

# MELD Protocol 3 - FreeSurfer Quality Control

Sophie Adler, Kirstie Whitaker, Konrad Wagstyl

## Abstract

*The MELD Project is an international collaboration aiming to create open-access, robust and generalisable tools for FCD detection. To this end, we will train a neural network classifier on MRI features from FCD patients from multiple centres worldwide.*

**Protocol 3 provides instructions on how to quality control the FreeSurfer reconstructions.**

These instructions are based on the freely available protocols on the ENIGMA-epilepsy website <http://enigma.ini.usc.edu>

We are very grateful to Derrek Hibar, Neda Jahanshad, Roberto Toro, Jerod Rasmussen, Theo van Erp, Esther Walton and Stefan Ehrlich who wrote the original ENIGMA protocols and offered them with an unlimited license without warranty!

The main change is correct co-registration of FLAIR to T1 is now checked.

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## Guidelines

These instructions are based on the freely available protocols on the ENIGMA-epilepsy website <http://enigma.ini.usc.edu/ongoing/enigma-epilepsy/enigma-epilepsy-protocols/>

We are very grateful to Derrek Hibar, Neda Jahanshad, Roberto Toro, Jerod Rasmussen, Theo van Erp, Esther Walton and Stefan Ehrlich who wrote the original ENIGMA protocol.

## Before start

Ensure that you have logged onto the MELD github (<https://github.com/MELDProject/meld>) and cloned the repository.

This will contain the essential MELD\_QC folder.

You also need to have MATLAB installed.

**If you have any questions or run into problems, please feel free to contact the MELD project: ([meld.study@gmail.com](mailto:meld.study@gmail.com))**

## Protocol

### Preparing for QC

#### Step 1.

#### Extract and Organize Cortical Measures (FreeSurfer)

Open the script for extracting SurfaceArea and Thickness Values: **extract.sh** located in the MELD\_QC folder.

We want to get the Surface Area and Thickness of each ROI for each subject from FreeSurfer.

The extract.sh script will extract and organize each of the values for each FreeSurfer ROI. The script assumes that your FreeSurfer output are organized as specified in protocol's 1 and 2.

### Preparing for QC

#### Step 2.

## Extract and Organize Cortical Measures (FreeSurfer)

Save the **extract.sh** script in the Parent Folder with your FreeSurfer output (`<path>/meld/output/`).

### Preparing for QC

#### Step 3.

Open the **extract.sh** and ensure that the for loop (line 6) selects the subject folder naming scheme used in this study (i.e. MELD):

```
for subj_id in $(ls -d MELD*); do  
#may need to change this so that it selects subjects with FS  
output
```

---

*Save the extract.sh script after editing.*

### Preparing for QC

#### Step 4.

On the command line, you can run the script directly by running:

```
sh <path>/meld/output/extract.sh
```

The result of this step will be two comma-separated (CSV) files that can be opened in your favorite spreadsheet application (i.e. Excel). The first row is a header describing the extracted regions and names for each column. Each row after the first gives the cortical thickness (or surface area) measures for each subject found in your FreeSurfer directory.

### Preparing for QC

#### Step 5.

**Note 1:** After running the extract.sh script, open both of the CSV files (CorticalMeasuresENIGMA\_ThickAvg.csv and CorticalMeasuresENIGMA\_SurfAvg.csv) and make sure that only subjects are listed in the rows of the file. Sometimes if there are other folders in your parent directory those folders can sometimes become included in your final files, if that happens just delete those from your CSV files and save.

## Preparing for QC

### Step 6.

**Note 2:** When you edit the files in Excel, be sure to keep them in CSV format when you save!

## Preparing for QC - Outlier Detection

### Step 7.

*This is a simple R script that will identify subjects with cortical thickness and surface area values that deviate from the rest of your subjects.*

*This step requires that you have R installed and that you have the outlier detection script found called outliers.R which is located in the MELD\_QC folder.*

Change directories to the location of your **CorticalMeasuresENIGMA\_ThickAvg.csv** and **CorticalMeasuresENIGMA\_SurfAvg.csv** generated in Step 1 of the protocols.

```
cd <path>/meld/output/
```

---

## Preparing for QC - Outlier Detection

### Step 8.

Save the outliers.R script that directory (<path>/meld/output)

## Preparing for QC - Outlier Detection

### Step 9.

Run:

```
R --no-save --slave < outliers.R > outliers.log
```

---

This will generate a log file that will tell you which subjects are outliers and for which structures they are outliers for.

