

# 1. Preprocessing MEG Python

Jupyter notebook with the **entire preprocessing and source reconstruction pipeline** for the SDMT dataset using the OSL Python library from the Oxford Centre for Human Brain Activity (OHBA) Ananalysis Group.

**Reference:** Quinn, A. J., van Es, M. W., Gohil, C., & Woolrich, M. W. (2023). OHBA Software Library in Python (OSL) (Version 0.2.0) [Computer software].

<https://doi.org/10.5281/zenodo.6875060>

## Required input files/folders to run this Notebook:

- **raw\_dataset\_dir:** folder containing all raw. fif files for the SDMT data
- **T1\_niifiles:** folder containing all the subjects' MRI scans (.nii files)
- **T1\_MNI\_niifiles:** folder containing all the subjects' already co-registered MRI scans in MATLAB (.nii.gz files)
- **parcellation file:**  
"fmri\_d100\_parcellation\_with\_PCC\_reduced\_2mm\_ss5mm\_ds8mm.nii.gz" **with 38 parcels** needs to be 4D (3 dimensions for spatial coordinates and 1 for each parcel) with sform\_code = 1
- **Parcels\_38\_names.txt (extra):** .txt file with the names of the 38 parcels of the parcellation file

## Install necessary packages

```
In [1]: #!pip install mne
        #!pip install fieldtrip2mne
        #!pip install fslypy
        #!pip install osl                    #(from https://github.com/OHBA-analysis/osl
        #!pip install numpy --upgrade
        #!pip install pymatreader
        #!pip install trame
        #!pip install ipywidgets
        #!pip install numpy==1.22.4
        #!pip install pandas~=1.3.5
        #!pip install numba~=0.53.1
```

## Import needed packages

```
In [8]: import os
        import glob
        import mne
        import numpy
        import shutil
        import osl
        # opencv_python should be in same directory as the Jupyterlab Notebook
        import opencv_python    # opencv_python is a manually edited package (cont
                                # stackoverflow.com/questions/68916893/typeerror-nu
                                # https://github.com/opencv/opencv/issues/23822
                                # should be fixed in release 4.9.x of opencv-pythor
```

## Load in dataset - raw .fif files for all patients

```
In [4]: raw_dataset_dir = "/home/data/vub_ms/MEGDATA_ANA/SDMT/DATA/"

all_fif_files = sorted(glob.glob(raw_dataset_dir + '*.fif'))
print(len(all_fif_files))
```

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## Remove subjects that should be excluded

```
In [5]: # The subjects excluded in the preprocessing phase relate to prior findings
# - issues with MRI scan
# - issues in the headmodelling step
# - missing DIODE channel (in the event extraction)
subject_IDs_to_exclude = ['2304', '2180', '2202', '2357', '2371', '1035', '0
                        '1063', '2174',
                        '2412', '2413', '2233', '2234']

file_IDs = []
for filename in all_fif_files:
    file_IDs.append(filename[-21:-17]) #gets the 'code' Number per patient

# Function to quickly find matching elements between 2 lists
def find_matching_index(list1, list2):

    inverse_index = { element: index for index, element in enumerate(list1)

    return [index
            for index,element in enumerate(list2) if element in inverse_index]

# Use the function to find which subject IDs have to be removed from the tot
indices_to_remove = find_matching_index(subject_IDs_to_exclude,file_IDs)
print(indices_to_remove)
```

[2, 29, 55, 57, 63, 75, 76, 94, 112, 116, 121, 128, 129]

```
In [9]: #Retained .fif files (filepaths)
fif_files = [ele for idx, ele in enumerate(all_fif_files) if idx not in indi

#Retained indices (subject ID
file_IDs_retained = [ele for idx, ele in enumerate(file_IDs) if idx not in i

print(len(fif_files))
```

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```
In [10]: # Quick example to see if a single .fif file can be loaded
parceled_data = mne.io.read_raw_fif(fif_files[0], preload=True)
```

Opening raw data file /home/data/vub\_ms/MEGDATA\_ANA/SDMT/DATA/meg\_0922\_SD  
MT\_tsss\_mc.fif...

Range : 56000 ... 1037999 = 56.000 ... 1037.999 secs

Ready.

Reading 0 ... 981999 = 0.000 ... 981.999 secs...

/tmp/ipykernel\_2716435/3664972164.py:2: RuntimeWarning: This filename (/home/data/vub\_ms/MEGDATA\_ANA/SDMT/DATA/meg\_0922\_SDMT\_tsss\_mc.fif) does not conform to MNE naming conventions. All raw files should end with raw.fif, raw\_sss.fif, raw\_tsss.fif, \_meg.fif, \_eeg.fif, \_ieeg.fif, raw.fif.gz, raw\_sss.fif.gz, raw\_tsss.fif.gz, \_meg.fif.gz, \_eeg.fif.gz or \_ieeg.fif.gz  
parceled\_data = mne.io.read\_raw\_fif(fif\_files[0], preload=True)

## Add MRI info/files

```
In [12]: DIR_MRI_FILES = "/home/data/vub_ms/MEGNII/"
T1_niifiles = sorted(glob.glob(DIR_MRI_FILES + 'MEG*_MRI_T1.nii'))
T1_MNI_niifiles = sorted(glob.glob(DIR_MRI_FILES + 'MEG*_MRI_T1_MNI.nii.gz'))

# We only retain subjects with existing T1_MNI.nii file:
file_IDs_MRI = []
for filename in T1_MNI_niifiles:
    file_IDs_MRI.append(filename[-22:-18]) # gets the 'code' from the .nii

# Match between the patients we want to keep (for preprocessing), and valid
indices_to_keep = find_matching_index(file_IDs_retained, file_IDs_MRI)

# Retained T1.nii files
T1_niifiles = [ele for idx, ele in enumerate(T1_niifiles) if idx in indices_to_keep]

# Retained T1_MNI.nii.gz files
T1_MNI_niifiles = [ele for idx, ele in enumerate(T1_MNI_niifiles) if idx in indices_to_keep]

print(len(T1_niifiles), f'amount of T1.nii files for {len(fif_files)} patients')
print(len(T1_MNI_niifiles), f'amount of T1_MNI.nii files for {len(fif_files)} patients')

126 amount of T1.nii files for 126 patients
126 amount of T1_MNI.nii files for 126 patients
```

## Processing Pipeline - 1 subject

First, the Python package and all processing steps are tested for 1 single subject. This allows to learn how the package works, how each input needs to be specified, and how to solve certain bugs/errors.

### Step 1: Preprocessing

Preprocessing WITH removal of bad segments and channels

```
In [3]: from osl import preprocessing, utils

# Directory in which we save the preprocessed data
PREPROC_DIR = "/home/olivierb/processed_data"

## Order of preprocessing steps actually matters in the list of dictionaries

# Settings
config = """
preproc:
- resample: {sfreq: 250}

- filter: {l_freq: 0.01, h_freq: 70, method: iir, iir_params: {order: 5,
- notch_filter: {freqs: 50, notch_widths: 2}

- bad_segments: {segment_len: 500, picks: mag, significance_level: 0.1}
- bad_segments: {segment_len: 500, picks: grad, significance_level: 0.1}
- bad_segments: {segment_len: 500, picks: mag, mode: diff, significance_level: 0.1}
- bad_segments: {segment_len: 500, picks: grad, mode: diff, significance_level: 0.1}
- bad_channels: {picks: mag, significance_level: 0.1}
- bad_channels: {picks: grad, significance_level: 0.1}
```

```
- ica_raw: {picks: meg, n_components: 64}
- ica_autoreject: {apply: true}

- interpolate_bads: {}
"""

