

DESIGN LAB

Documentation

To Visualize the EEG Microstates

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Documentation for the Microstates

In this documentation we are using the microstates module to "Segment a continuous signal into microstates" and then we have implemented and visualized the microstates in topographical maps. Also, the code plots "auto scaled data" just to visualize the raw data that was selected by the user. The preferred file format is ".mul" and the preferred channel location file can have the following formats:

```
'.elc','.txt','.loc','.locs','csd','elp','hpts',' sfp','eloc' or '.bvref'.
```

Here we have used the concept of *Global field power* [2]. First the user is prompted to give the desired data file and channel location file of the EEG data. Then using the microstates module, the segment method can be called to find the Peaks in the global field power (GFP). These are used to find microstates, using a modified K-means algorithm. Several runs of the modified K-means algorithm are performed, using different random initializations. The run that resulted in the best segmentation, as measured by global explained variance (GEV), is used. The following parameters are given to call the method(function) from the microstate module.

Method (function) Parameters:

data: ndarray, shape (n_channels, n_samples)

The data for finding the EEG microstates where the data has 63 channels(by default) and 30000 samples

n states:int

The number of unique microstates to find. Defaults to 4.

n_inits:int

The number of random initializations to use for the k-means algorithm. The best fitting segmentation across all initializations is used. Defaults to 10 but can be also 30/40/50.

max_iter : int

The maximum number of iterations to perform in the k-means algorithm. Defaults to 1000.

thresh: float

The threshold of convergence for the k-means algorithm, based on relative change in noise variance. Defaults to 1e-6.

normalize: bool

Whether to normalize the data across time before running the k-means algorithm. Defaults to ``False``.

min_peak_dist:int

Minimum distance (in samples) between peaks in the GFP. Defaults to 2.

```
max_n_peaks : int
```

Maximum number of GFP peaks to use in the k-means algorithm. Chosen randomly. Defaults to 10000.

```
random_state : int | numpy.random.RandomState | None
```

The seed or ``RandomState`` for the random number generator. Defaults to ``None``, in which case a different seed is chosen each time this function is called.

verbose: input type: int(integer) | bool | None => This controls the verbosity.

The method(function) returns the followings:

Returns:

```
maps : ndarray, shape (n_channels, n_states)
```

The topographic maps of the found unique microstates.

```
segmentation: ndarray, shape (n_samples,)
```

For each sample, the index of the microstate to which the sample has

been assigned.

Next we want to plot the topographical map of the microstates. For this we called the *plot_maps* method(function) from the microstate module.

"""Plotting prototypical microstate maps.

Parameters:

```
maps: ndarray, shape (n_channels, n_maps)
```

The prototypical microstate maps.

info: instance of mne.io.Info. The info structure of the dataset, containing the location of the sensors.

.....

Lastly the module can be used to plot the microstate segmentation for the data.

[&]quot;""Plot a microstate segmentation.

Parameters:

segmentation: list of int (integer)

For each sample in time, the index of the state to which the sample has been assigned.

times: list of float => The time-stamp for each sample.

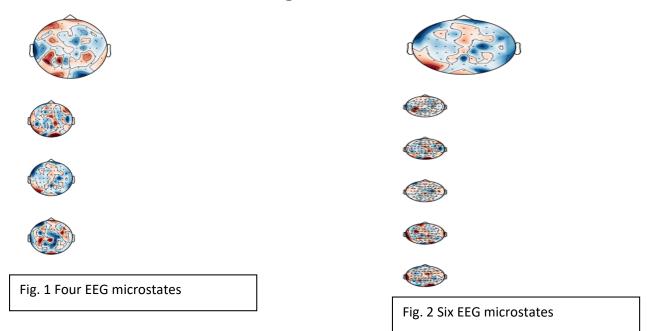
References:

[1] Pascual-Marqui, R. D., Michel, C. M., & Lehmann, D. (1995). Segmentation of brain electrical activity into microstates: model estimation and validation. IEEE Transactions on Biomedical Engineering.

[2] Global Field Power is a related, parametric assessment of map strength, defined as the sum of the absolute microvolt values measured at all electrodes divided by the number of electrodes; the assessment must be done after the values in each map have been expressed as deviations from the mean of all momentary values (spatial DC offset removal, 'average reference') computed as standard deviation of the momentary potential values (Lehmann and Skrandies, 1980).

http://www.scholarpedia.org/article/EEG_microstates#Brain_electric_fields

Visualizing EEG Microstates:



Topographic maps of EEG microstates in a continuous EEG signal. This depends on the number of microstates as given by the user.

