QUEEN'S NEUROECONOMICS LABORATORY



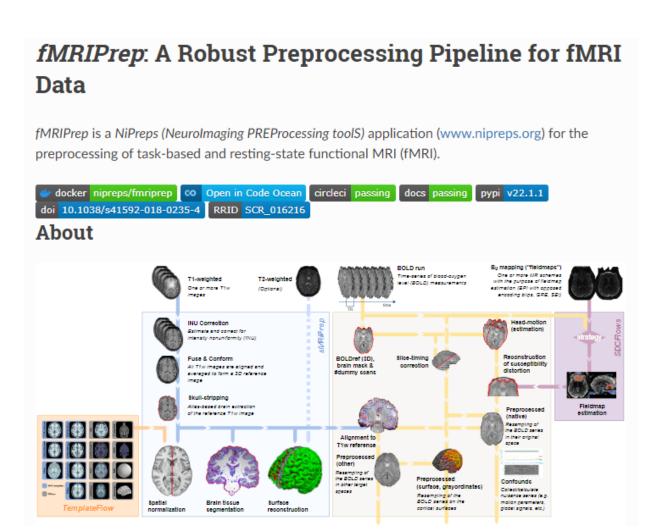
Neuroscience of Decision-Making | Lab Director: Dr. Anita Tusche

fMRIprep: A Robust Preprocessing Pipeline for fMRI Data

QUEEN'S NEUROECONOMICS LABORATORY



Neuroscience of Decision-Making | Lab Director: Dr. Anita Tusche



- ➤ Installation
- **▶** BIDs format
- A brief overview of how to use fmriprep

 Preprocessing steps Running the analysis

 Output Examining the preprocessed data

 Tips
- ➤ Conclusion/discussion



Installation



Installation







```
#!/bin/bash
#SBATCH -J fmriprep
#SBATCH --account=def-hpcg1879
#SBATCH --partition=tusch
#SBATCH -- gos=tusch
#SBATCH --time=72:0:0
#SBATCH --cpus-per-task=16
#SBATCH --mem-per-cpu=2GB
#SBATCH --job-name=fmriprep
#SBATCH --mail-type=ALL
#SBATCH --mail-user=remi.janet@queensu.ca
#SBATCH -o log %x-%A-%a.out
#SBATCH -e log_%x-%A-%a.err
# For Singularity version >= 2.5
LOAD="module load singularity/3.6"
BUILD="singularity build fmriprep-20.2.3.simg \
        docker://nipreps/fmriprep:20.2.3"
eval ${LOAD}
eval ${BUILD}
```

https://www.nipreps.org/apps/singularity/



```
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#SBATCH --account=def-hpcg1879
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Python 3.7+



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eval ${LOAD}
eval ${BUILD}
```

https://www.nipreps.org/apps/singularity/

Python 3.7+

- \$ python -m pip install fmriprep
- # Check your installation
- \$ fmriprep --version

Note

If you try running the command above, you may get the following error:

ImportError: cannot import name md5. This can happen sometimes with Python version 2.7; to fix this error, install a more recent version of Python, and then rerun the command:

https://fmriprep.org/en/stable/installation.html



```
#!/bin/bash
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https://www.nipreps.org/apps/singularity/

Python 3.7+

\$ python -m pip install fmriprep

Check your installation

\$ fmriprep --version

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If you try running the command above, you may get the following error:

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https://fmriprep.org/en/stable/installation.html

Docker

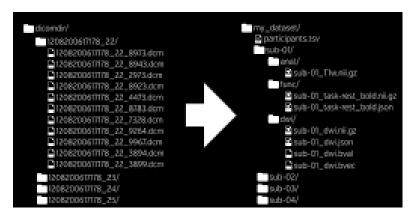
https://www.nipreps.org/apps/docker/

https://andysbrainbook.readthedocs.io/en/latest/OpenScience/OS/fMRIPrep.html#fmriprep



BIDs format







BIDS (Brain Imaging Data Structure)

Standarized format for the organization and description of neuroimaging and corresponding behavioral data.

Easily **shared** and **understood** by other researchers. **Reproducibility and Data Sharing**

Different packages that can be used to convert your data into BIDS format, such as dcm2bids, heudiconv, bidscoin, bidskit, etc.



- 1- Create Project folder
- 2- Download data
- 3- CMake and pip Installation
- 4- dcm2niix Installation
- 5- dcm2bids Installation
- 6- Setting Up Your Configuration File
- 7- Running the dcm2bids command
- 8- Double-check BIDS



1- Create Project folder

cd \$HOME mkdir BIDS_tutorial

2- Download data

The data can be found here. https://drive.google.com/file/d/1Gx4GdWJEvT5O-2MYjDoyJIiirSpVFV4O/view

mv/global/project/hpcg1879/Remi/BIDS_tutorial/OpenScience \$HOME/BIDS_tutorial



3- CMake and pip Installation

If you are working on a HPC (cluster) then they are already installed.

which cmake which pip

Otherwise, follow these steps

```
cd curl https://bootstrap.pypa.io/get-pip.py -o get-pip.py python get-pip.py --user export PATH= "/Users/$USER/Library/Python/2.7/bin/:$PATH" cd ~/Downloads tar -zxvf cmake-3.16.3-*-x86_64.tar.gz export PATH="~/Downloads/cmake-3.16.3-Darwin-x86_64/CMake.app/Contents/bin/:$PATH" pip install cmake
```



4- dcm2niix Installation

```
cd ~
git clone https://github.com/rordenlab/dcm2niix.git
cd dcm2niix
mkdir build
cd build
cmake ..
make
```

Add dcm2nix to your path

export PATH="\$HOME/dcm2niix/build/bin/:\$PATH"

5- dcm2bids Installation

cd \$HOME/BIDS_tutorial
module load python
pip install --user dcm2bids



6- Setting Up Your Configuration File

See the file I shared with you.

It is a .txt file. However, you can just save it and change the extension to .json.

7- Running the dcm2bids command

dcm2bids -d \$HOME/BIDS_tutorial/BIDS_tutorial_data -p 001 -c \$HOME/BIDS_config.json -o \$HOME/BIDS_tutorial --forceDcm2niix



This is true if you only have one session per subject. If you have multiple session then copy this line



8- Double-check BIDS

You can use the <u>BIDS validator</u> to ensure that your data are BIDS-compliant https://bids-standard.github.io/bids-validator/

BIDs format



Files



Anatomical scan

- sub-101_T1w.nii.gz
- sub-101_T1w.json

Functional scans

- sub-102_task-choose_run-01_events.tsv
- sub-102_task-choose_run-01_bold.nii.gz
- sub-102 task-choose run-01 bold.nii
- sub-102_task-choose_run-01_bold.json

Fieldmaps

- sub-101_phasediff.nii.gz
- sub-101_phasediff.json
- sub-101_magnitude2.nii.gz
- sub-101_magnitude2.json
- sub-101_magnitude1.nii.gz
- sub-101_magnitude1.json

Metadata files

- participants.tsv
- dataset_description.json
- 🗐 bids_filter.json
- .bidsignore

dataset_description.json

```
{
    "Authors": [
        "Remi Janet",
        "John-Dennis (Jack) Parsons",
        "Anita Tusche",
        "Hilke Plassman"
],
    "Acknowledgements": "HP supplied raw data to AT which was converted to BIDS by JP. and RJ",
    "Name": "fMRI Food Regulation Study",
    "BIDSVersion": "1.6.0"
}
```

participants.tsv

```
participant id scanner id
                                        female
                                age
sub-101 CC0058
                        0
sub-102 CC0003
sub-103 CC0007
sub-107 CC0025
sub-108 CC0012 28
sub-109 CC0033
                        0
sub-110 CC0054 32
sub-112 CC0027
sub-114 CC0049
                        0
sub-116 CC0029
sub-118 CC0005
sub-120 CC0024 28
                        0
sub-122 CC0053
sub-123 CC0021 25
sub-126 CC0041 29
                        0
sub-130 CC0008 26
                        0
```

.bidsignore