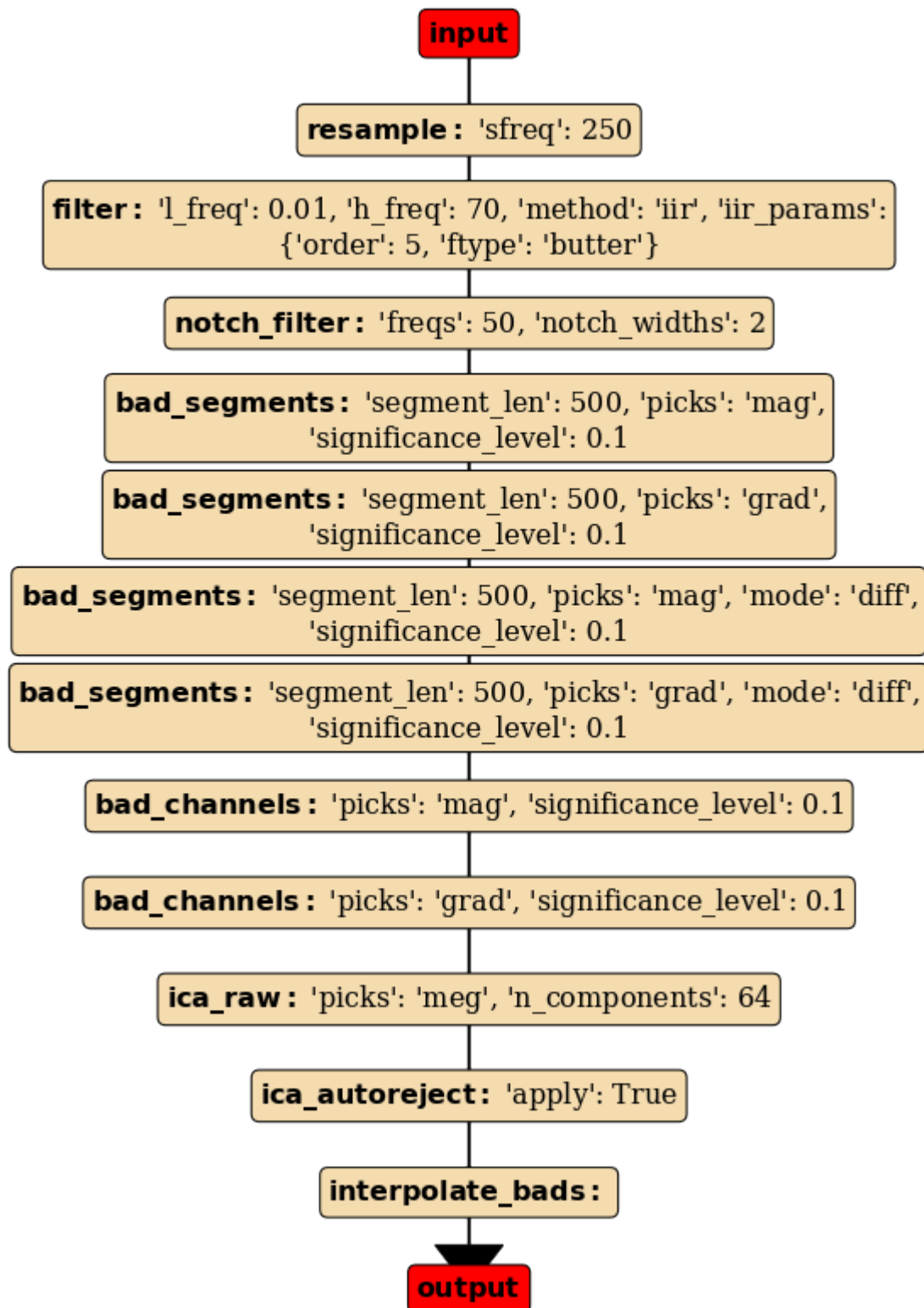
# Main preprocessing
preprocessing.run_proc_batch(
    config,
    fif_files[0],
    outdir=PREPROC_DIR,
    overwrite=True,
)
```

In [4]: `osl.preprocessing.plot_preproc_flowchart(config)`

```
/home/olivierb/osl/preprocessing/batch.py:566: UserWarning: FigureCanvasAgg is non-interactive, and thus cannot be shown
    fig.show()
```

Out[4]: (<Figure size 576x864 with 1 Axes>,  
<Axes: title={'center': 'OSL Preprocessing Recipe'}>)

# OSL Preprocessing Recipe



## Step 2: Coregistration

```

In [5]: #From: https://github.com/OHBA-analysis/osl/blob/main/examples/mrc\_meguk/can

from osl import source_recon, utils

# Settings
config = """
    source_recon:
    - extract_fiducials_from_fif: {}      # ok
    - compute_surfaces:

```

```

        include_nose: False

    - coregister:
        use_nose: False
        use_headshape: True
        n_init: 50
    """

# Directories

subject = "/home/olivierb/test/meg_0925_SDMT_tsss_mc/meg_0925_SDMT_tsss_mc_p

# Directory in which we will save the coregistered data
COREG_DIR = "/home/olivierb/coreg"

# Directory to the FSL library, same as the one used in MATLAB, supported by
FSL_DIR = "/usr/lib/fsl/6.0"

# Run coregistration

source_recon.setup_fsl(FSL_DIR)
source_recon.run_src_batch(
    config,
    src_dir=COREG_DIR,
    subjects=['meg_0925_SDMT_tsss_mc_preproc_raw.fif'], # name of the subj
    preproc_files=[subject], # path to preproc
    smri_files=[T1_MNI_niifiles[0]], # path to .nii file
)

```

### Step 3: Source reconstruction (Forward Modelling/Beamforming/Parcellation/Orthogonalisation)

```

In [ ]: from osl import source_recon
# '/home/olivierb/38_parcel.nii' => 38 parcels / sform_code=1

# Settings
config = """
    source_recon:
    - forward_model:
        model: Single Layer # ok: Single Shell in M
    - beamform_and_parcellate:
        freq_range: [1, 80]
        chantypes: [mag, grad] # standard for MEG
        rank: {meg: 60}
        parcellation_file: '/home/olivierb/38_parcel.nii'
        method: spatial_basis # ok => spatial_basis or
        orthogonalisation: symmetric # ok
    """

# Run beamforming and parcellation
FSL_DIR = "/usr/lib/fsl/6.0"
COREG_DIR = "/home/olivierb/coreg"
SUBJECT = subject = "/home/olivierb/processed_data/meg_0925_SDMT_tsss_mc/meg

source_recon.setup_fsl(FSL_DIR)
source_recon.run_src_batch(
    config,
    src_dir=COREG_DIR, # path to all the
    subjects=['meg_0925_SDMT_tsss_mc_preproc_raw.fif'], # name of the sub