Code:

The coding is done with Python 3.7. We have used the *ANACONDA* package for the python version 3.7. However, one can use the following packages, libraries and modules to implement the code. The setup file is given as "setup.py" for the easy installation of *certain* libraries

Required packages, Libraries and modules:

We adopted the following modules, packages and libraries:

- 1. Numpy
- 2. MNE python
- 3. Matplotlib
- 4. Pathlib library and Path module
- 5. Microstates
- 6. SciPy for scientific computing and technical computing
- 7. Tkinter package for python 3.7
- 8. Operation Sytem (Os) module

Python 3.7 CODE:

MAIN:

File name: "MAIN.py"

```
import tkinter as tk
                                            lines = f.readlines()
from tkinter import filedialog
                                            matrix = []
                                            for line in lines:
import numpy as np
import mne
                                                 res=[]
import microstates
                                                 temp
                                    line.split(' ')
import os
                                                 for num in temp:
from pathlib import Path
                                                     if num:
#Function to read the MUL data
file
                                    res.append(float(num))
def read data(path):
                                                matrix.append(res)
    with open (path, 'r') as f:
```

#Loading of the data: return Prefereable format .MUL np.asarray(matrix) data1 read data(data file path) #Tkinter package for user selection of files: DATA and data Channel Location file np.resize(data1, (63, 30000)) root = tk.Tk()root.withdraw() #The name of the channels. It can be modified as desired. By default 63 print("Please select the data ch names = ['FP1', 'Fz', 'F3',file") 'F7', 'FT9', 'FC5', 'FC1', 'C3','T7','TP9', data file path filedialog.askopenfilename() 'CP5', 'CP1', 'Pz', 'P3', 'P7', 'O1 print("Please select the ','Oz','O2','P4','P8','TP10',' channel location file") CP6', channel file path filedialog.askopenfilename() 'CP2','C4','T8','FT10','FC6',' FC2', 'F4', 'F8', 'FP2', 'AF7', 'AF 3', # "kind" and "path" variables for mne.channels read montage function 'AFz','F1','F5','FT7','FC3','F Cz','C1','C5','TP7','CP3','P1' p = Path(channel file path) ,'P5', k = p.parts[-1]d = k.find('.')'PO7', 'PO3', 'POz', 'PO4', 'PO8', 'P6', 'P2', 'CPz', 'CP4', 'TP8', 'C kind = k[0:d]6', f = p.parts'C2', 'FC4', 'FT8', 'F6', 'F2', 'AF 4','AF8'] f=f[0:len(f)-1]path = os.path.join(*f) #In the unit parameter "cm"/"m" can be given

#EEG Microstates montage mne.channels.read montage(kind # Segment the data in number of ='Cap63', ch names = ch names, microstates n states = int(input("Please path path, unit='cm', provide the number of transform=False) Microstates: ")) #creating the channel info if n states <2: instance print("The number info microstates must be equal mne.create info(ch names greater than or equal to 2") sfreq=250,ch types ch names, ='eeg', montage n inits = int(input("Please = montage, verbose = None) give the number of random initializations to use for the #Creating the raw instance of k-means algorithm: ")) the data mne.io.RawArray(data,info,firs maps, segmentation t samp= 0, verbose = None) microstates.segment(raw.get da n states= n states, ta(), n inits = n inits) #""" OPTIONAL PARTS Raw data # Plot the topographic maps of visualization """ the microstates and the segmentation #Auto scaling option scalings ='auto' print(" Visualizing the topographical maps of the EEG raw.plot(n channels Micrsotates ") scalings=scalings, title='Autoscaled Data from arrays', show=True,block=True) microstates.plot maps (maps, raw.info) #Plotting the segementation for first 600 time samples microstates.plot segmentation(segmentation[:600], raw.get data()[:, :600],raw.times[:600])

Microstates Module Code:

The required code for Microstates module: "microstates.py"

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Functions to segment EEG into microstates. Based on the Microsegment toolbox

for EEGlab, written by Andreas
Trier Poulsen [1] .

Reference:

Author: Marijn van Vliet <w.m.vanvliet@gmail.com>

References

.. [1] Poulsen, A. T., Pedroni, A., Langer, N., & Hansen, L. K. (2018).