## Step 10.

**There are three major steps for quality checking the cortical surface segmentations outputted from FreeSurfer:**

1. **Internal Surface Method:** This method uses a Matlab function to plot cortical surface segmentations directly on a subject's scan and collates snapshots from internal slices of the brain into a webpage for easy checking.
2. **External Surface Method:** This is loosely based on the QATools for FreeSurfer. This creates a webpage with external views of the segmentations from different angles.
3. **Check FLAIR co-registration (if applicable)**

**The following steps will take you through these 3 methods**

## Step 11.

Start Matlab: `/usr/local/matlab/bin/matlab`

## Step 12.

Folder containing scripts required is: `<path>/meld/MELD_QC/ENIGMA_QC`

Add the folder containing all of the required scripts to Matlab's path.

Select: File -> Set Path -> Add Folder -> {OK} -> {Save} -> {Close}

## Step 13.

In the Matlab console window change directories to the folder with all of your FreeSurfer subject folders.

```
cd /meld/output/
```

## Step 14.

Make a directory to store all of the QC output.

## Quality Checking - The Internal Surface Method

### Step 15.

The script we want to run is called **func\_make\_corticalpngs\_ENIGMA\_QC.m** with the following parameters:

```
func_make_corticalpngs_ENIGMA_QC(output_QC_directory, subject_name, select_MRI_image,
select_Segmented_image)
```

We want to set 'subject\_name' such that 'subject\_name' + 'select\_Segmented\_image' will form the full name of the segmentation label files (e.g. subj1/mri/aparc+aseg.mgz) and similarly for 'select\_MRI\_image' we want it to give the full name of the registered MRI scan outputted by FreeSurfer (e.g. subj1/mri/orig.mgz).

In the Matlab command window we can do:

```
QC_output_directory='<path>/meld/output/QC/';
FS_directory='<path>/meld/output/';
a=dir(char(strcat(FS_directory,'/MELD_*')));%Choose this so that it
selects %only your subject folders that contain FS output

for x = 1:size(a,1)
    [c,b,d]=fileparts(a(x,1).name); %b becomes the subject_name
    try
        func_make_corticalpngs_ENIGMA_QC(QC_output_directory, b,
[FS_directory, '/', b, '/mri/orig.mgz'], [FS_directory, '/', b,
'/mri/aparc+aseg.mgz']);
    end
    display(['Done with subject: ', b, ': ', num2str(x-2), ' of ',
num2str(size(a,1)-2)]);
end
```

---

The func\_make\_corticalpngs\_ENIGMA\_QC script should take approximately 7 seconds/subject and will output a series of \*.png image files separated by individual subject folders.

**NB:** if you run into problems with this Matlab loop try removing the last "/" in the QC\_output\_directory variable. So, QC\_output\_directory='/meld/output/QC/'; would become QC\_output\_directory='/meld/output/QC';

## Quality Checking - The Internal Surface Method

### Step 16.

#### Making the QC Webpage:

To create a webpage for easy viewing of the QC output you just generated in Matlab. Go to the directory where you stored the script **make\_ENIGMA\_QC\_webpage.sh** and make sure it is executable:

```
chmod 777 make_ENIGMA_QC_webpage.sh
```

---

## Quality Checking - The Internal Surface Method

### Step 17.

Now to run the script, just give the script the full path to the directory where you stored the Matlab QC output files:

```
./make_ENIGMA_QC_webpage.sh /meld/output/QC/
```

---

**NB:** If you have trouble running this script, it's possible that you need to fix the line endings in the script before running. You can do this by running this command: `sed -i -e 's/\r$//' make_ENIGMA_QC_webpage.sh`

This script will create a webpage called **ENIGMA\_Cortical\_QC.html** in the same folder as your QC output.

**Step 18.**

To open the webpage in a browser in a Linux environment you can probably just type:

```
firefox /meld/output/QC/ENIGMA_Cortical_QC.html
```

---

In a mac environment you can type

```
open /meld/output/QC/ENIGMA_Cortical_QC.html
```