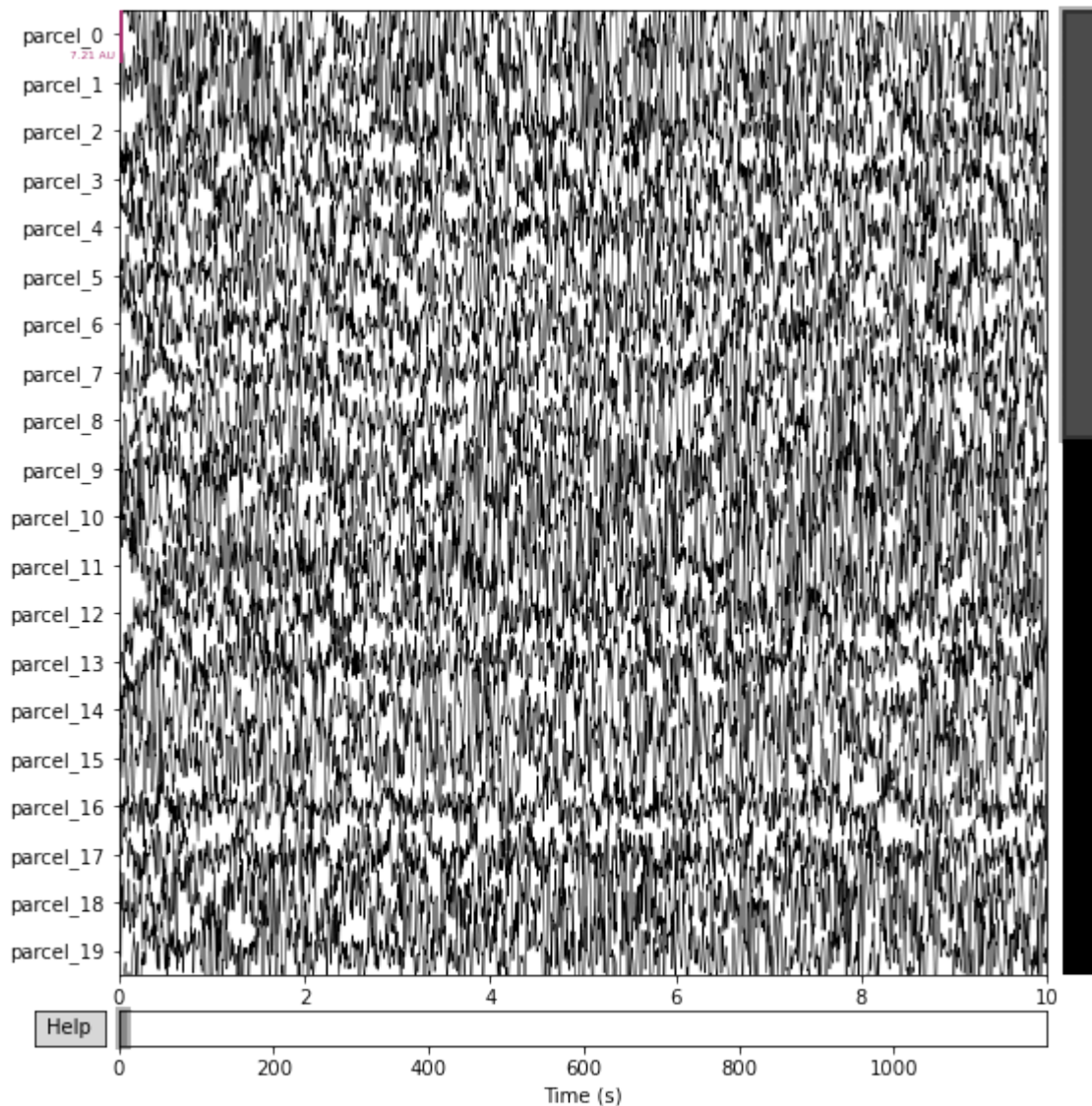
```

```
preproc_files=[subject]                                     # path to preproc  
)
```

### Plot result of one of parcelated file

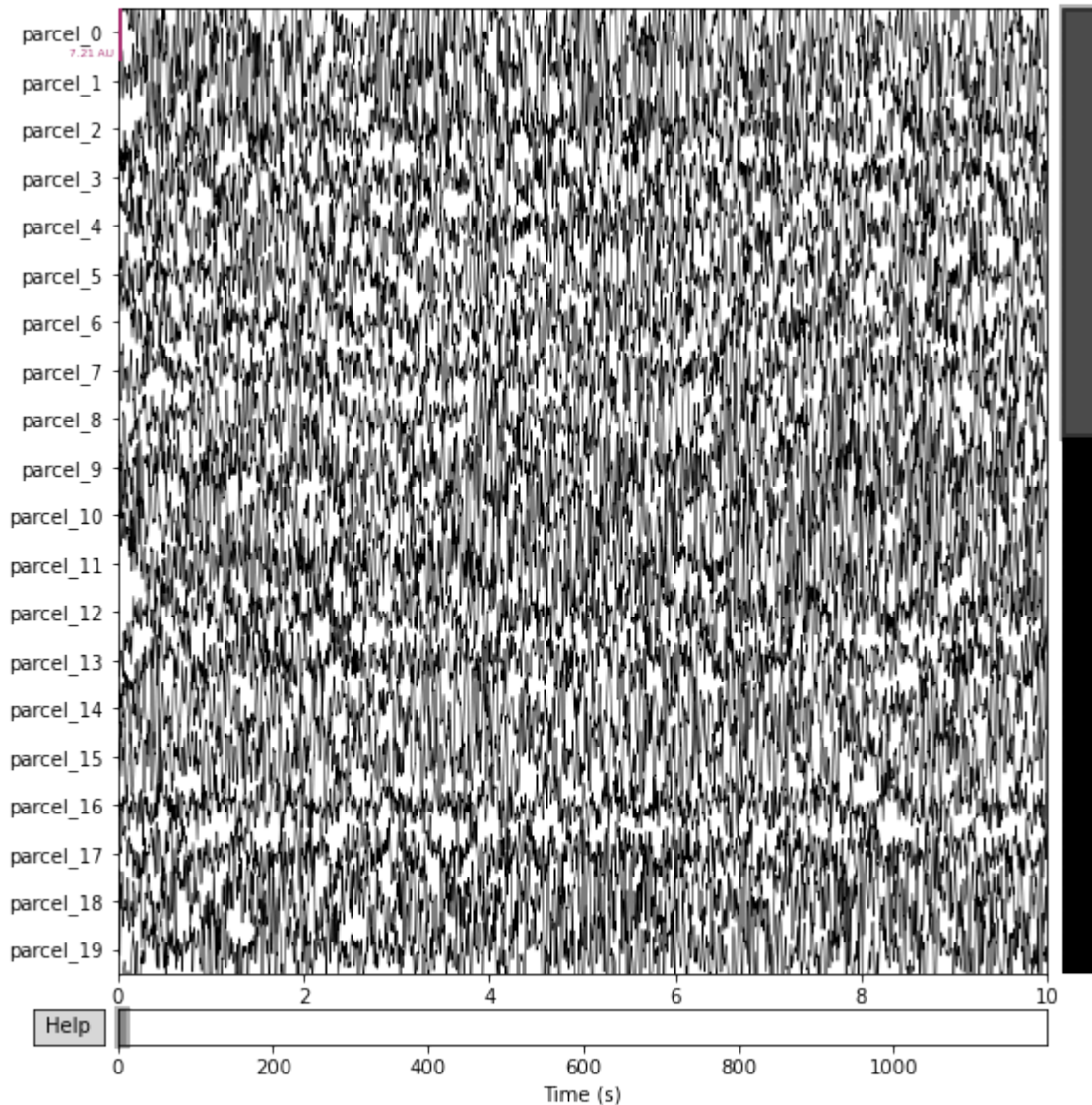
```
In [47]: raw = mne.io.read_raw_fif("/home/olivierb/coreg/meg_0925_SDMT_tsss_mc_preproc")
```

```
In [48]: raw.plot()
```





Out[48]:



## Processing Pipeline - different subject groups in Parallel processing

Here the entire pipeline is ran for a larger amount of subjects at once. The goal is to process all subject files. I avoid processing all subjects in a single run, in case that an unexpected error occurs.

Several CPU cores can be allocated at the same time to perform the processing of several subjects in parallel (faster).

Select subject group to perform analysis on:

```
In [15]: from dask.distributed import Client

## To check for correct .fif/.nii pairs
#for i in range(len(fif_files)):
    #print(i, fif_files[i][-21:-17], T1_MNI_niifiles[i][-22:-18])

## 1. index 0-59
#idx_start = 18
#idx_end = 60
#input_fif_files = fif_files[idx_start:idx_end]
```



```
## 2. index 60-115
#idx_start = 60
#idx_end = len(fif_files)
#input_fif_files = fif_files[idx_start:idx_end]

## 3. index 90-115
idx_start = 90
idx_end = len(fif_files)
input_fif_files = fif_files[idx_start:idx_end]
```

## Step 1: Preprocessing

```
In [ ]: from osl import preprocessing, utils

# Directory in which we save the preprocessed data
PREPROC_DIR = "/home/olivierb/processed_data"
inputs = input_fif_files

## Order of preprocessing steps actually matters in the list of dictionaries

# Settings
config = """
    preproc:
    - resample: {sfreq: 250}

    - filter: {l_freq: 0.01, h_freq: 70, method: iir, iir_params: {order: 5,
    - notch_filter: {freqs: 50, notch_widths: 2}

    - bad_segments: {segment_len: 500, picks: mag, significance_level: 0.1}
    - bad_segments: {segment_len: 500, picks: grad, significance_level: 0.1}
    - bad_segments: {segment_len: 500, picks: mag, mode: diff, significance_
    - bad_segments: {segment_len: 500, picks: grad, mode: diff, significance
    - bad_channels: {picks: mag, significance_level: 0.1}
    - bad_channels: {picks: grad, significance_level: 0.1}

    - ica_raw: {picks: meg, n_components: 64}
    - ica_autoreject: {apply: true}

    - interpolate_bads: {}
    """
# Setup parallel processing
client = Client(n_workers=4, threads_per_worker=1)

# Main preprocessing
preprocessing.run_proc_batch(
    config,
    inputs,
    outdir=PREPROC_DIR,
    overwrite=True,
    dask_client = True
)
```

```
In [18]: # Prepare subjects and subject names for the remaining steps in the pipeline

# Get the group of preprocessed files
DIR_processed = "/home/olivierb/processed_data/"
all_processed = sorted(glob.glob(DIR_processed + 'meg*_SDMT_tsss_mc/'))

# get the actual fif files for the current group of subjects
group_processed_fif = []
for i in range(idx_end - idx_start):
```

```

group_processed_fif.append(sorted(glob.glob(all_processed[idx_start + i]

# Get the names of the subjects of each preprocessed file (the last of part
group_processed_names = []
for i in range(idx_end - idx_start):
    group_processed_names.append(os.path.basename(group_processed_fif[i]))

# Get the group of T1_MNI.nii files
group_smri = T1_MNI_niifiles[idx_start:idx_end]
print(len(group_smri))

```

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## Step 2: Coregistration

```

In [ ]: from osl import source_recon, utils
import numpy as np

# Settings
config = """
    source_recon:
    - extract_fiducials_from_fif: {}      # ok
    - fix_headshape_points: {}          # ok
    - compute_surfaces:
        include_nose: False

    - coregister:
        use_nose: False
        use_headshape: True
        n_init: 50
"""

# Directories

# Directory in which we will save the coregistered data
COREG_DIR = "/home/olivierb/coreg"
# Directory to the FSL library, same as the one used in MATLAB, supported by
FSL_DIR = "/usr/lib/fsl/6.0"