Microstate EEGlab toolbox: An introductionary guide. bioRxiv.

** ** **

import warnings

import numpy as np

from scipy.stats import zscore

from scipy.signal import
find peaks

from scipy.linalg import eigh

import matplotlib as mpl

from matplotlib import pyplot as
plt

import mne

from mne.utils import logger,
verbose

@verbose

def segment(data, n_states=4,
n_inits=10, max_iter=1000,
thresh=1e-6,

normalize=False,
min_peak_dist=2,
max n peaks=10000,

random_state=None,
verbose=None):

"""Segment a continuous signal into microstates.

Peaks in the global field power (GFP) are used to find microstates, using a

 $\,$ modified K-means algorithm. Several runs of the modified K-means algorithm

are performed, using different random initializations. The run that

resulted in the best segmentation, as measured by global explained variance

(GEV), is used.

Parameters

data : ndarray, shape
(n channels, n samples)

The data to find the microstates in

n_states : int

The number of unique microstates to find. Defaults to $4. \ \ \,$

n inits : int

 $\begin{tabular}{ll} The number of random \\ initializations to use for the \\ k-means algorithm. \end{tabular}$

The best fitting segmentation across all initializations is used.

Defaults to 10.

max iter : int

The maximum number of iterations to perform in the k-means algorithm.

Defaults to 1000.

thresh : float

The threshold of convergence for the k-means algorithm, based on

relative change in noise variance. Defaults to 1e-6.

normalize : bool

Whether to normalize (z-score) the data across time before running the

 $$k{\mbox{-}means}$$ algorithm. Defaults to ''False''.

min_peak_dist : int

Minimum distance (in samples) between peaks in the GFP. Defaults to 2.

max n peaks : int

Maximum number of GFP peaks to use in the k-means algorithm. Chosen

randomly. Defaults to 10000.

random_state : int |
numpy.random.RandomState | None

The seed or ``RandomState`` for the random number generator. Defaults

to ``None``, in which case a different seed is chosen each time this

function is called.

verbose: int | bool | None
Controls the verbosity.

Returns

maps : ndarray, shape
(n_channels, n_states)

The topographic maps of the found unique microstates.

segmentation : ndarray,
shape (n samples,)

For each sample, the index of the microstate to which the sample has

been assigned.

References

.. [1] Pascual-Marqui, R. D., Michel, C. M., & Lehmann, D. (1995).

Segmentation of brain random state = electrical activity into np.random.RandomState(random sta microstates: model estimation and chosen peaks = validation. IEEE Transactions on random state.choice(n peaks, Biomedical size=max n peaks, Engineering. replace=False) 11 11 11 peaks = logger.info('Finding %d peaks[chosen peaks] microstates, using %d random intitializations for the k-means algorithm' % # Run microstates analysis (n states, on selected data n inits)) if normalize: data = zscore(data, # Convert min peak dist to axis=1) samples # min peak dist = 1 + # Cache this value for later int(round(min peak dist * raw.info['sfreq'])) gfp_sum_sq = np.sum(gfp ** 2) # Find peaks in the global field power (GFP) # Do several runs of the kmeans algorithm, keep track of gfp = data.std(axis=0) the best peaks, = find peaks(gfp, # segmentation. distance=min peak dist) best gev = 0n peaks = len(peaks) best maps = None best segmentation = None # Limit the number of peaks by randomly selecting them for in range(n inits): if max_n_peaks is not None: maps, segmentation = mod kmeans (data, n states, max n peaks = n inits, max iter, min(n peaks, max n peaks) if not thresh, random state, verbose) isinstance(random state,

np.random.RandomState):

if not map corr = corr vectors (data, isinstance(random state, maps[segmentation].T) np.random.RandomState): random state = np.random.RandomState(random sta # Compare across te) iterations using global explained variance (GEV) of n channels, n samples = data.shape # the found microstates. qev = sum((qfp *map_corr) ** 2) / gfp_sum_sq # Cache this value for later logger.info('GEV of data sum sq = np.sum(data ** found microstates: %f' % gev) 2) if gev > best gev: best gev, best maps, # Select random timepoints best segmentation = gev, maps, for our initial topographic maps segmentation init times = random state.choice(n samples, size=n states, replace=False) return best_maps, best segmentation maps = data[:, init times].T maps /= np.linalg.norm(maps, axis=1, keepdims=True) # Normalize the maps @verbose def mod kmeans(data, prev residual = np.inf n states=4, n inits=10, max iter=1000, thresh=1e-6, for iteration in range(max iter): random state=None, # Assign each sample to verbose=None): the best matching microstate """The modified K-means activation = clustering algorithm. maps.dot(data) segmentation = np.argmax(np.abs(activation), See :func:`segment` for the axis=0)meaning of the parameters and # assigned activations = return np.choose(segmentations, values.