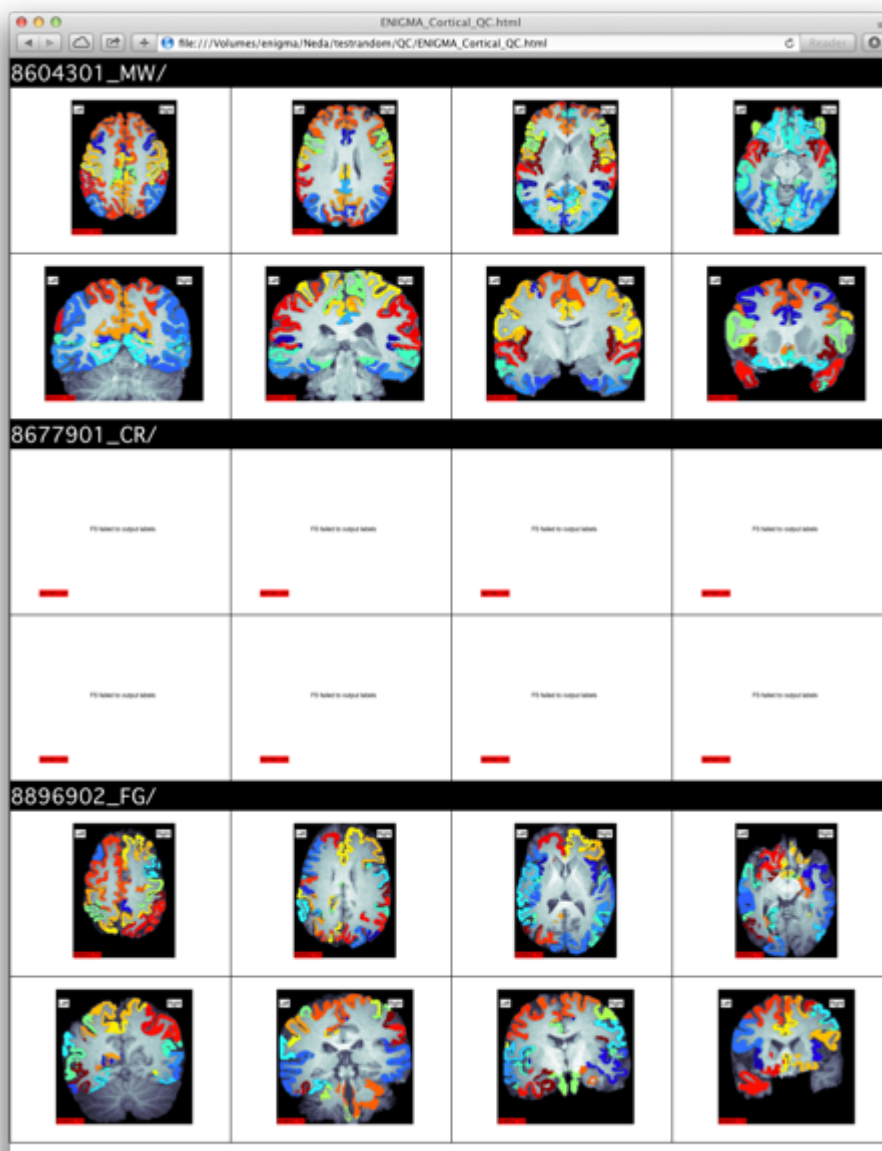
---

**Note:** if you want to check the segmentation on another computer, you can just copy over the whole /meld/output/QC/ output folder to your computer and open the webpage from there.

Scroll through each set of images. Note that you can click on a subject's files to see a larger version.

**NOTE: you can use the legend.jpg file found in the ENIGMA\_QC/ folder as a colored coded reference of each FreeSurfer ROI (split by left/right).**

