3, col.y = alpha("red", 0.5), te
```
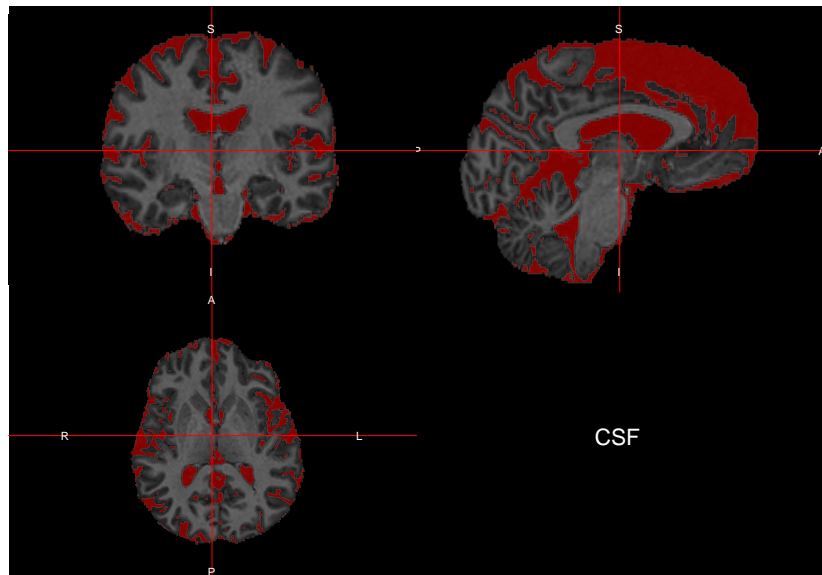

White Matter

Gray Matter

```
ortho2(t1, robust_t1seg == 2, col.y = alpha("red", 0.5), te
```


Gray Matter

CSF

```
ortho2(t1, robust_t1seg == 1, col.y = alpha("red", 0.5), te
```

References

Avants, Brian B, Nicholas J Tustison, Jue Wu, Philip A Cook, and James C Gee. 2011. "An Open Source Multivariate Framework for N-Tissue Segmentation with Evaluation on Public Data." *Neuroinformatics* 9 (4). Springer: 381–400.

Zhang, Yongyue, Michael Brady, and Stephen Smith. 2001. "Segmentation of Brain MR Images Through a Hidden Markov Random Field Model and the Expectation-Maximization Algorithm." *Medical Imaging, IEEE Transactions on* 20 (1): 45–57. `http://ieeexplore.ieee.org/xpls/abs_all.jsp?arnumber=906424`.