Sharif University Of Technology

Neuroscience Lab

Introduction to fMRI (AFNI)

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Section 1: Brain Extraction

1.1 Removing the Skull and Surrounding Area of the Brain

By using the function 1, the skull is precisely separated from surrounding. The result show in figure 1.

$$3dSkullStrip - inputsub - 13_T1w.nii.gz - prefix FT_anat_brain$$
 (1)

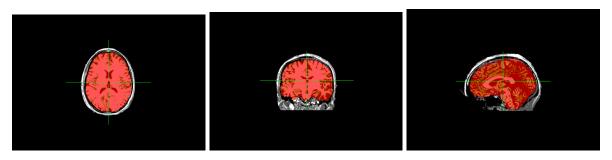


Figure 1: Skull- axial, coronal and sagittal view.

1.2 Making a Gray Matter Mask

By using the function 2 and the range is shown in figure 2, the gray matter mask is created.

 $3dcalc - aFT_anat_brain + orig - expr'within(a, 600, 2900)' - prefix anat_3dcalc_result$ (2)

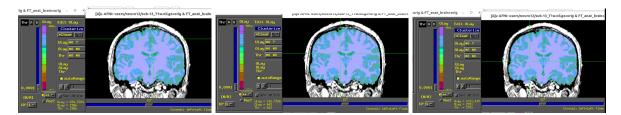


Figure 2: In order to: gray_matter,ventricle and white_matter.

Final mask:

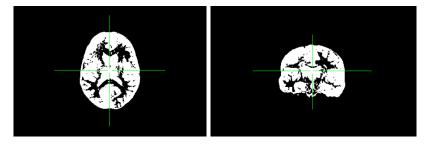


Figure 3: Gray Matter Mask.

Section 2: Pre-process Functional Image

I run the steps as below:

(About the slice time correction, we have the slice order, not time. So, I use the '3dinfo' function. This data has a time step equal 2s and the number of slice is 32. So dt equal 0.066s. I save the slice time in a file and use 'tpattern' option in '3dTshift' function.)

step 1: slice time correction

```
3dTshift -tpattern '@time_slice.txt' -prefix sub - 13_t ask_t shift
sub - 13_t ask - visualoddballwithbuttonresponsetotargetstimuli_run - <math>01_b old.nii.gz (3)
```

step 2: motion correction

```
3dvolreg - verbose - zpad1 - base \quad sub - 13_t ask_t shift + orig \\ - 1Dfile \quad dfile.FT1.1D - prefix \quad sub - 13_t ask.volreg \\ - cubic \quad - 1Dmatrix_s avemat.FT1.vr.aff12.1D \quad sub - 13_t ask_t shift + orig \quad \textbf{(4)}
```

step 3: normalize to mean

```
3dTstat - mean - prefix meanfunc1.nii.gz sub - 13_task.volreg + orig \\ 3dcalc - a sub - 13_task.volreg + orig - b meanfunc1.nii.gz \\ - expr'(a-b)/(abs(b)+1) * astep(b,200) * 100' - prefixsub - 13_task.scale \\ 3dAutomask - clfrac 0.4 - prefix clean_base.nii.gzmeanfunc1.nii.gz \\ 3dBandpass - input sub - 13_task.scale + orig - mask clean_base.nii.gz - blur 5 \\ - band 0.01 0.1 - prefix sub - 13_task.scale.smooth (5)
```

In figure, you can see the functional image before and after pre-processing:

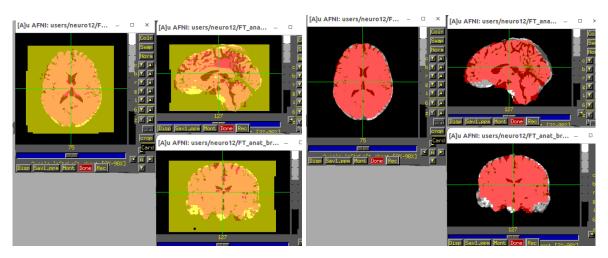


Figure 4: Pre-processing functional image; left: before, right: after pre-processing (clean base)

Section 3: Align functional and structural image

Align EPI and anatomical datasets by using a below function:

```
align\_epi\_anat.py - anat2epi - anat FT_anat_brain + orig
- anat_has_skull \quad no \quad - suffix_al_junk
- epiclean_base.nii.gz \quad - epi_base0
- epi_strip \quad 3dAutomask
- cost \quad nmi \quad - giant_move \quad - rigid_body
- volreg \quad off \quad - tshiftoff
(6)
```

Align anatomical to epi:

```
cat_{m}atvec \quad FT_{a}nat_{b}rain_{a}l_{j}unk_{m}at.aff12.1D - I > func2mri_{w}arp.1D
3dAllineate - baseFT_{a}nat_{b}rain + orig
-input \quad clean_{b}ase.nii.gz
-1Dmatrix_{a}pply \quad func2mri_{w}arp.1D
-prefix \quad epimask2anat.nii.gz
(7)
```