```
*.html
logs/
figures/
*_xfm.*
*.surf.gii
*_boldref.nii.gz
*_bold.func.gii
*_mixing.tsv
*_AROMAnoiseICs.csv
*_timeseries.tsv
```

BIDs format



Useful links to go further:

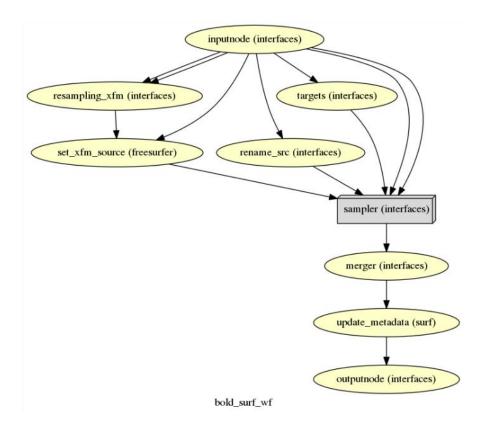
- https://openneuro.org/
- https://www.mathworks.com/matlabcentral/fileexchange/42997-xiangruili-dicm2nii
- https://bids-apps.neuroimaging.io/
- https://bids-standard.github.io/bids-validator/
- https://openneuro.org/
- https://bids-standard.github.io/bids-starter-kit/dataset_examples.html

https://www.mathworks.com/matlabcentral/fileexchange/42997-xiangruili-dicm2nii

setpref('dicm2nii_gui_para', 'bidsForceGUI', true)
dicm2nii(DCfilesList, 'F:\path_to_your_files\rawfiles', 'bids')



fMRIprep usage





Preprocessing steps – Running the analysis



With slurm. Use the ".slurm" files and launch them with the command sbatch --array=[sub_start-sub_end] fmriprep_scrip.slurm



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```
export STUDY="/global/project/hpcg1879/Remi/Altruism AT"
BIDS DIR="$STUDY"
DERIVS DIR="derivatives/fmriprep-20.2.1"
FS_DIR="${BIDS_DIR}/${DERIVS_DIR}/freesurfer"
# Prepare some writeable bind-mount points.
TEMPLATEFLOW HOST HOME=$HOME/.cache/templateflow
FMRIPREP HOST CACHE=$HOME/.cache/fmriprep
mkdir -p ${TEMPLATEFLOW HOST HOME}
mkdir -p ${FMRIPREP HOST CACHE}
# Prepare derivatives folder
mkdir -p ${DERIVS_DIR}
# Make sure FS LICENSE is defined in the container.
export SINGULARITYENV FS LICENSE=$HOME/free surfer license.txt
# Designate a templateflow bind-mount point
export SINGULARITYENV TEMPLATEFLOW HOME="/templateflow"
SINGULARITY_CMD="singularity run --cleanenv -B ${BIDS_DIR}:/Altruism_AT -B ${TEMPLATEFLOW_HOST_HOME}:${SINGULARITYENV_TEMPLATEFLOW_HOME} fmriprep-20.2.1.simg"
# Parse the participants.tsv file and extract one subject ID from the line corresponding to this SLURM task.
subject = \$ ( sed -n -E "\$ ((\$\{SLURM\_ARRAY\_TASK\_ID\} + 1))s/sub - (\$) \cdot */1/gp" \$\{STUDY\}/participants.tsv )
echo $subject
# Remove IsRunning files from FreeSurfer
# find ${FS_DIR}/sub-$subject/ -name "*IsRunning*" -delete
SINGULARITY LOAD="module load singularity/3.8"
# DON"T FUCK WITH THIS LINE RIGHT NOW
RUN FMRIPREP CMD="${SINGULARITY CMD} /Altruism AT /Altruism AT/${DERIVS DIR} participant --participant-label ${subject} --skip bids validation --ignore fieldmaps --output-spaces MNI152NLin2009cAsym"
eval ${SINGULARITY LOAD}
eval ${RUN FMRIPREP CMD}
# Output results to a table
echo "sub-$subject ${SLURM_ARRAY_TASK_ID} $exitcode" \
      >> ${SLURM JOB NAME}.${SLURM ARRAY JOB ID}.tsv
echo Finished tasks ${SLURM ARRAY TASK ID} with exit code $exitcode
exit $exitcode
```