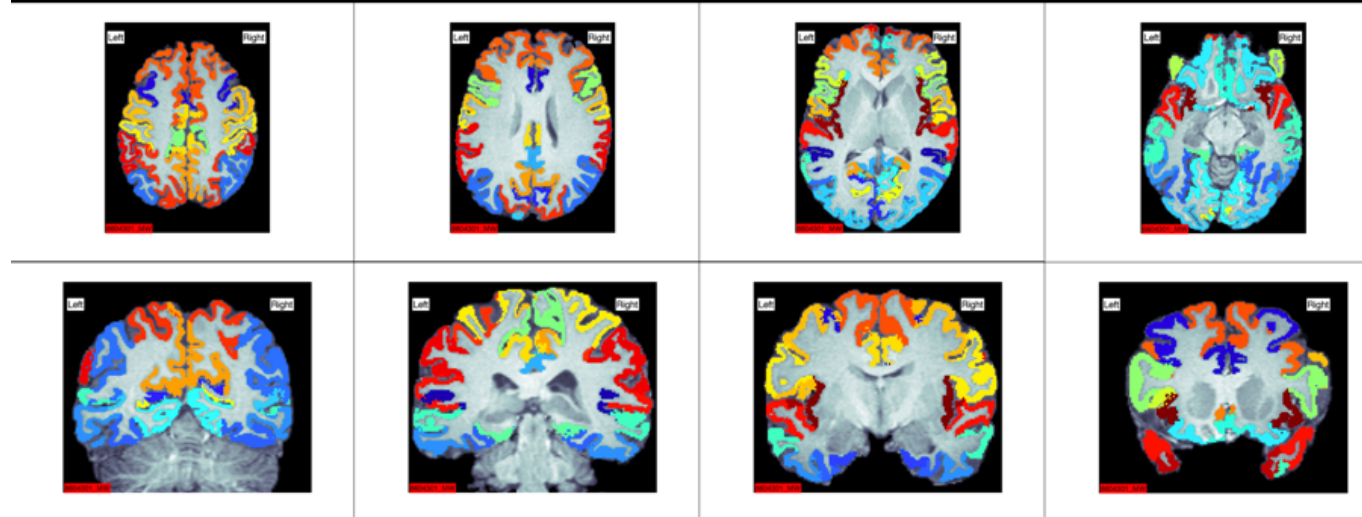




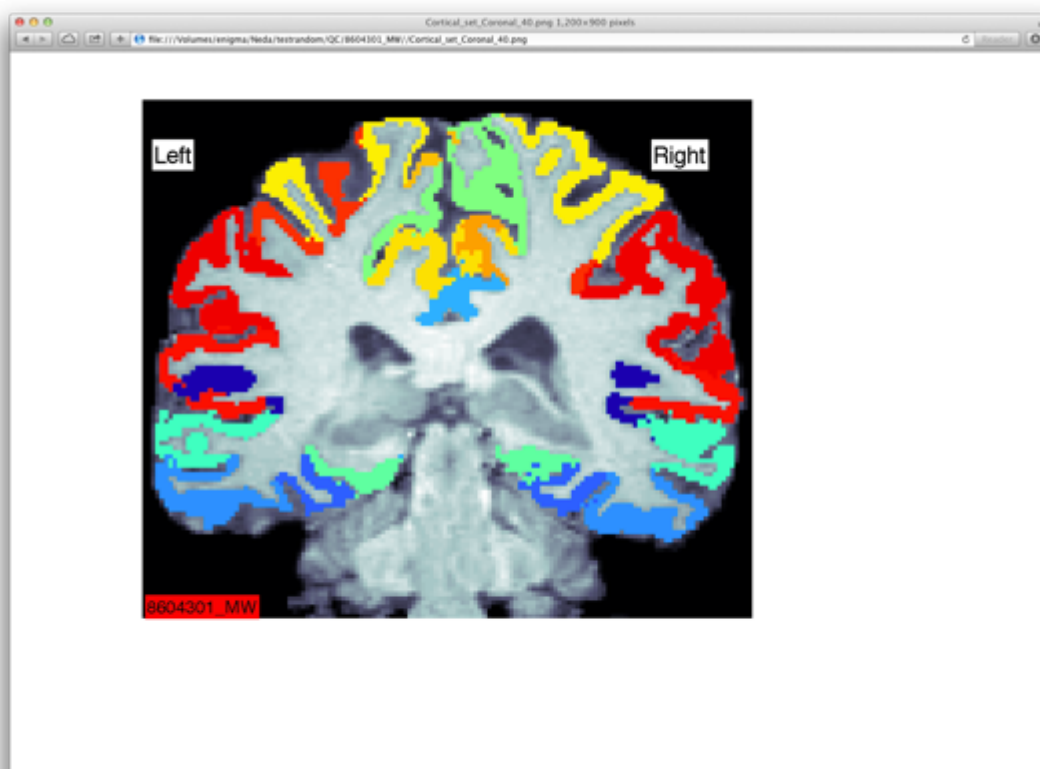
## Quality Checking - The Internal Surface Method

### Step 19.

Here is an example of a good segmentation:



Here is a close up of a good segmentation:



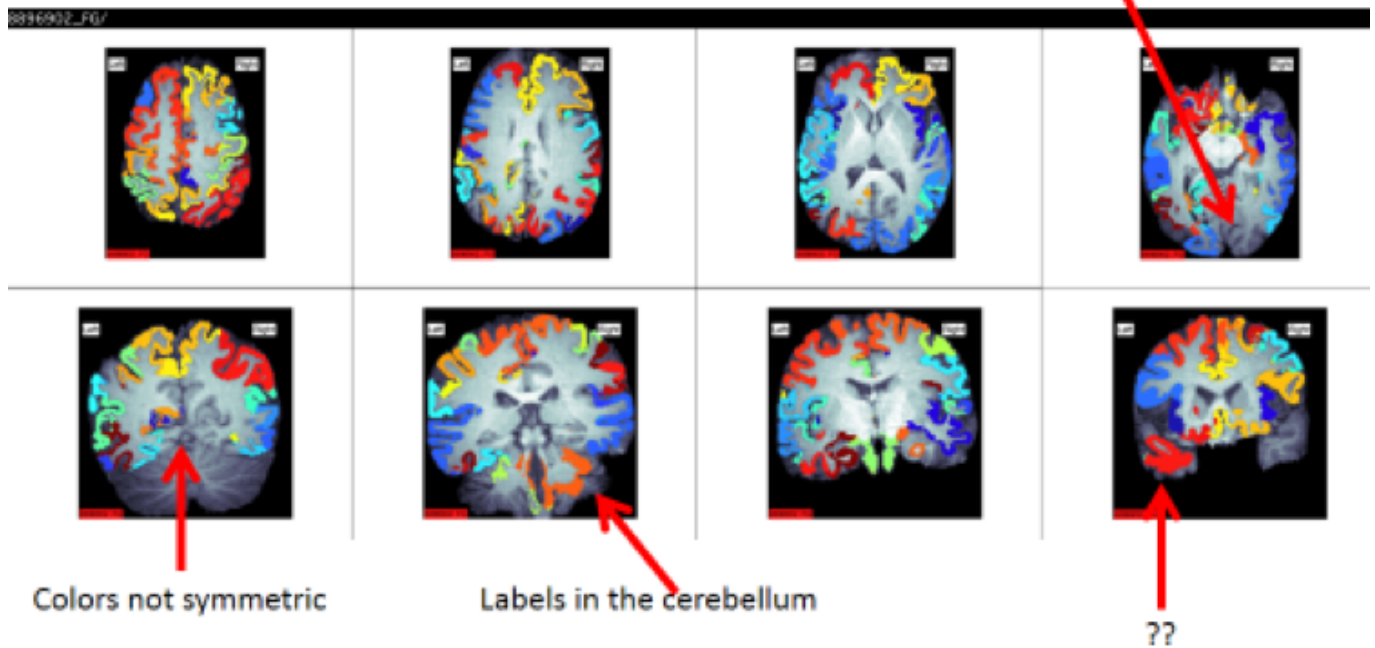
## Quality Checking - The Internal Surface Method