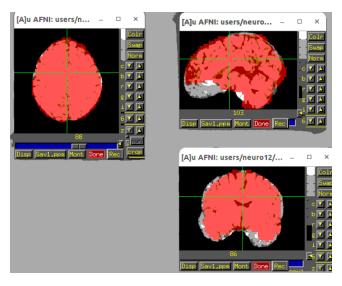


Figure 5: Align epi to anat; underlay: FT_anat_brain_al_junk, overlay: clean_base

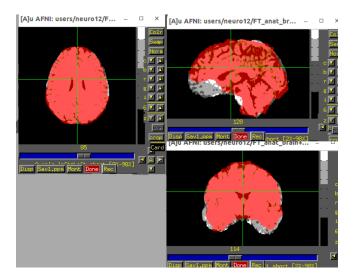


Figure 6: Align anat to epi; underlay: FT_anat_brain, overlay: epimask2anat

Section 4: Regressors & GLM

The onset time of each stimulus (usual and unusual), the subject response, the contrast and the mean of these two type of stimulus is considered as a regressors. I use the '3dDeconvolve' function as below:

```
3dDeconvolve - input sub - 13_task.scale.smooth + orig
                                   -polort 0
                                 -num_s timts 3
                 -stim_times 1 visual_time.txt 'BLOCK(0.2, 1)'
                              -stim_label 1 visual
                 -stim_times 2 oddball<sub>t</sub>ime.txt 'BLOCK(0.2, 1)'
                              -stim_label 2 oddball
              -stim_times_FSL 3 response_time.txt 'dmBLOCK(1)'
                             -stim_label 3 response
                                    -jobs 2
                      -qltsym 'SYM: visual -oddball'
                              - glt_1abel \quad 1 \quad V - Odd
                 -gltsym 'SYM: 0.5*visual +0.5*oddball'
                            -qlt_label 2 mean.VOdd
              - fout - tout - x1D \quad X.xmat.1D \quad - xjpeg \quad X.jpg
                              -errts errts.sub - 13
                             -bucket stats.sub - 13
                                                                                (8)
```

The GLM coefficient is mapped to anatomical image by using '3dAllineate' function:

```
3dAllineate -base FT_anat_brain + orig \\ -input stats.sub - 13 + orig \\ -1Dmatrix_apply func2mri_warp.1D \\ -prefix GLM_stats2anat.nii.gz 
(9)
3dAllineate -baseFT_anat_brain + orig \\ -input errts.sub - 13 + orig \\ -1Dmatrix_apply func2mri_warp.1D \\ -prefix GLM_errts2anat.nii.gz
```

(10)

Now, I show each regressors coefficient on the anatomical images. The region with higher value were more active for the task.

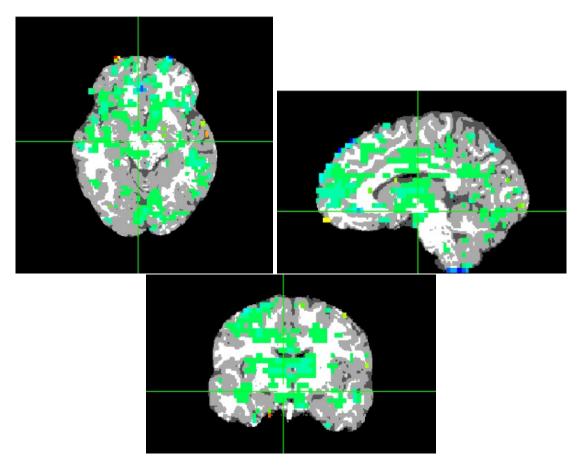


Figure 7: Active region when the normal stimulus is shown. ((p < 0.05))

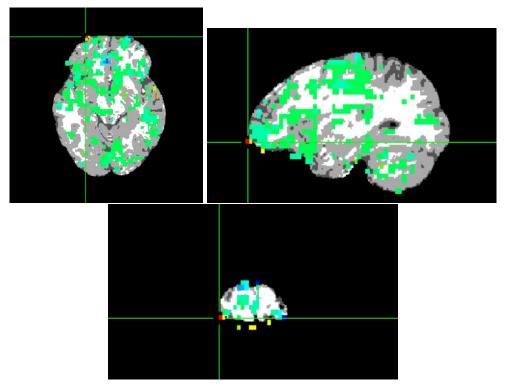


Figure 8: The most active region when the normal stimulus is shown. ((p < 0.05))