>> \${SLURM JOB NAME}.\${SLURM ARRAY JOB ID}.tsv

exit \$exitcode

echo Finished tasks \${SLURM ARRAY TASK ID} with exit code \$exitcode



With slurm. Use the ".slurm" files and launch them with the command sbatch --array=[sub_start-sub_end] fmriprep_scrip.slurm

```
export STUDY="/global/project/hpcg1879/Remi/Altruism AT"
BIDS DIR="$STUDY"
DERIVS DIR="derivatives/fmriprep-20.2.1"
FS_DIR="${BIDS_DIR}/${DERIVS_DIR}/freesurfer"
# Prepare some writeable bind-mount points.
TEMPLATEFLOW HOST HOME=$HOME/.cache/templateflow
FMRIPREP HOST CACHE=$HOME/.cache/fmriprep
mkdir -p ${TEMPLATEFLOW HOST HOME}
mkdir -p ${FMRIPREP HOST CACHE}
# Prepare derivatives folder
mkdir -p ${DERIVS_DIR}
# Make sure FS LICENSE is defined in the container.
export SINGULARITYENV FS LICENSE=$HOME/free surfer license.txt
# Designate a templateflow bind-mount point
export SINGULARITYENV TEMPLATEFLOW HOME="/templateflow"
SINGULARITY CMD="singularity run --cleanenv -B ${BIDS_DIR}:/Altruism_AT -B ${TEMPLATEFLOW_HOST_HOME}:${SINGULARITYENV_TEMPLATEFLOW_HOME} fmriprep-20.2.1.simg"
# Parse the participants.tsv file and extract one subject ID from the line corresponding to this SLURM task.
subject=\{(sed -n -E "$((\{SLURM_ARRAY_TASK_ID\} + 1))s/sub-((S*)).*/(1/gp" $\{STUDY\}/participants.tsv )\}
echo $subject
# Remove IsRunning files from FreeSurfer
# find ${FS DIR}/sub-$subject/ -name "*IsRunning*" -delete
SINGULARITY LOAD="module load singularity/3.8"
# DON"T FUCK WITH THIS LINE RIGHT NOW
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eval ${SINGULARITY LOAD}
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# Output results to a table
echo "sub-$subject ${SLURM_ARRAY_TASK_ID} $exitcode" \
```



With Python. Use bash code: bash frmiprepcode.sh



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```
#User inputs:
bids_root_dir=$HOME/Desktop/Flanker
subj=08
nthreads=4
mem=20 #gb
container=docker #docker or singularity
#Convert virtual memory from gb to mb
mem='echo "${mem//[!0-9]/}"\ #remove gb at end
mem_mb=`echo $(((mem*1000)-5000))` #reduce some memory for buffer space during pre-processing
export FS LICENSE=$HOME/Desktop/Flanker/derivatives/license.txt
#Run fmriprep
if [ $container == singularity ]; then
  unset PYTHONPATH; singularity run -B $HOME/.cache/templateflow:/opt/templateflow $HOME/fmriprep.simg
    $bids root dir $bids root dir/derivatives \
    participant \
    --participant-label $subj \
    --skip-bids-validation \
    --md-only-boilerplate \
    --fs-license-file $HOME/Desktop/Flanker/derivatives/license.txt \
    --fs-no-reconall \
    --output-spaces MNI152NLin2009cAsym:res-2 \
    --nthreads $nthreads \
    --stop-on-first-crash \
    --mem_mb $mem_mb \
    -w $HOME
  fmriprep-docker $bids_root_dir $bids_root_dir/derivatives \
    participant \
    --participant-label $subj \
    --skip-bids-validation \
    --md-only-boilerplate \
    --fs-license-file $HOME/Desktop/Flanker/derivatives/license.txt \
    --fs-no-reconall \
    --output-spaces MNI152NLin2009cAsym:res-2 \
    --nthreads $nthreads \
    --stop-on-first-crash \
    --mem_mb $mem_mb \
    -w $HOME
```

 $https://andysbrainbook.readthedocs.io/en/latest/OpenScience/OS/fMRIPrep_Demo_2_RunningAnalysis.html\\$



With Python. Use bash code: bash frmiprepcode.sh