# Setup parallel processing
client = Client(n_workers=4, threads_per_worker=1)

# Run coregistration
source_recon.setup_fsl(FSL_DIR)
source_recon.run_src_batch(
    config,
    src_dir=COREG_DIR,
    subjects=group_processed_names,      # name of the subject (pr
    preproc_files=group_processed_fif,   # path to preprocessed fi
    smri_files=group_smri,               # path to .nii file of th
    dask_client=True
)

```

## Step 3: Source reconstruction (Forward Modelling/Beamforming/Parcellation/Orthogonalisation)

```

In [ ]: from osl import source_recon
#'/home/olivierb/38_parcel.nii' => 38 parcels / sform_code=1

# Settings
config = """

```

```

source_recon:
- forward_model:
    model: Single Layer                                # ok: Single
- beamform_and_parcellate:
    freq_range: [1, 80]
    chantypes: [mag, grad]                            # standard f
    rank: {meg: 60}
    parcellation_file: '/home/olivierb/38_parcel.nii'
    method: spatial_basis                             # ok => spat
    orthogonalisation: None                           # ok (if you

"""

# Run beamforming and parcellation
FSL_DIR = "/usr/lib/fsl/6.0"
COREG_DIR = "/home/olivierb/coreg"

# Setup parallel processing
client = Client(n_workers=4, threads_per_worker=1)

## !!!!!!!!!!!!!!! Parallel processing fails for source reconstruction for some

source_recon.setup_fsl(FSL_DIR)
source_recon.run_src_batch(
    config,
    src_dir=COREG_DIR,                                # path to all the
    subjects=group_processed_names,                   # name of the sub
    preproc_files=group_processed_fif,                 # path to preproc
    #dask_client=True
)

```

**Before we can do the Sign-Flipping (Step 4):**

**1 subject (ID 0995) is missing ECG/EOG channels, the automated ICA artifact rejection can not be executed**

For the single subject that did not have EOG/ECG channels

```

In [ ]: from osl import preprocessing, utils

# Directory in which we save the preprocessed data
PREPROC_DIR = "/home/olivierb/processed_data"

## Order of preprocessing steps actually matters in the list of dictionaries

# Settings
config = """
preproc:
- resample: {sfreq: 250}

- filter: {l_freq: 0.01, h_freq: 70, method: iir, iir_params: {order: 5,
- notch_filter: {freqs: 50, notch_widths: 2}

- bad_segments: {segment_len: 500, picks: mag, significance_level: 0.1}
- bad_segments: {segment_len: 500, picks: grad, significance_level: 0.1}
- bad_segments: {segment_len: 500, picks: mag, mode: diff, significance_
- bad_segments: {segment_len: 500, picks: grad, mode: diff, significance
- bad_channels: {picks: mag, significance_level: 0.1}
- bad_channels: {picks: grad, significance_level: 0.1}

- interpolate_bads: {}

```

```

# NO ICA FOR SUBJECT 0995 => NO EOG CHANNELS FOUND
#- ica_raw: {picks: meg, n_components: 64}
#- ica_autoreject: {apply: true}

"""

# Main preprocessing
preprocessing.run_proc_batch(
    config,
    fif_files[7],
    outdir=PREPROC_DIR,
    overwrite=True,
)

```

## Manual ICA - subject 0995

be careful! ICA should be performed on the .fif file after filtering/referencing etc. (Don't accidentally select the file with the already applied manual ICA on it)

```

In [9]: # Load the single subject
raw = mne.io.read_raw_fif('/home/olivierb/processed_data/meg_0995_SDMT_tsss_

```

```

In [10]: ica = mne.preprocessing.ICA(n_components=64, random_state = 0)
raw_ica = raw.copy()
ica.fit(raw_ica)

```

```

Out[10]:

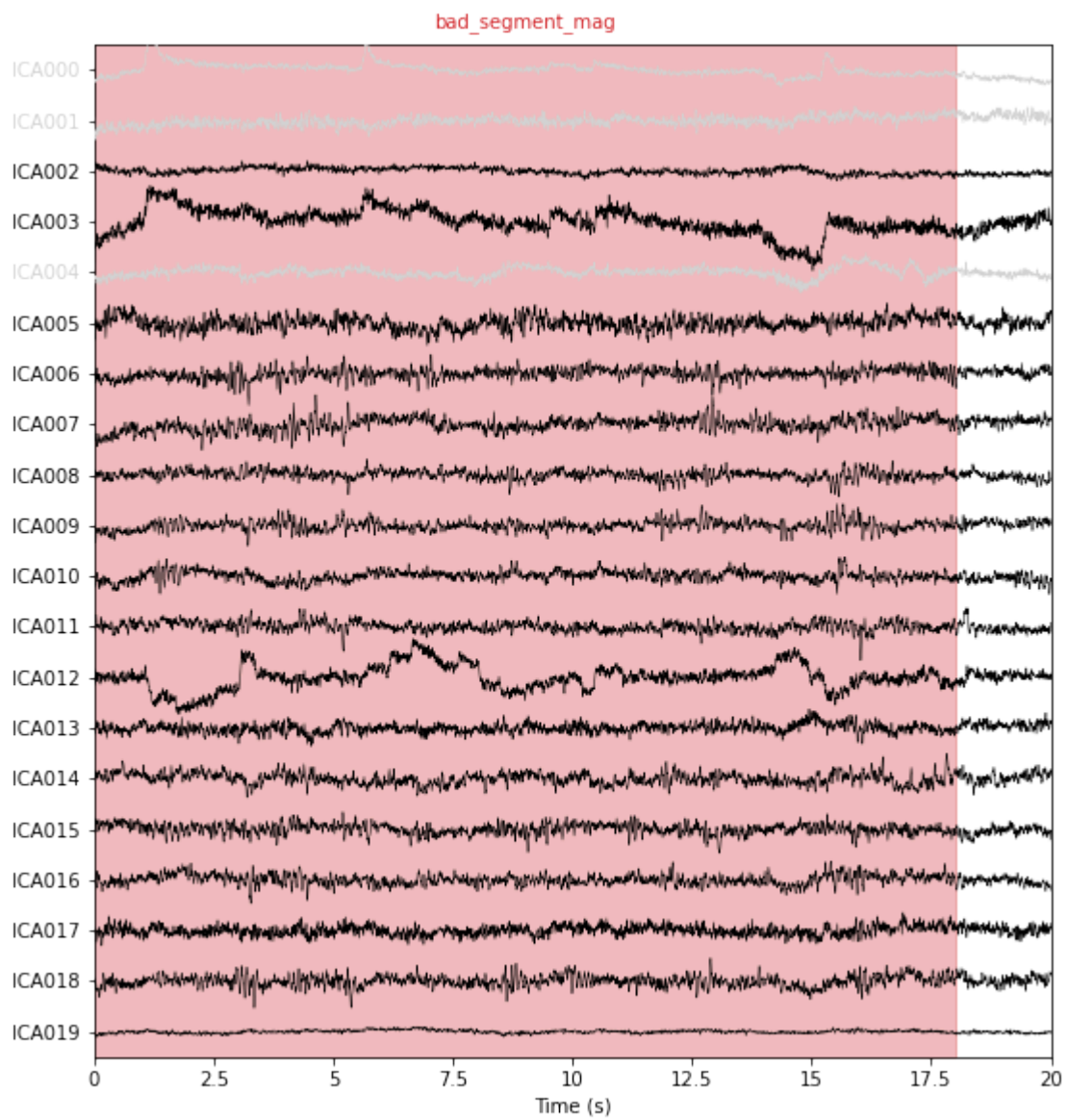
```

	Method	fastica
	Fit	76 iterations on raw data (239001 samples)
	ICA components	64
	Available PCA components	306
	Channel types	mag, grad
	ICA components marked for exclusion	—

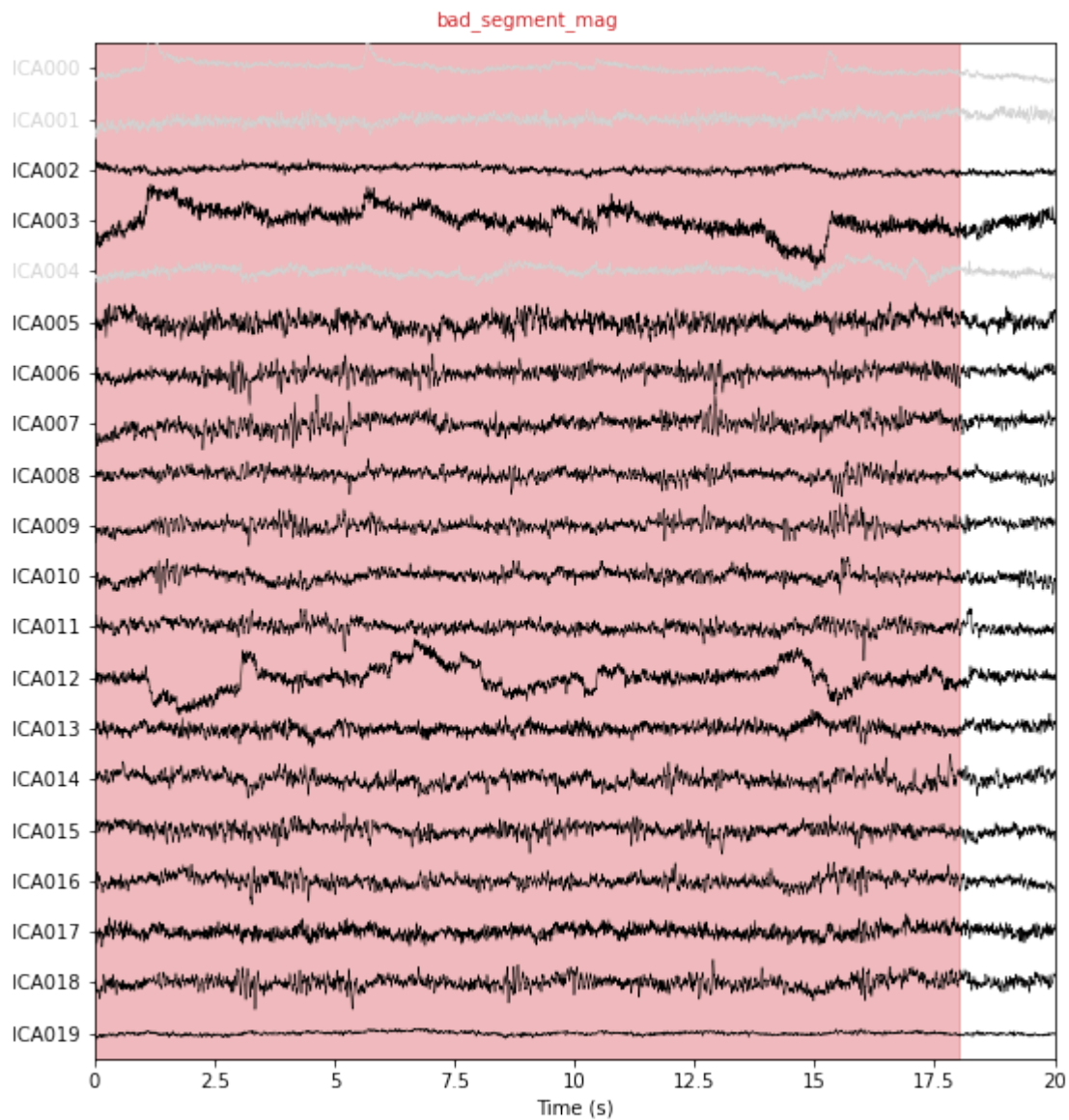
```