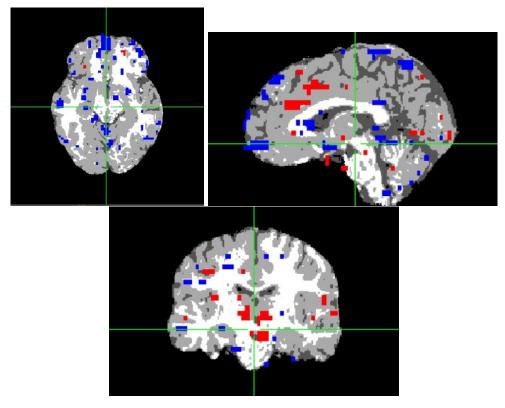


Figure 9: Active region when the oddball stimulus is shown. ((p < 0.05))

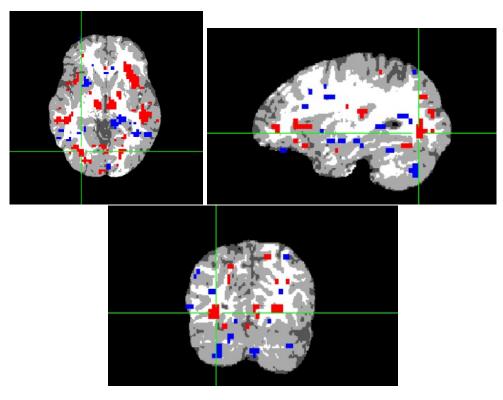


Figure 10: The most active region when the oddball stimulus is shown. ((p < 0.05))

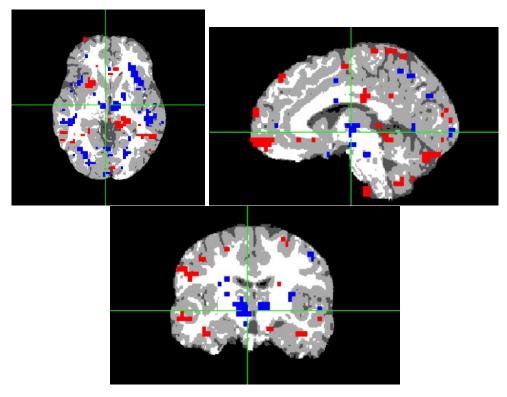


Figure 11: The most active region in the response time. ((p < 0.05))

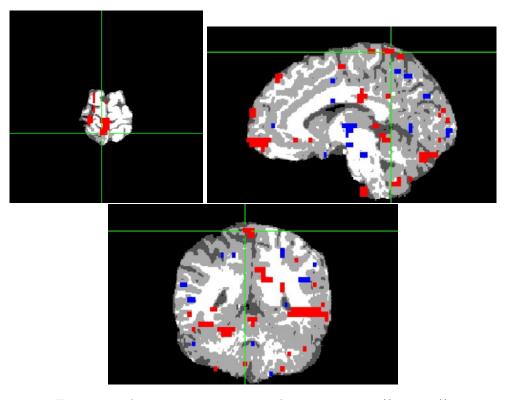


Figure 12: The most active region in the response time. ((p < 0.05))

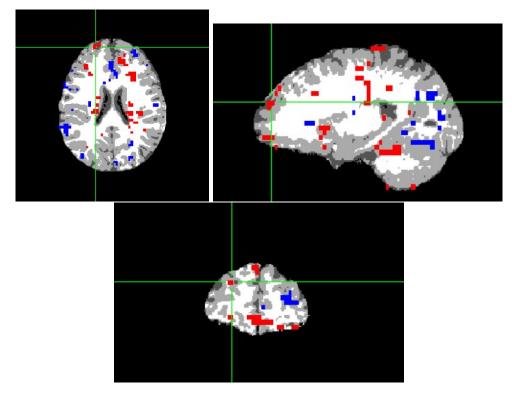


Figure 13: The difference regions' activities (normal - oddball). ((p < 0.05))

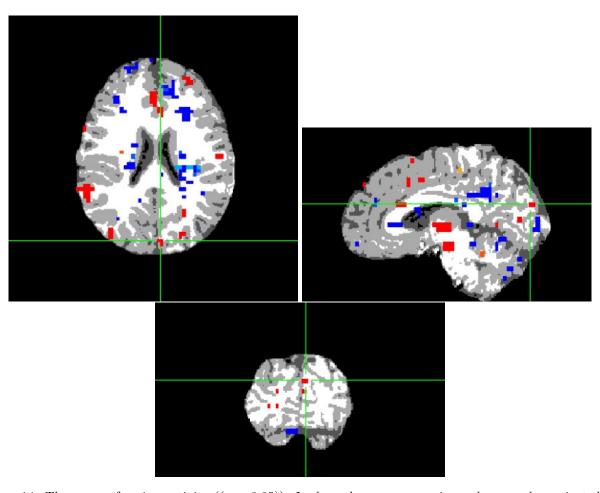


Figure 14: The mean of region activity ((p < 0.05)). It show the common regions where are be activated for both stimulus.