```
#User inputs:
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nthreads=4
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    $bids root dir $bids root dir/derivatives \
    participant \
    --participant-label $subj \
    --skip-bids-validation \
    --md-only-boilerplate \
    --fs-license-file $HOME/Desktop/Flanker/derivatives/license.txt \
    --fs-no-reconall \
    --output-spaces MNI152NLin2009cAsym:res-2 \
    --nthreads $nthreads \
    --stop-on-first-crash \
    --mem_mb $mem_mb \
    -w $HOME
  fmriprep-docker $bids_root_dir $bids_root_dir/derivatives \
    participant \
    --participant-label $subj \
    --skip-bids-validation \
    --md-only-boilerplate \
    --fs-license-file $HOME/Desktop/Flanker/derivatives/license.txt \
    --fs-no-reconall \
    --output-spaces MNI152NLin2009cAsym:res-2 \
    --nthreads $nthreads \
    --stop-on-first-crash \
    --mem_mb $mem_mb \
    -w $HOME
```

It will process only subject 8 in this example.

https://andysbrainbook.readthedocs.io/en/latest/OpenScience/OS/fMRIPrep_Demo_2_RunningAnalysis.html



Output – Examining the preprocessed data



Output – Examining the preprocessed data

sub-162
sub-163
sub-164
sub-164
sub-164
dataset_description.json
desc-aparcaseg_dseg.tsv
desc-aseg_dseg.tsv
sub-101.html
sub-102.html

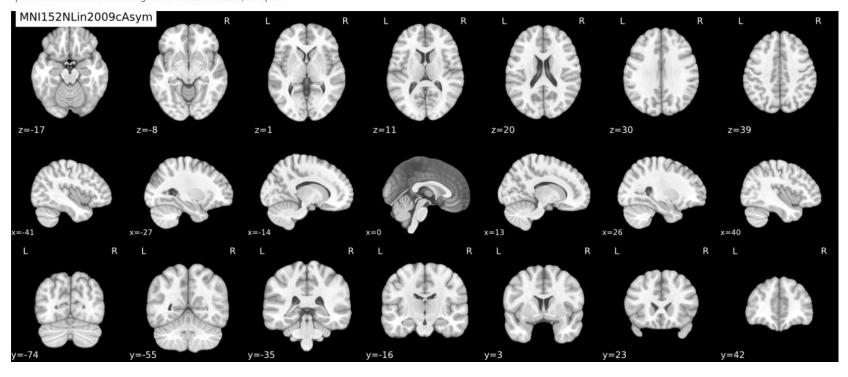


Anatomical output

Spatial normalization of the anatomical T1w reference

Results of nonlinear alignment of the T1w reference one or more template space(s). Hover on the panels with the mouse pointer to transition between both spaces.

Spatial normalization of the T1w image to the MNI152NLin2009cAsym template.



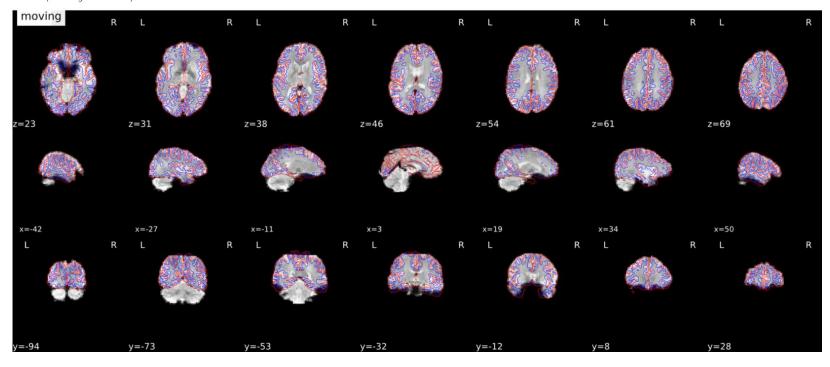