In [17]: ica.plot_sources(raw_ica, show_scrollbars=False)

```



Out[17]:

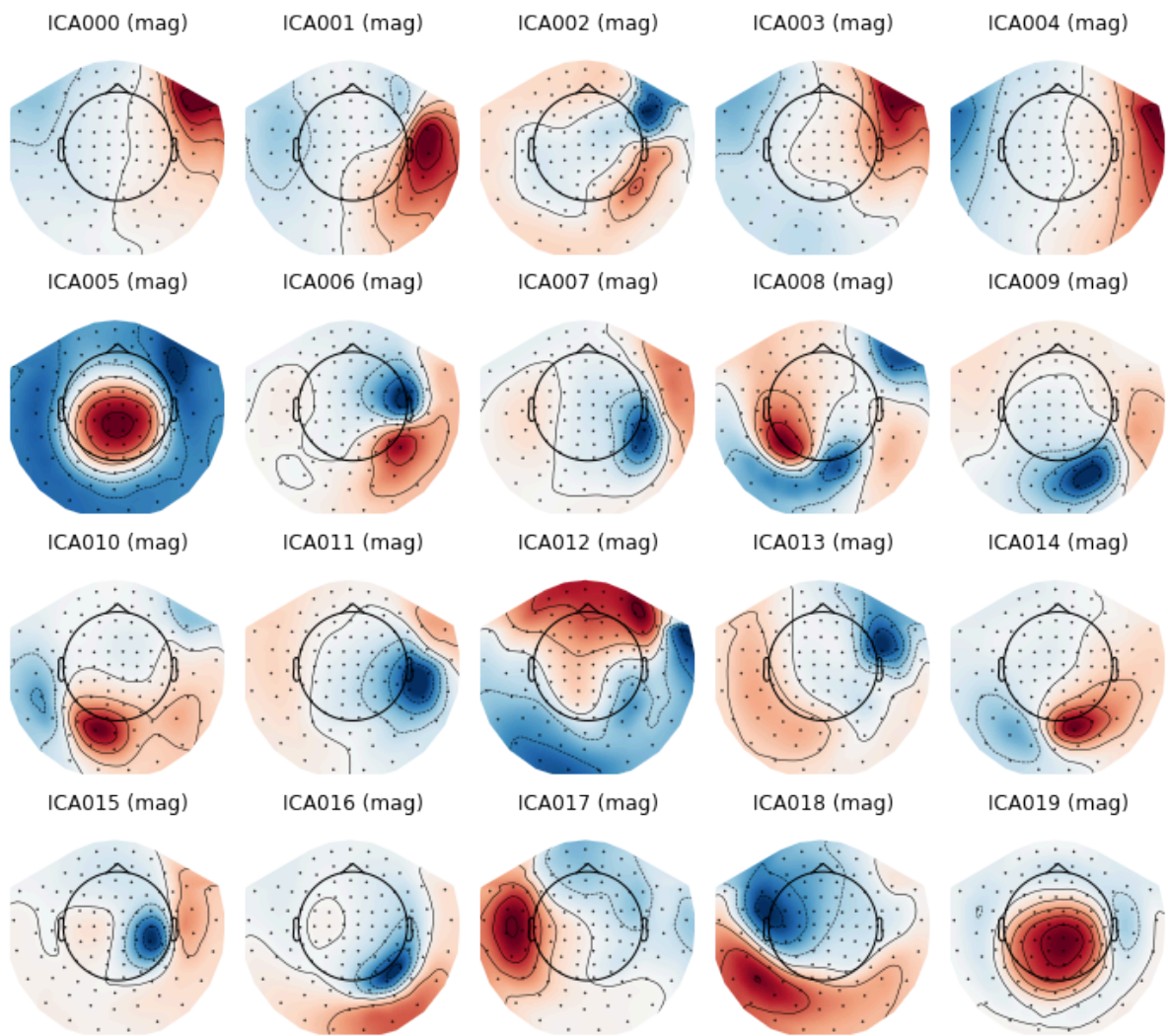