### Step 20.

Here is an example of a BAD segmentation:

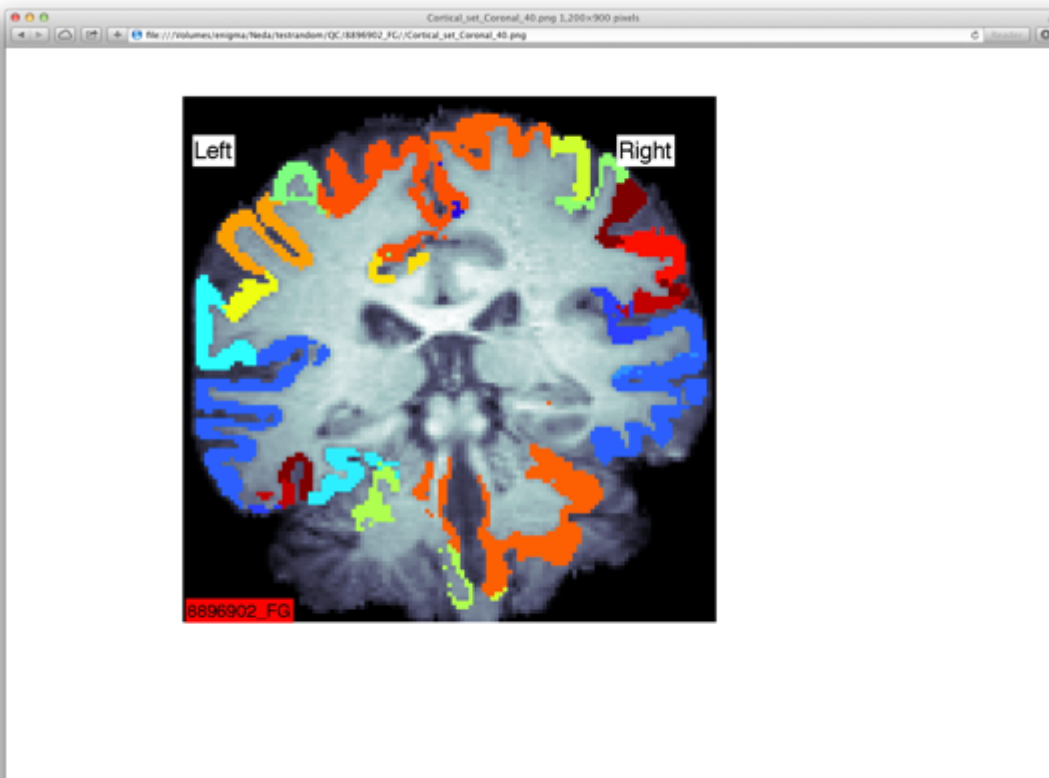
# Bad

Cortical regions missing



**rerun or remove entire subject**

Here is a close-up of a poor segmentation:



## Quality Checking - The Internal Surface Method

### Step 21.

Open the data file MELD\_[site code]\_participants.csv

### Rate the quality of the segmentation:

If good, code as 1 in the QC column

If borderline, code as 2 in the QC column

If poor, either attempt to rerun the FreeSurfer segmentation (protocol 2) OR code as 3 in the QC column.

