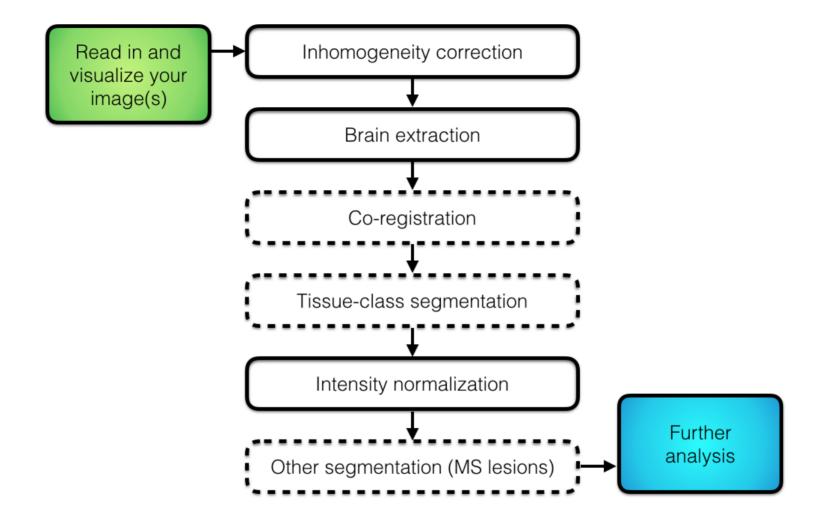
MS Lesion Segmentation

Processing math: 100%

Overall Pipeline

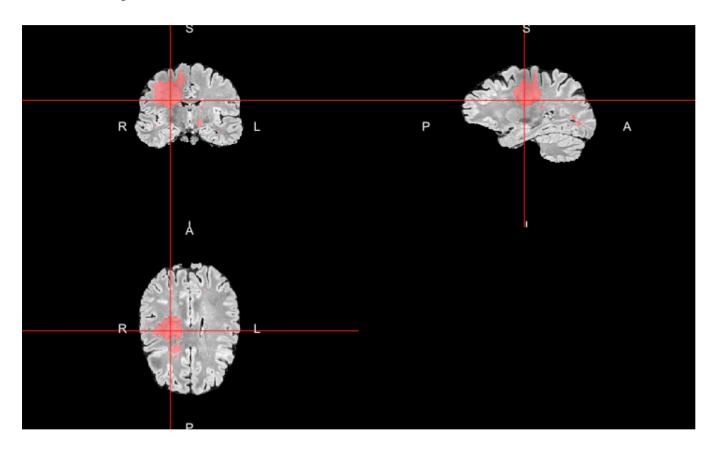


Background

- Obtaining manual lesion segmentations is often resource intensive.
- "Gold standard": Inter- and Intra-rater variability
- Accurate and efficient methods for automatic segmentation are necessary for scalability and research progress.
- · In this tutorial, we will learn how to train and apply OASIS (Sweeney et al. 2013), an automatic lesion segmentation model, to obtain predicted lesion probability maps.
 - relies on intensity-normalized data

Visualization

 Here's the FLAIR volume for training subject 05 with a manual lesion segmentation overlayed.



MS Lesion Segmentation with OASIS

- OASIS is Automated Statistical Inference for Segmentation (Sweeney et al. 2013).
- OASIS takes FLAIR, T1, T2, and PD (optional) images.
 - Produces OASIS probability maps of MS lesion presence.
 - These can be thresholded into a binary lesion segmentation.
- The OASIS model is based on a logistic regression.
 - Regress binary manual segmentation labels on the images, smoothed versions of the images, and some interaction terms (e.g., supervised learning).
 - Performed well compared to common machine learning models (Sweeney et al. 2014)