```
In [15]: %matplotlib inline  
ica.plot_components()
```

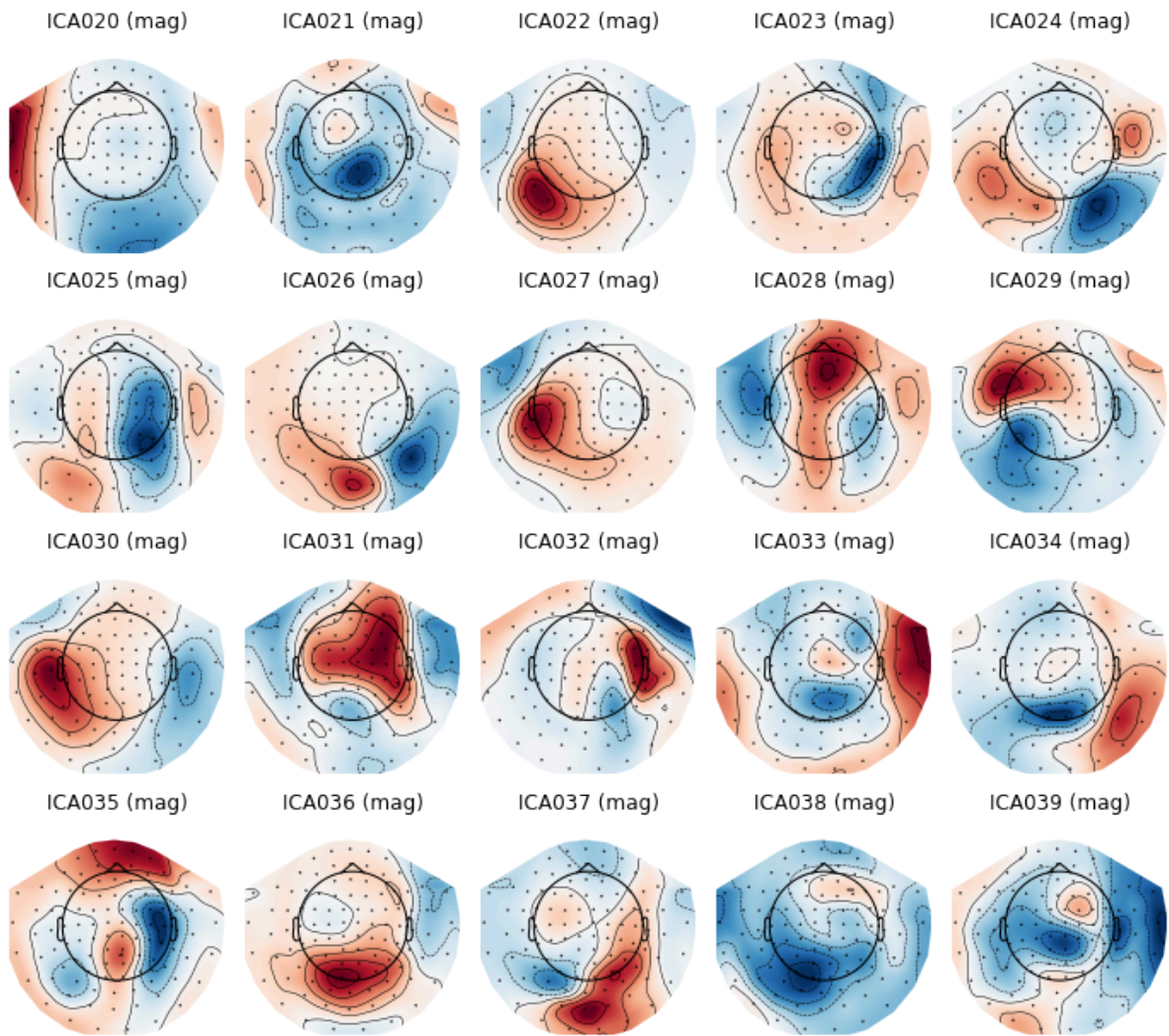
```
/tmp/ipykernel_945248/150138342.py:2: RuntimeWarning: (X, Y) fit (3.1, 2  
0.1) more than 20 mm from head frame origin  
ica.plot_components()
```



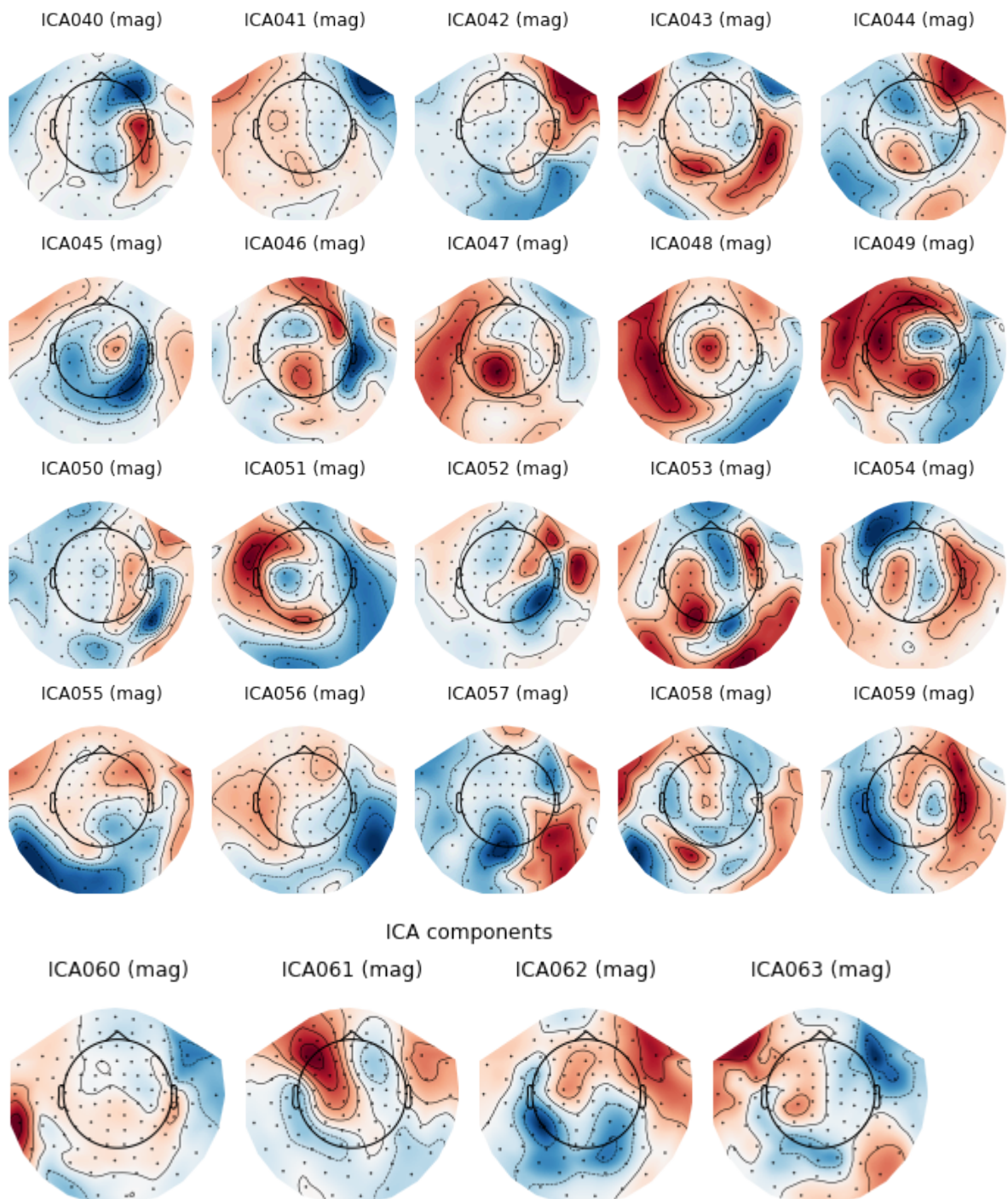
## ICA components



## ICA components



## ICA components

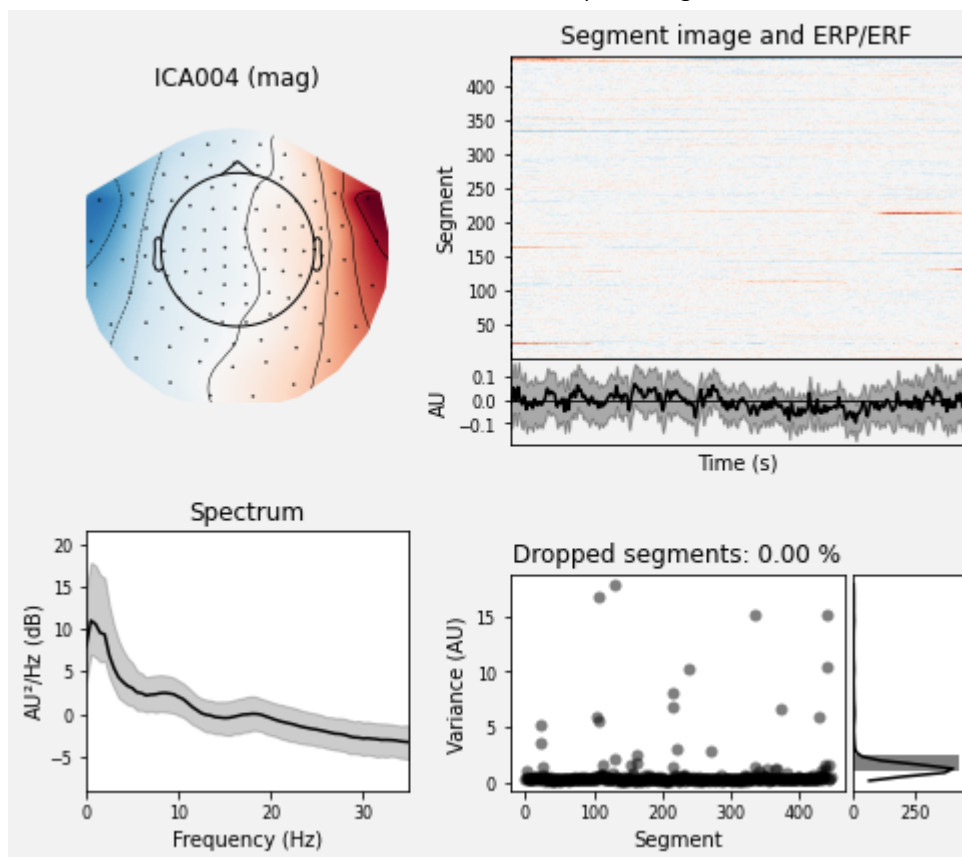