**Make sure to save the file back in CSV format!**

## Quality Checking - The External Surface Method

### Step 22.

**Use the script fsqc.sh:**

**NB: FreeSurfer and its utilities need to be in your path or this script will not run properly.**

**You can type `tksurfer` on the command line to make sure it is available.**

Before you want to work with FreeSurfer, you must make sure three things have happened:

1. The variable `FREESURFER_HOME` is set (so your computer knows where FreeSurfer is installed):

```
setenv FREESURFER_HOME <freesurfer_installation_directory>/freesurfer
```

2. The FreeSurfer set up script must be sourced (so FreeSurfer knows the location of everything it needs):

```
source $FREESURFER_HOME/SetUpFreeSurfer.csh
```

3. FreeSurfer has been pointed to a directory of subjects to work on:

```
setenv SUBJECTS_DIR <path>/meld/output
```

## Quality Checking - The External Surface Method

### Step 23.

The script **fsqc.sh** will create a webpage with lateral and medial snapshots of pial surface reconstructions colored with cortical labels. Clicking on the images will display a larger version.

To run the script, first source FreeSurfer's environment variable `$SUBJECTS_DIR` to point to your subjects directory. For example:

```
bash
export SUBJECTS_DIR=/meld/output
```

## Quality Checking - The External Surface Method

### Step 24.

Next create a directory to contain the snapshots (.tif image files), here we will call it `fsqcdir/`. Change

the working directory to fsqcdir and run the **fsqc.sh** script from there:

```
mkdir /meld/path/output/QC/fsqcdir
cd fsqcdir
source /meld/MELD_QC/fsqc.sh
```

---

This script will call `tksurfer` in a loop for each subject and output a series of 4 images (.tif) files for each subject. It will also create a website called index.html so that you can easily view the images. You can open the index.html file in any browser, just make sure all of the .tif files are in the same folder if you decide to move the index.html file to a different location (like a local computer).

## Quality Checking - The External Surface Method

### Step 25.

If you are in a Linux environment you should be able to just type on the command line:

#while in the output fsqcdir output folder

```
firefox index.html
```

---

In a mac environment you can type

```
open /meld/output/QC/fsqcdir/index.html
```

---

Please note that .tiff is not supported by all browsers (e.g. Google Chrome). It is supported by Safari.