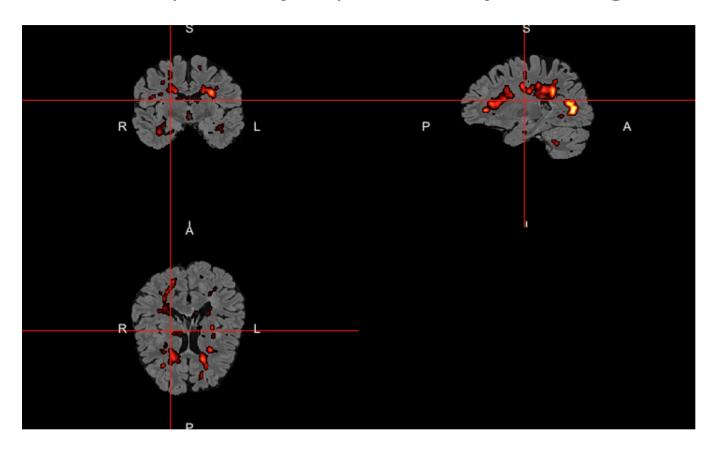
Default OASIS Model

- The OASIS library comes with default parameters that can be used to generate probability maps for new test subjects.
 - The default model was trained on approximately 100 MS subjects and 30 healthy subjects with manual segmentations.
- Here we apply oasis_predict with the default model to obtain OASIS probability maps for the test subjects.

```
library(oasis)
default_predict_ts = function(x) {
    res = oasis_predict(
        flair=ts_flairs[[x]], t1=ts_t1s[[x]],
        t2=ts_t2s[[x]], pd=ts_pds[[x]],
        brain_mask = ts_masks[[x]],
        preproc=FALSE, normalize=TRUE,
        model=oasis::oasis_model)
    return(res)
}
default_probs_ts = lapply(1:3, default_predict_ts)
```

Vizualization of probability map

• Here's the probability map for test subject 01 (no gold standard):

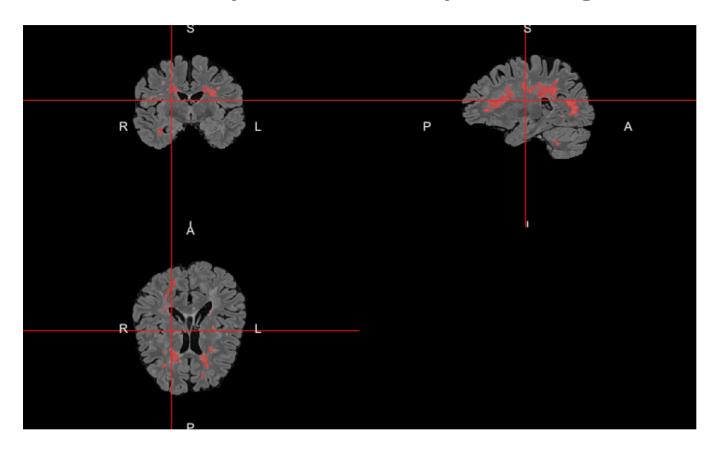


Thresholding: Getting a binary map

- We must choose a cutoff to binarize the OASIS probability maps.
- The binary argument in the oasis_predict function is FALSE by default, resulting in the output being the probability map.
 - Setting binary=TRUE will return the thresholded version, using the input to the threshold argument (default = 0.16).
 - 0.16 was obtained via a validation set allowing for a 0.5% false positive rate.
- In practice, we might want to use a grid search over thresholds and cross validation to choose the cutoff.

Vizualization of binary map

• Here's the binary mask for test subject 01, using the default 0.16 threshold:



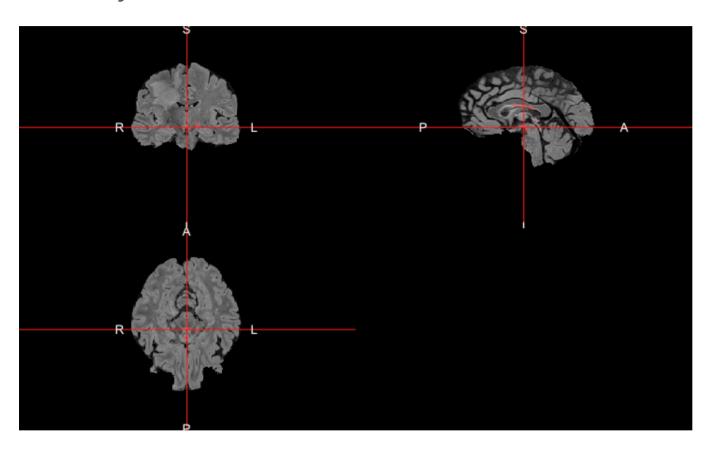
Default OASIS Model

- To evaluate how the default model performs, we need to compare the predictions to a gold standard.
- · Let's therefore obtain OASIS probability maps for our training subjects.
- · We will use the default threshold to binarize.

```
default_predict_tr = function(x) {
    res = oasis_predict(
        flair=tr_flairs[[x]], t1=tr_t1s[[x]],
        t2=tr_t2s[[x]], pd=tr_pds[[x]],
        brain_mask=tr_masks[[x]],
        preproc=FALSE, normalize=TRUE,
        model=oasis::oasis_model, binary=TRUE)
    return(res)
}
default_probs_tr = lapply(1:5, default_predict_tr)
```

Default OASIS Model Results

 Here's the FLAIR volume for training subject 05 with the OASIS segmentation overlayed.



Default OASIS Model Results

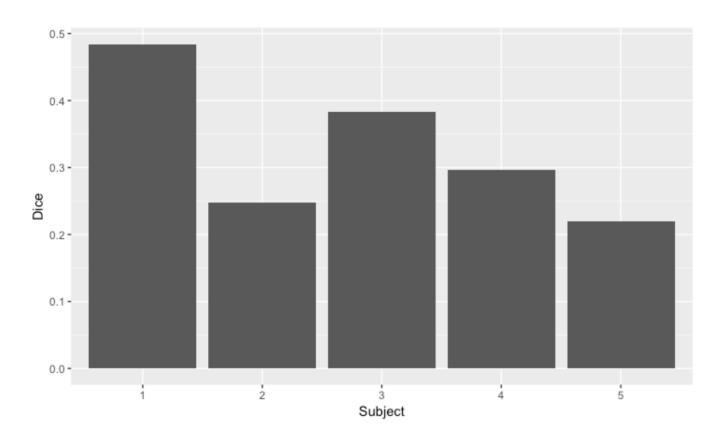
Sorensen-Dice coefficient

- · Similarity measure between two samples
- · Ranges from 0 to 1
- · (TP) true positive, (FP) false positive, (FN) false negative

$$D = 2TP2TP + FP + FN$$

Default OASIS Model Results

Dice coeffients for the training subjects compared to raters 1 and 2



Improving Results

- The default model is picking up a lot of false positives in the spinal cord.
- We might improve the results by re-training the OASIS model using our five training subjects.
- To re-train using new data, binary masks of gold standard lesion segmentations are needed and should be in T1 space.

Making OASIS data frames

- OASIS requires a particular data frame format, which we create using the function oasis_train_dataframe.
- Includes an option to preprocess your data (preproc), which does (1) inhomogeneity correction using fsl_biascorrect and (2) rigid coregistration using flirt to the T1 space.
- Includes an option to whole-brain intensity normalize (normalize).
- make df() below is a helper function.

```
make_df = function(x) {
    res = oasis_train_dataframe(
        flair=tr_flairs[[x]], t1=tr_t1s[[x]], t2=tr_t2s[[x]],
        pd=tr_pds[[x]], gold_standard=tr_golds[[x]],
        brain_mask=tr_masks[[x]],
        preproc=FALSE, normalize=TRUE, return_preproc=FALSE)
    return(res$oasis_dataframe)
}
oasis_dfs = lapply(1:5, make_df)
```

Training OASIS

- The function oasis_training takes the data frames we made and fits a logistic regression using labels and features from a subset of voxels in each subject's brain mask (top 15% in FLAIR intensity).
- The function do.call is a useful R function that applies the function named in the first argument to all elements of the list specified in the second argument.