Make sure to check the alignment not only between the outlines of the brain, but also the internal structures such as the ventricles.



Functional outputs

Alignment of functional and anatomical MRI data (surface driven)

bbregister was used to generate transformations from EPI-space to T1w-space. Note that Nearest Neighbor interpolation is used in the reportlets in order to highlight potential spin-history and other artifacts, whereas final images are resampled using Lanczos interpolation.

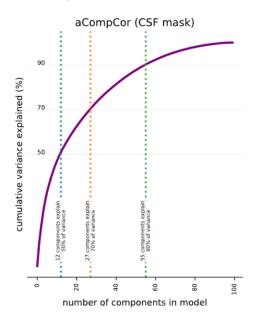


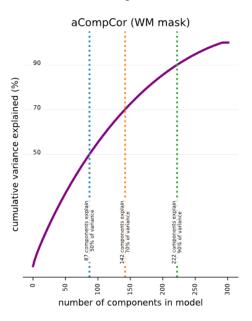
Make sure that the internal structures are well aligned

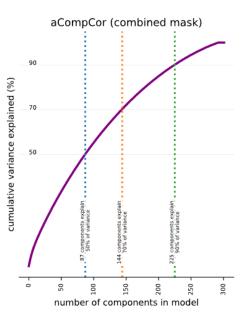


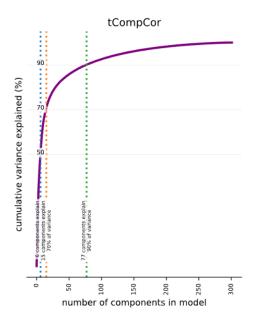
Variance explained by t/aCompCor components

The cumulative variance explained by the first k components of the *t/aCompCor* decomposition, plotted for all values of *k*. The number of components that must be included in the model in order to explain some fraction of variance in the decomposition mask can be used as a feature selection criterion for confound regression.

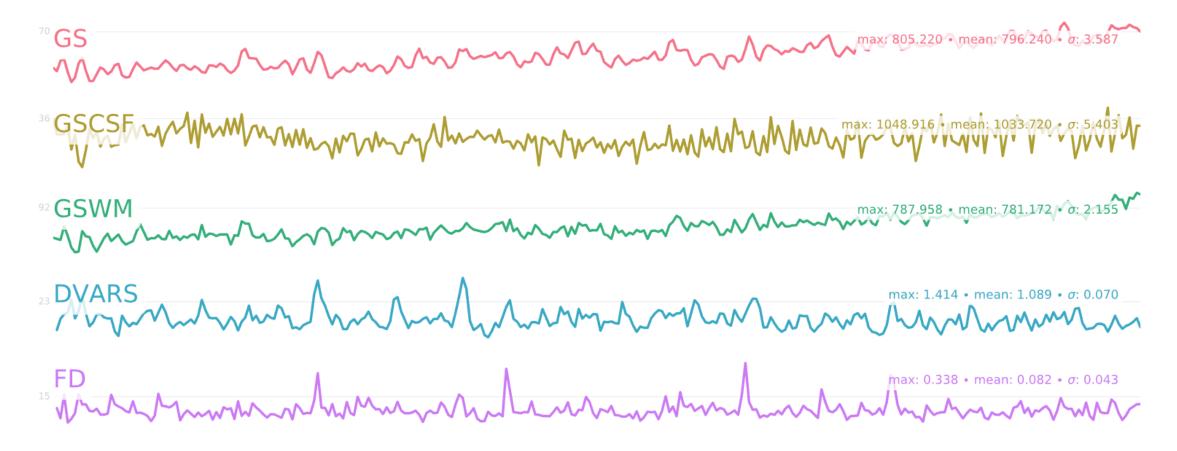






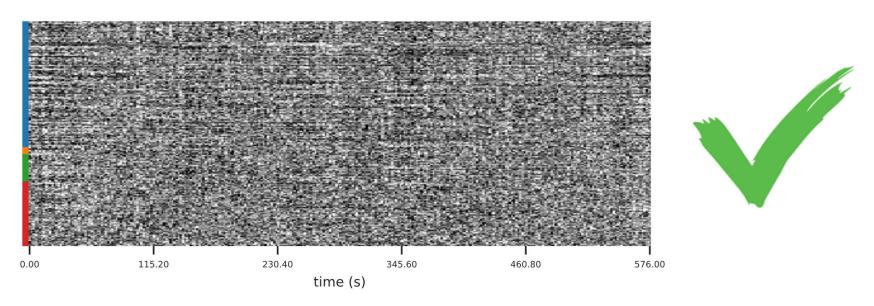




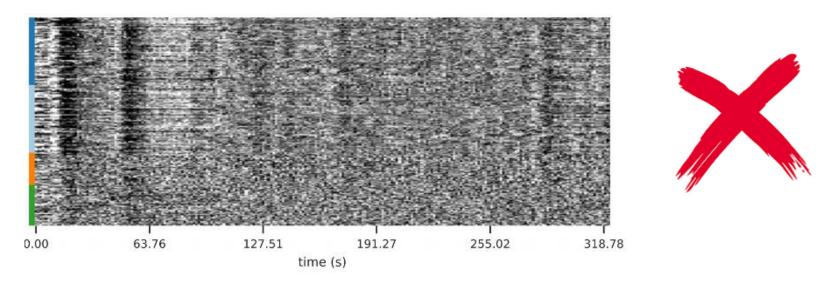


Rule of thumb = warning if mean FD (σ FD) > 0.2

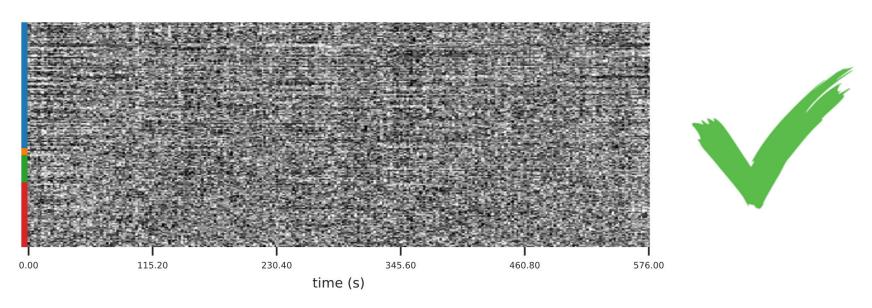




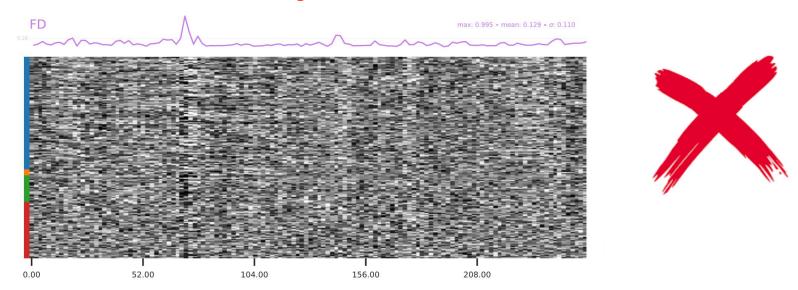
Any sudden changes in motion may be reflected in uniform changes across the entire column for that timepoint.







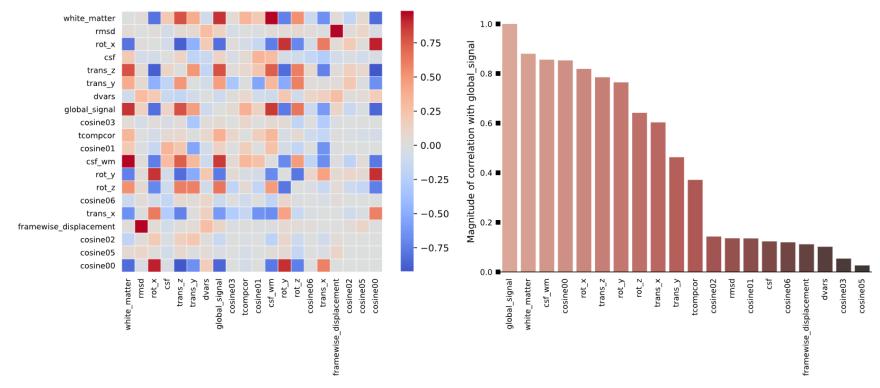
Any sudden changes in motion may be reflected in uniform changes across the entire column for that timepoint.





Correlations among nuisance regressors