## ■ ANNOTATIONS

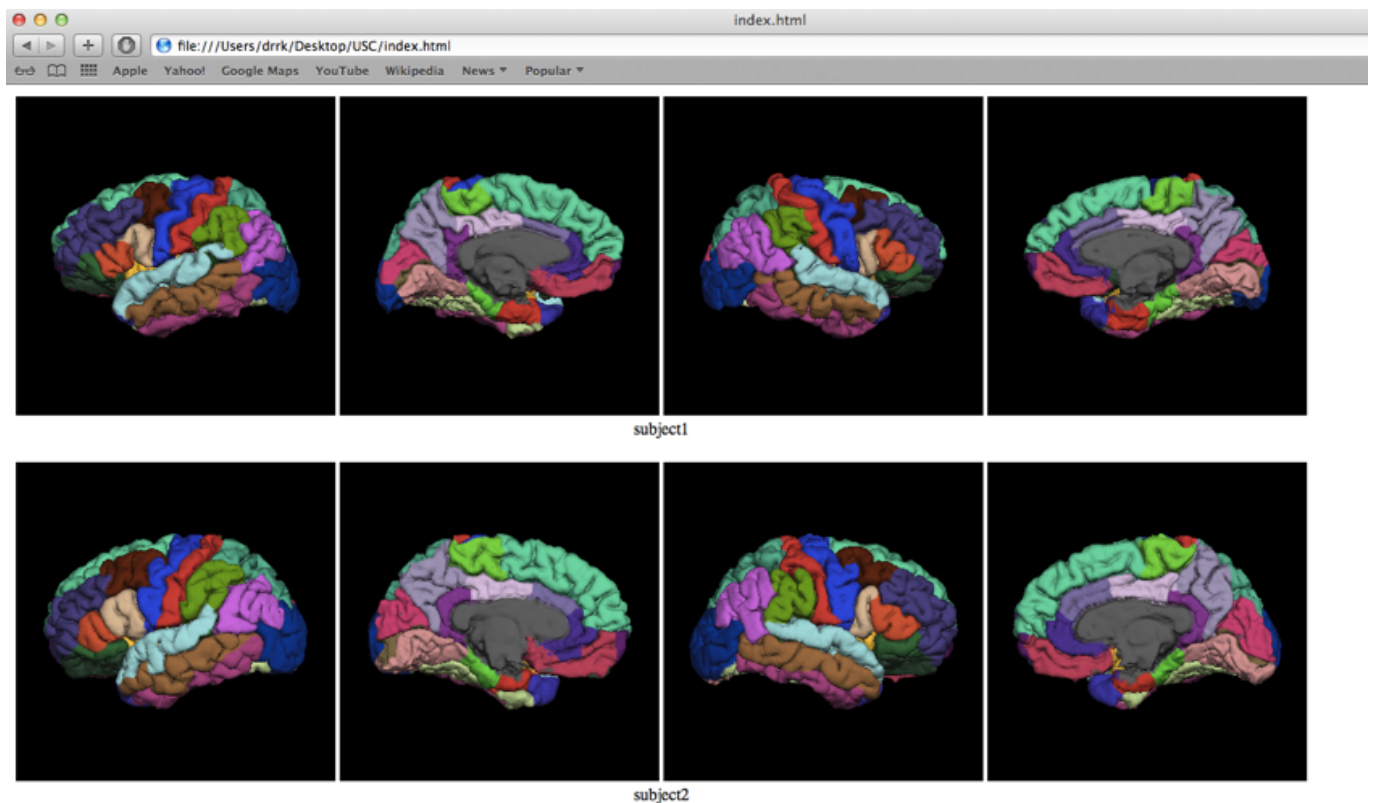
**Mira Semmelroch** 20 Mar 2018

This does not seem to work for me but it is easy to view the results otherwise

## Quality Checking - The External Surface Method

### Step 26.

These are examples of good external surfaces:



### Make sure to:

- Check that all lobes are present, especially the ventral part of the temporal lobe
- Check that labels positions are not different from the general case

### ■ ANNOTATIONS

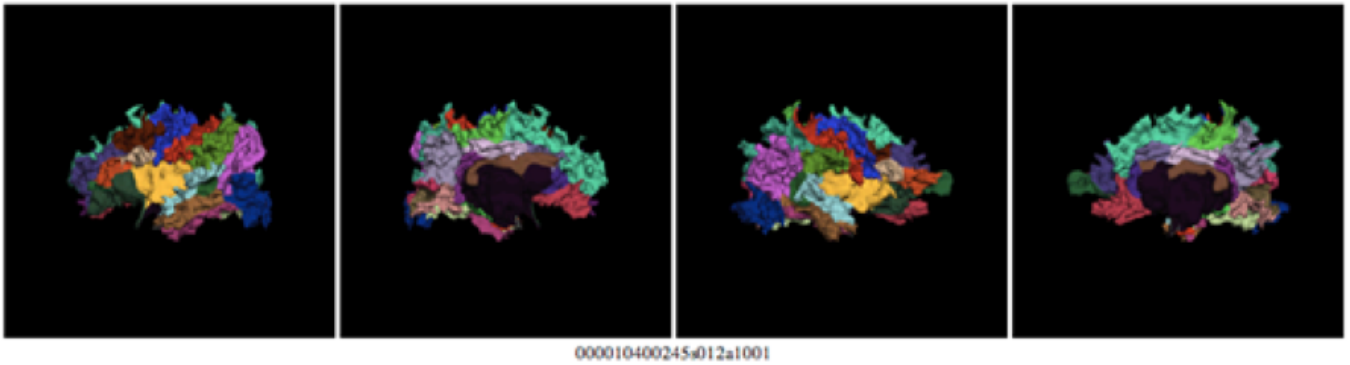
**Mira Semmelroch** 20 Mar 2018

Maybe add further examples (like ENIGMA) eg regarding banks of superior temporal sulcus, ventricles, insula etc (probably only necessary if people have joint this project whopreviously have not been involved with the ENIGMA Freesurfer analysis)

## Quality Checking - The External Surface Method

### Step 27.

This is an example of a very poor external surface:



## Quality Checking - The External Surface Method

### Step 28.

Open the data file MELD\_[site code]\_participants.csv

**Check that the quality of the external surface is the same as the internal surface. If the quality is poorer, re-code:**

If good, code as 1 in the QC column

If borderline, code as 2 in the QC column

If poor, either attempt to rerun the FreeSurfer segmentation (protocol 2) OR code as 3 in the QC column.

**Make sure to save the file back in CSV format!**

### ■ ANNOTATIONS

**Mira Semmelroch** 20 Mar 2018

I assume you would want those regions not well segmented removed from the CorticaMeasuresENIGMA\_SurfAvg.csv spreadsheet in this step???

## Quality Checking - FLAIR coregistration

### Step 29.

**NB: FreeSurfer and its utilities need to be in your path or this script will not run properly. You can type tkmedit on the command line to make sure it is available.**

The script **FLAIR\_coreg.sh** will create a webpage with coronal, sagittal and axial snapshots of T1 and FLAIR images with the pial and grey-white matter boundary surfaces superimposed. Clicking on the images will display a larger version.