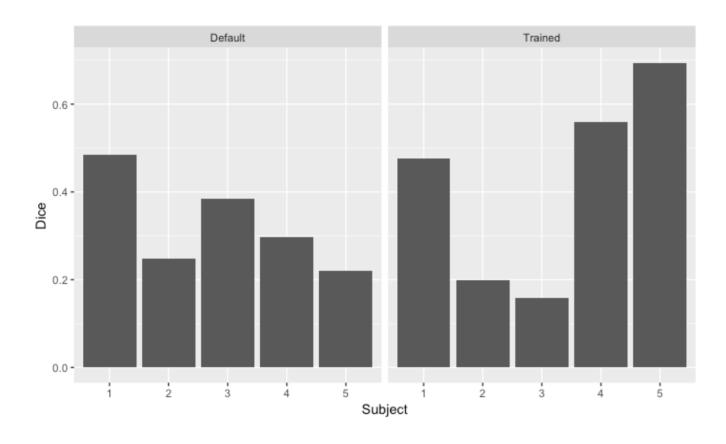
```
ms_model = do.call("oasis_training", oasis_dfs)
```

OASIS model object

```
print(ms.lesion::ms model)
Call: glm(formula = form, family = binomial, data = df)
Coefficients:
   (Intercept)
                     FLAIR 10
                                        FLAIR FLAIR 20
                                     1.14120
                      13.10\overline{3}86
                                                     -18.77\overline{0}10
      -4.79369
                                         T2 20
         T2 10
                                                         T1 10
                                      -7.06\overline{7}50 13.63\overline{5}54
       4.85\overline{3}70
                    1.09444
                         T1 20 FLAIR 10:FLAIR FLAIR:FLAIR 20
            T1
       1.04771 -21.09\overline{8}48
                                     -1.28891 1.03\overline{1}21
      T2 10:T2
                  T2:T2 20
                                      T1 10:T1 T1:T1 20
       0.09151
                    3.18\overline{9}03
                                      -1.04701 3.14\overline{2}65
Degrees of Freedom: 3930444 Total (i.e. Null); 3930429 Residual
Null Deviance: 2691000
Residual Deviance: 1842000 AIC: 1842000
```

Trained OASIS Model Results

- Using the same threshold of 0.16.
- · Dice coeffients for default vs. re-trained OASIS model



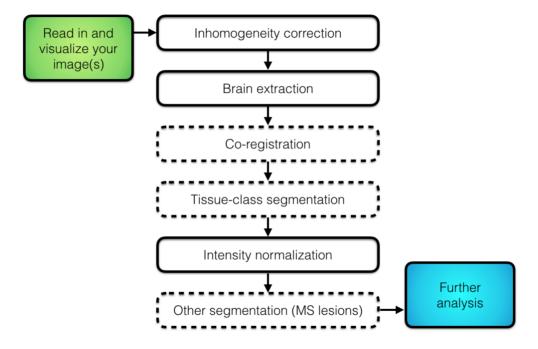
Improvement

· Percent improvement in Dice over the default model:

ID	Dice
01	-1.8
02	-19.8
03	-58.6
04	88.1
05	215.2

Wrap-up

- We've covered all (or most) image pre-procssing steps in a typical image preprocessing pipeline, starting with raw nifti images.
- · Everything was done in R!



What we didn't cover

- fMRI: see fmri library
- · Other imaging modalities, e.g., CT, PET
 - MALF segmentation is robust
- Voxel-wise testing: see voxel library
- · Other population-level statistical inference:
- Statistical/machine learning: see caret library

What can you do next?

- Further modeling and statistical analysis.
- Register images to a template to do population inference.
- General R
 - Build your own R libraries for image analysis.
 - Rmarkdown for reproducible reports
 - R shiny apps

Resources

- Neurohacking tutorial on Coursera
- · Neuroconductor: central repository for image analysis R libraries

Website

http://johnmuschelli.com/imaging_in_r

References

Sweeney, Elizabeth M, Russell T Shinohara, Navid Shiee, Farrah J Mateen, Avni A Chudgar, Jennifer L Cuzzocreo, Peter A Calabresi, Dzung L Pham, Daniel S Reich, and Ciprian M Crainiceanu. 2013. "OASIS Is Automated Statistical Inference for Segmentation, with Applications to Multiple Sclerosis Lesion Segmentation in Mri." 2. Elsevier:402–13.

Sweeney, Elizabeth M, Joshua T Vogelstein, Jennifer L Cuzzocreo, Peter A Calabresi, Daniel S Reich, Ciprian M Crainiceanu, and Russell T Shinohara. 2014. "A Comparison of Supervised Machine Learning Algorithms and Feature Vectors for MS Lesion Segmentation Using Multimodal Structural MRI." 9 (4). Public Library of Science:e95753.