Left: Heatmap summarizing the correlation structure among confound variables. (Cosine bases and PCA-derived CompCor components are inherently orthogonal.) Right: magnitude of the correlation between each confound time series and the mean global signal. Strong correlations might be indicative of partial volume effects and can inform decisions about feature orthogonalization prior to confound regression.



The bar chart on the right shows the correlation of different regressors with respect to global signal; those components that show a high degree of correlation may be candidates for nuisance regression.



Tips





Danger

Slice timing correction in *fMRIPrep* is referenced to the middle slice by default, which leads to a time shift in the volume onsets by 0.5 TR (repetition time). For example, assuming a TR of 2s, original onsets of 0, 2, and 4s would be shifted to 1, 3, and 5s, respectively. In case you did execute slice timing correction, you must check that subsequent analyses (e.g., general linear modeling) consider the right onset shifts. For example, when specifying a first-level model, you should set parameters in your software package or first-level model function accordingly (e.g., select the middle slice as reference). Alternatively, you could manually adjust the volume onsets (e.g. as mentioned in the example above from [0, 2, 4] to [1, 3, 5]) or the event onsets accordingly.

Further information on this issue is found at this blog post (with thanks to Russell Poldrack and Jeanette Mumford).



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TL/DR: If you are analyzing fMRIPrepped data with SPM or FSL, you are good to go. If you are analyzing it using nilearn, AFNI, or custom code, then you need to do some extra work to ensure that your statistical model is properly aligned with your slice-time-corrected data.