To run the script, first source FreeSurfer's environment variable \$SUBJECTS\_DIR to point to your subjects directory. For example:

```
bash
export SUBJECTS_DIR=/meld/output
```

---

## Quality Checking - FLAIR coregistration

### Step 30.

Next create a directory to contain the snapshots (.tif image files), here we will call it FLAIR\_coreg/. Change the working directory to FLAIR\_coreg and run the **FLAIR\_coreg.sh** script from there:

```
mkdir meld/output/QC/FLAIR_coreg
cd FLAIR_coreg
source /meld/MELD_QC/FLAIR_coreg.sh
```

---

This script will call `tkmedit` in a loop for each subject and output a series of 4 images (.tif) files for each subject. It will also create a website called index.html so that you can easily view the images. You can open the index.html file in any browser, just make sure all of the .tif files are in the same folder if you decide to move the index.html file to a different location (like a local computer).

## Quality Checking - FLAIR coregistration

### Step 31.

If you are in a Linux environment you should be able to just type on the command line:

#while in the output fsqcdir output folder

```
firefox index.html
```

---

In a mac environment you can type

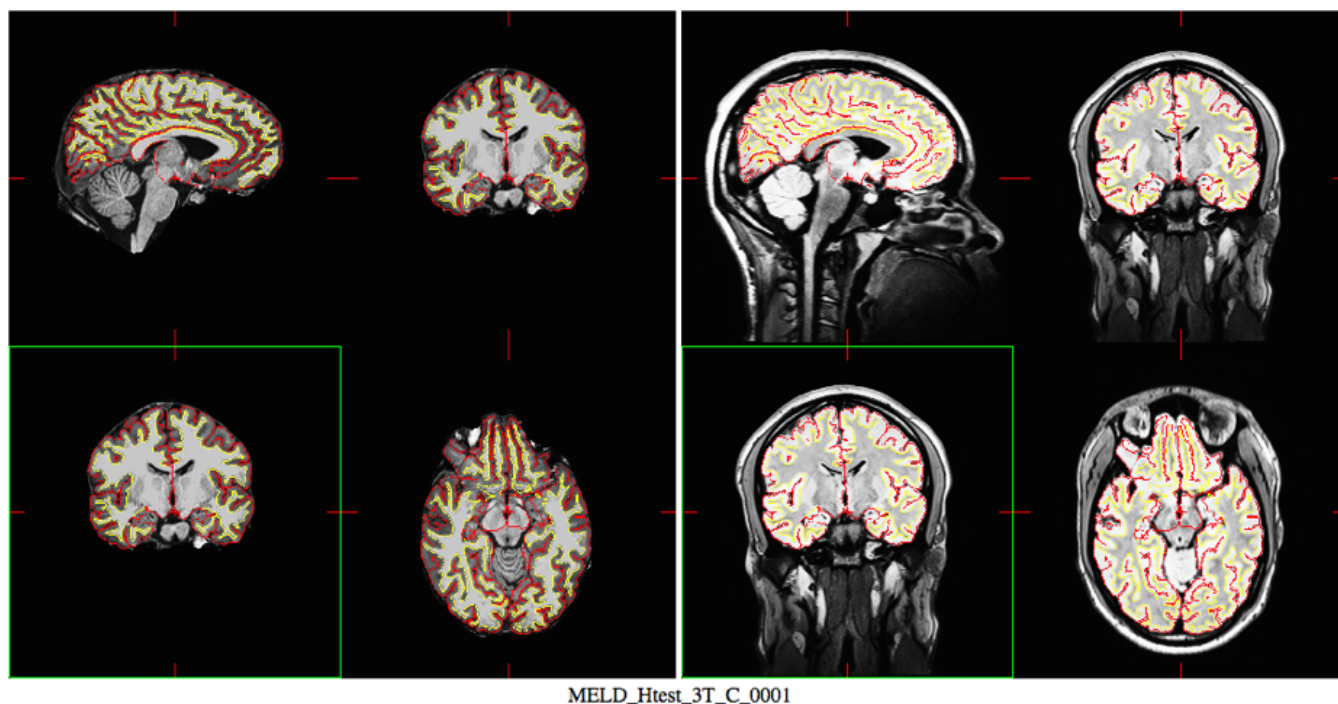
```
open /meld/output/QC/FLAIR_coreg/index.html
```

---

Please note that .tiff is not supported by all browsers. It is supported by Safari.

### Step 32.

This is an example of a correctly co-registered FLAIR scan:



If the FLAIR is not correctly co-registered you will need to re-run the FreeSurfer reconstruction (Protocol 2) but change some parameters to ensure that the FLAIR is correctly co-registered this time. **Please see supplementary protocol on FLAIR coregistration issues.**

#### ■ ANNOTATIONS

**Mira Semmelroch** 22 Mar 2018

You are referring to a supplementary protocol on Flair coregistration issues. Where do I find this? (sorry if I missed something obvious here)

### Step 33.

## Warnings

PLEASE DO NOT SHARE ANY IDENTIFIABLE DATA

Data sharing only occurs at the level of anonymised demographics information and anonymised data matrices. These are in a template space that cannot be traced back to an individual.