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all activations)

Recompute the residual = topographic maps of the abs(data_sum_sq - act_sum_sq) microstates, based on the residual /= float(n samples * (n channels -# samples that were assigned to each state. for state in range(n states): # Have we converged? idx = (segmentation)if (prev residual -== state) residual) < (thresh * residual):</pre> if np.sum(idx) == 0: logger.info('Converged at %d iterations.' % iteration) warnings.warn('Some microstates are never activated') break maps[state] = 0continue prev residual = residual else: # Find largest warnings.warn('Modified eigenvector K-means algorithm failed to converge.') #cov = data[:, idx].dot(data[:, idx].T) # , vec = eigh(cov, # Compute final microstate eigvals=(n_channels - 1, segmentations n channels - 1)) activation = maps.dot(data) #maps[state] = vec.ravel() segmentation = np.argmax(activation ** 2, maps[state] = axis=0)data[:, idx].dot(activation[state, idx]) maps[state] /= return maps, segmentation np.linalg.norm(maps[state]) # Estimate residual def corr vectors(A, B, axis=0): noise """Compute pairwise act sum sq = correlation of multiple pairs of np.sum(np.sum(maps[segmentation] vectors. .T * data, axis=0) ** 2)

Fast way to compute correlation of multiple pairs of vectors without

computing all pairs as would with corr(A,B). Borrowed from Oli at Stack

overflow. Note the resulting coefficients vary slightly from the ones

obtained from corr due differences in the order of the calculations.

(Differences are of a magnitude of 1e-9 to 1e-17 depending of the tested

data).

Parameters

A : ndarray, shape (n, m)

The first collection of vectors

B : ndarray, shape (n, m)

The second collection of vectors

axis : int

The axis that contains the elements of each vector. Defaults to 0.

Returns

corr : ndarray, shape (m,)

For each pair of vectors, the correlation between them.

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An = A - np.mean(A, axis=axis)

Bn = B - np.mean(B,
axis=axis)

An /= np.linalg.norm(An, axis=axis)

Bn /= np.linalg.norm(Bn, axis=axis)

return np.sum(An * Bn,
axis=axis)

def

plot_segmentation(segmentation,
data, times):

"""Plot a microstate segmentation.

Parameters

segmentation : list of int

For each sample in time, the index of the state to which the sample has been assigned.

times : list of float

 $\label{eq:the_time-stamp} \mbox{ for each } \\ \mbox{sample.}$

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gfp = data.std(axis=0)

```
n states =
                                           Parameters
len(np.unique(segmentation))
                                           _____
    plt.figure(figsize=(6 *
                                           maps : ndarray, shape
np.ptp(times), 2))
                                       (n channels, n maps)
    cmap =
                                               The prototypical
plt.cm.get cmap('plasma',
                                       microstate maps.
n states)
                                           info : instance of
    plt.plot(times, gfp,
                                       mne.io.Info
color='black', linewidth=1)
                                               The info structure of
    for state, color in
                                       the dataset, containing the
zip(range(n states),
                                       location of the
cmap.colors):
                                               sensors.
        plt.fill between(times,
gfp, color=color,
where=(segmentation == state))
                                           plt.figure(figsize=(5*
   norm =
                                       len(maps), 2)
mpl.colors.Normalize(vmin=0,
                                           layout =
vmax=n states)
                                       mne.channels.find layout(info)
    sm =
                                           for i, map in
plt.cm.ScalarMappable(cmap=cmap,
                                       enumerate (maps):
norm=norm)
                                       plt.subplot(1, len(maps), i+1)
    sm.set array([])
                                       mne.viz.plot topomap(map,
   plt.colorbar(sm)
                                       layout.pos[:, :2])
   plt.yticks([])
                                       # plt.title('%d' % i)
   plt.xlabel('Time (s)')
    plt.title('Segmentation into
%d microstates' % n states)
   plt.autoscale(tight=True)
   plt.tight layout()
def plot maps(maps, info):
    """Plot prototypical
microstate maps.
```