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Further information on this issue is found at this blog post (with thanks to Russell Poldrack and Jeanette Mumford).



TL/DR: If you are analyzing fMRIPrepped data with SPM or FSL, you are good to go. If you are analyzing it using nilearn, AFNI, or custom code, then you need to do some extra work to ensure that your statistical model is properly aligned with your slice-time-corrected data.

Imagine you have 45 subjects. You already preprocessed all of them, and your supervisor realized he has 6 more subjects you can add to your sample.

Then it can be possible that you get error message if you add these six subjects to your previous folder.

=> Create a new folder from scratch and launch the preprocessing again



Danger

Slice timing correction in *fMRIPrep* is referenced to the middle slice by default, which leads to a time shift in the volume onsets by 0.5 TR (repetition time). For example, assuming a TR of 2s, original onsets of 0, 2, and 4s would be shifted to 1, 3, and 5s, respectively. In case you did execute slice timing correction, you must check that subsequent analyses (e.g., general linear modeling) consider the right onset shifts. For example, when specifying a first-level model, you should set parameters in your software package or first-level model function accordingly (e.g., select the middle slice as reference). Alternatively, you could manually adjust the volume onsets (e.g. as mentioned in the example above from [0, 2, 4] to [1, 3, 5]) or the event onsets accordingly.

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It take approximatively a day to process one subject.

Parallelization if recommended.

If you want to speed up the process use:
--fs-no=reconall