```
Out[15]: [<MNEFigure size 702x696.24 with 20 Axes>,
<MNEFigure size 702x696.24 with 20 Axes>,
<MNEFigure size 702x696.24 with 20 Axes>,
<MNEFigure size 561.6x187.56 with 4 Axes>]
```

```
In [45]: %matplotlib inline
ica.plot_properties(raw, picks=4, psd_args={'fmax': 35.})
```

```
/tmp/ipykernel_945248/2658260692.py:2: RuntimeWarning: (X, Y) fit (3.1, 2
0.1) more than 20 mm from head frame origin
ica.plot_properties(raw, picks=4, psd_args={'fmax': 35.})
```





Out[45]: [**Figure** size 504x432 with 6 Axes]

```
In [46]: #ICA000, ICA001, ICA003, ICA004
         artifact_picks = 0,1,3,4

         #Removing the ICA components containing eyeblinks and eyemovements
         ica.exclude = artifact_picks # indices chosen based on various plots above

         reconst_raw_1 = raw.copy()
         ica.apply(reconst_raw_1)
```

Out[46]: **Measurement date** August 26, 2015 11:01:12 GMT

<b>Experimenter</b>	neuromag
<b>Participant</b>	
<b>Digitized points</b>	299 points
<b>Good channels</b>	204 Gradiometers, 102 Magnetometers, 3 Stimulus, 12 misc, 9 CHPI
<b>Bad channels</b>	None
<b>EOG channels</b>	Not available
<b>ECG channels</b>	Not available
<b>Sampling frequency</b>	250.00 Hz
<b>Highpass</b>	0.10 Hz
<b>Lowpass</b>	70.00 Hz
<b>Filenames</b>	meg_0995_SDMT_tsss_mc_preproc_raw.fif
<b>Duration</b>	00:18:18 (HH:MM:SS)

```
In [47]: reconst_raw_1.save('/home/olivierb/processed_data/meg_0995_SDMT_tsss_mc/meg_
```

```
In [5]: from osl import source_recon, utils

# Settings
config = """
    source_recon:
    - extract_fiducials_from_fif: {}      # ok
    - compute_surfaces:
        include_nose: False

    - coregister:
        use_nose: False
        use_headshape: True
        n_init: 50
"""

# Directories

subject = "/home/olivierb/test/meg_0995_SDMT_tsss_mc/meg_0995_SDMT_tsss_mc_p

# Directory in which we will save the coregistered data
COREG_DIR = "/home/olivierb/coreg"

# Directory to the FSL library, same as the one used in MATLAB, supported by
FSL_DIR = "/usr/lib/fsl/6.0"

# Run coregistration

source_recon.setup_fsl(FSL_DIR)
source_recon.run_src_batch(
    config,
    src_dir=COREG_DIR,
    subjects=['meg_0995_SDMT_tsss_mc_preproc_raw.fif'], # name of the subj
    preproc_files=[subject], # path to preproc
    smri_files=[T1_MNI_niifiles[7]], # path to .nii fi
)
```

```
In [ ]: from osl import source_recon
# '/home/olivierb/38_parcel.nii' => 38 parcels / sform_code=1

# Settings
config = """
    source_recon:
    - forward_model:
        model: Single Layer # ok: Single Shell in MA
    - beamform_and_parcellate:
        freq_range: [1, 80]
        chantypes: [mag, grad] # standard for MEG
        rank: {meg: 60}
        parcellation_file: '/home/olivierb/38_parcel.nii'
        method: spatial_basis # ok => spatial_basis or
        orthogonalisation: symmetric # ok
"""

# Run beamforming and parcellation
FSL_DIR = "/usr/lib/fsl/6.0"
COREG_DIR = "/home/olivierb/coreg"
SUBJECT = subject = "/home/olivierb/processed_data/meg_0995_SDMT_tsss_mc/meg

source_recon.setup_fsl(FSL_DIR)
source_recon.run_src_batch(
    config,
    src_dir=COREG_DIR, # path to all the
```

```

subjects=['meg_0995_SDMT_tsss_mc_preproc_raw.fif'],      # name of the sub
preproc_files=[subject]                                  # path to preproc
)

```

## Step 4: Sign-flipping (all files together)

```

In [ ]: # To fix dipole sign ambiguity.

from dask.distributed import Client
from osl import utils
from osl.source_recon import find_template_subject, run_src_batch, setup_fsl

# Directories
SRC_DIR = "/home/olivierb/coreg/"
FSL_DIR = "/usr/lib/fsl/6.0"

# Get the parceled data
DIR_parceled = "/home/olivierb/coreg/"
all_parceled = sorted(glob.glob(DIR_parceled + 'meg*_SDMT_tsss_mc_preproc_*.fif'))
subjects = all_parceled

setup_fsl(FSL_DIR)

# Find a good template subject to align other subjects to
template = find_template_subject(SRC_DIR, subjects, n_embeddings=15, standardize=True)

```

Get the subject names for all processed files

```

In [ ]: all_successful_parceled = sorted(glob.glob('/home/olivierb/coreg/' + 'meg*_SDMT_tsss_mc_preproc_*.fif'))

names_subjects = []

# Gets you all the subject names for which the data was successfully parceled
for file_path in all_successful_parceled:
    # Split the string by '/'
    parts = file_path.split('/')
    # Extract the desired part (second-to-last element)
    desired_part = parts[-3]
    names_subjects.append(desired_part)

```

```

In [ ]: # Settings for batch processing
config = f"""
    source_recon:
    - fix_sign_ambiguity:
        template: {template}
        n_embeddings: 15
        standardize: True
        n_init: 5
        n_iter: 500
        max_flips: 20
"""

# Setup parallel processing
DIR_parceled = "/home/olivierb/coreg/"

# Do the sign flipping - Apply the template
run_src_batch(config, SRC_DIR, names_subjects) #dask_client=True)

```

**Extra: Function to overwrite sform\_code of a .nii file (can be used on both MRI scans & parcellation files)**



This is required due to a mismatch in `sform_code` for the **parcellation file** (gives an error in osl python):

- Original `sform_code` = 4 (normalized coordinates)
- Required `sform_code` = 1 (scanner)based anatomical coordinates)

Meaning of the `sform` codes: [https://nifti.nimh.nih.gov/nifti-1/documentation/nifti1fields/nifti1fields\\_pages/qsform.html/document\\_view](https://nifti.nimh.nih.gov/nifti-1/documentation/nifti1fields/nifti1fields_pages/qsform.html/document_view)

```
In [20]: import nibabel as nib
import numpy as np
import pandas as pd

## Parcellation files
#(original_file_path = "/home/olivierb/osl/source_recon/parcellation/files/1

# Function to overwrite the sform_code of an existing .nii file
def overwrite_sformcode(original_file_path, output_file_path):

    # Load the NIfTI image
    nii_img = nib.load(original_file_path)

    # Get the image data and affine matrix
    data = nii_img.get_fdata()
    affine = nii_img.affine

    # Create a new NIfTI image with the modified sform_code
    new_sformcode = 1 # Set to the desired value
    new_header = nii_img.header.copy()
    new_header.set_sform(affine, code=new_sformcode)

    # Create a new image using the modified data and header
    nii_img_with_modified_header = nib.Nifti1Image(data, affine, header=new_

    # Save the new image to a file
    nib.save(nii_img_with_modified_header, output_file_path)

    # Reload the saved image
    loaded_img = nib.load(output_file_path)

    # Check the sform_code of the loaded image
    print(loaded_img.header['sform_code'])

    return
```

## Extra: Explore and visualise parcellation file

Plot parcellation for visualisation

```
In [50]: from nilearn import image

nii_img = image.load_img("/home/fahimeha/PhD/preprocessing/parcellation/fmri

print(nii_img.shape)

one_volume = image.index_img(nii_img, 3)
print(one_volume.shape)
```

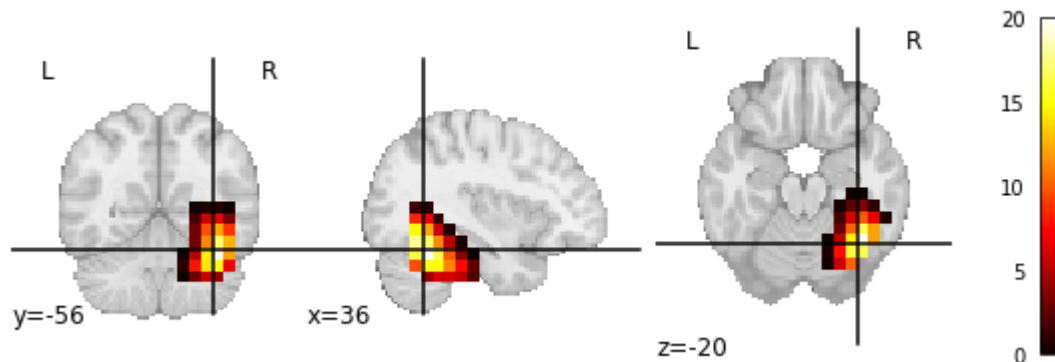
```
(23, 27, 23, 42)
(23, 27, 23)
```

```
In [51]: from nilearn import image
path_parcel = "/home/olivierb/osl/source_recon/parcellation/files/fmri_d100_
nii_img = image.load_img(path_parcel)
print(nii_img.shape)
```

```
(23, 27, 23, 38)
```

```
In [52]: from nilearn import plotting
plotting.plot_stat_map(one_volume)
```

```
Out[52]: <nilearn.plotting.displays._slicers.OrthoSlicer at 0x7f6ed5594640>
```



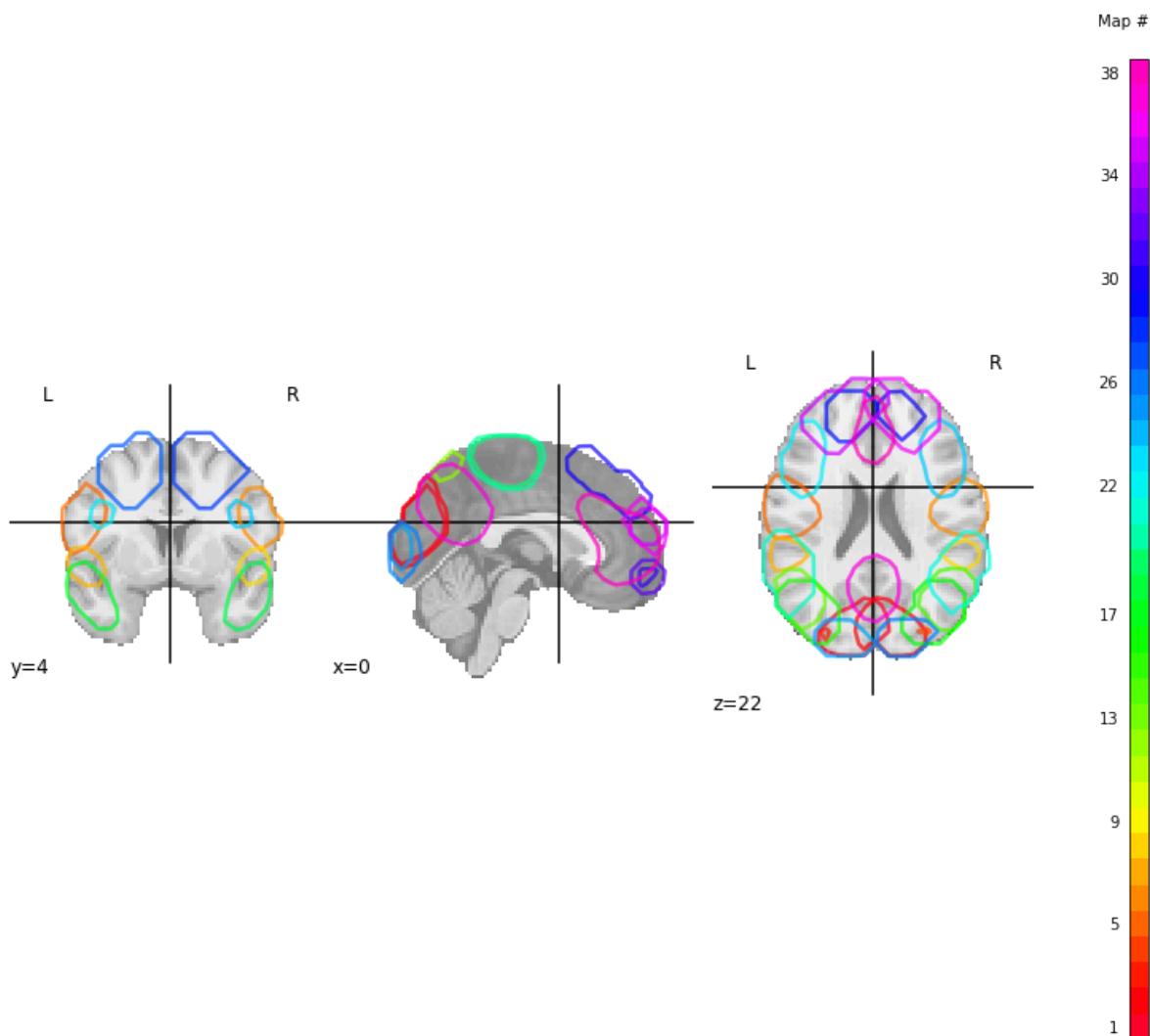
```
In [21]: import nilearn
```

```
In [22]: from nilearn.regions import connected_regions
from nilearn import datasets, plotting
import nibabel as nib
atlas = nib.load("/home/olivierb/38_parcel.nii")
```

```
In [23]: region_labels = connected_regions(atlas)
```

```
In [24]: import matplotlib.pyplot as plt
nilearn.plotting.plot_prob_atlas(atlas, figure=plt.figure(figsize=(10,10)),
```

```
Out[24]: <nilearn.plotting.displays._slicers.OrthoSlicer at 0x7f695e0115e0>
```



Extract parcel names for the parcellation file

```
In [25]: ## Parcel names extract

# Open the .txt file for reading
file_path = "/home/olivierb/Downloads/Parcels_38_names.txt"
with open(file_path, 'r') as file:
    lines = file.readlines()

# Strip any leading/trailing whitespace and create a list of strings
parcel_names = [line.strip() for line in lines]

# Print the list of strings
print(parcel_names)

# Parcel indices (how the channel names are stored in the Python parceled .i
parcel_idx = []
for i in range(len(parcel_names)):      # 38 parcels
    parcel_idx.append('parcel_' + str(i))
print(parcel_idx)
```

```
['L Cuneus', 'R Cuneus', 'L Inf Occ', 'R Inf Occ', 'L Supramarginal', 'R
Supramarginal', 'L Sup Temp', 'R Sup Temp', 'L Lat SMC', 'R Lat SMC', 'L
Sup Parietal', 'R Sup Parietal', 'L Middle Occ', 'R Middle Occ', 'L Sup O
cc', 'R Sup Occ', 'L Ant Temp', 'R Ant Temp', 'L Medial SMC', 'R Medial S
MC', 'L Angular', 'R Angular', 'L VL PFC', 'R VL PFC', 'L Occ pole', 'R O
cc pole', 'L Sup PFC', 'R Sup PFC', 'L Sup Dorsal PFC', 'R Sup Dorsal PF
C', 'L Orbitofrontal', 'R Orbitofrontal', 'L Post Temp', 'R Post Temp',
'L Inf Dorsal PFC', 'R Inf Dorsal PFC', 'Medial PFC', 'Posterior Cingulat
e Cortex']
['parcel_0', 'parcel_1', 'parcel_2', 'parcel_3', 'parcel_4', 'parcel_5',
'parcel_6', 'parcel_7', 'parcel_8', 'parcel_9', 'parcel_10', 'parcel_11',
'parcel_12', 'parcel_13', 'parcel_14', 'parcel_15', 'parcel_16', 'parcel
_17', 'parcel_18', 'parcel_19', 'parcel_20', 'parcel_21', 'parcel_22', 'pa
rcel_23', 'parcel_24', 'parcel_25', 'parcel_26', 'parcel_27', 'parcel_2
8', 'parcel_29', 'parcel_30', 'parcel_31', 'parcel_32', 'parcel_33', 'pa
rcel_34', 'parcel_35', 'parcel_36', 'parcel_37']
```

```
In [26]: from nilearn.image import iter_img
from nilearn.plotting import plot_stat_map, show
```

```
for i, cur_img in enumerate(iter_img(atlas)):
    plot_stat_map(
        cur_img,
        title=f"{parcel_idx[i]} - {parcel_names[i]}",
        colorbar=False,
    )
```

*# takes a little while to plot everything*

/home/olivierb/.local/lib/python3.9/site-packages/nilearn/plotting/displays/\_slicers.py:144: RuntimeWarning: More than 20 figures have been opened. Figures created through the pyplot interface (`matplotlib.pyplot.figure`) are retained until explicitly closed and may consume too much memory. (To control this warning, see the rcParam `figure.max\_open\_warning`). Consider using `matplotlib.pyplot.close()`.

```
figure = plt.figure(figure, figsize=figsize,
```

