



Stimulating States of Parkinsonian Tremor

Internship Report



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Problem Statement / Motivation

- Tremor is defined as a rhythmic, involuntary oscillatory movement of a body part
- AFFECTING AS MANY AS 41 MILLION PATIENTS WORLDWIDE
- Caused by neurological degeneration

Patients cannot even do **basic tasks**No permanent cure possible until today
Medications work but lose their effectiveness

PROBLEM:

Tremor is <u>variable</u> -> This makes it hard to treat (**optimal stimulation parameters** are changing) -> so we need to define states -> understand how to treat



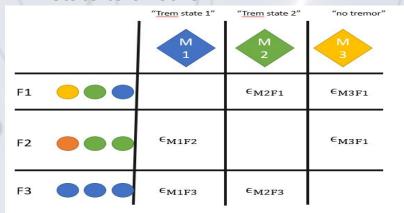




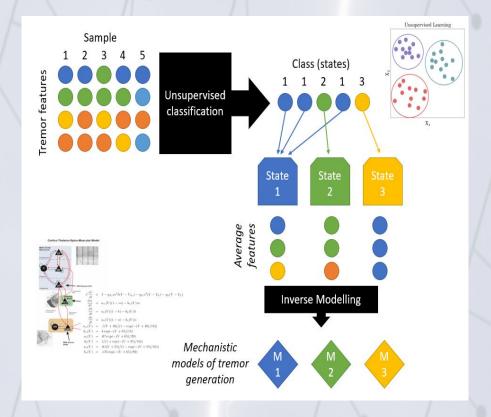


Tremor changes throughout the day and depending on activity/medication.

- Detect tremor states and their characteristics
- Find parameters to change from one state to another.



Need to find/learn transition matrix



Presentation Overview

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- Preprocessing of datasets
- Details of feature space calculated
- Clustering algorithm
- Optimization of clustering algorithm
- Validation on simulated data(from oscillators)
- Results on unsupervised data under:
 - O Single subject with unique data
 - O Single subject with overlapping data (80%)
 - O Nine subjects with unique data
- Future work / Other points



Methodology

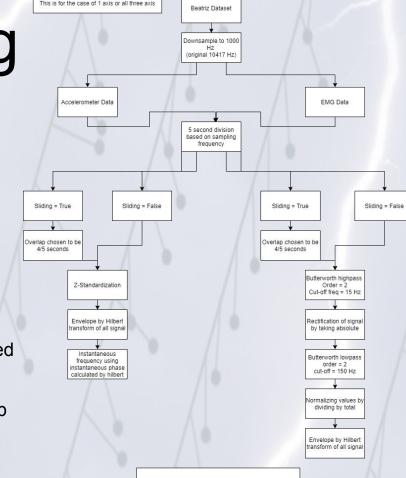
Preprocessing pipeline



Dataset (5-6 minutes of recordings) extraction:

- Total patients/samples 10
- Accelerometer
- EMG
- Accelerometer data downsampled -> 1000 Hz, so each sample has 1/1000 second duration (original 10417 Hz)
- High pass filtered the EMG at 15 Hz -> remove DC-offset - output signal centered around zero
- Low pass filtered the EMG at 150 Hz -> EMG signals are mostly dominant until up to 150 Hz

Window size + overlap is parameter to be optimized



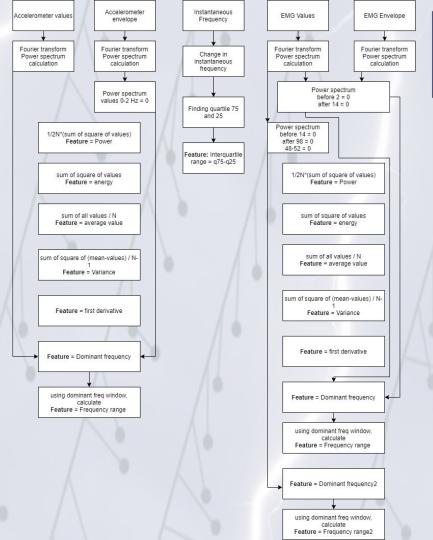
Outputs: Accelerometer values Accelerometer Envelope Instantaneous frequency EMG values EMG Envelope

Feature Extraction

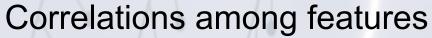
- Features are:
 - Power
 - Energy
 - Average value
 - Variance
 - First derivative
 - Dominant frequency
 - Frequency range
- 31 features in total
 - 7 of Accelerometer values
 - 7 of Accelerometer Envelope
 - 1 of Tremor Stability Index
 - 9 of EMG values
 - 7 of EMG Envelope
- Computed for each of the 5 second instance

The features have correlations with each other

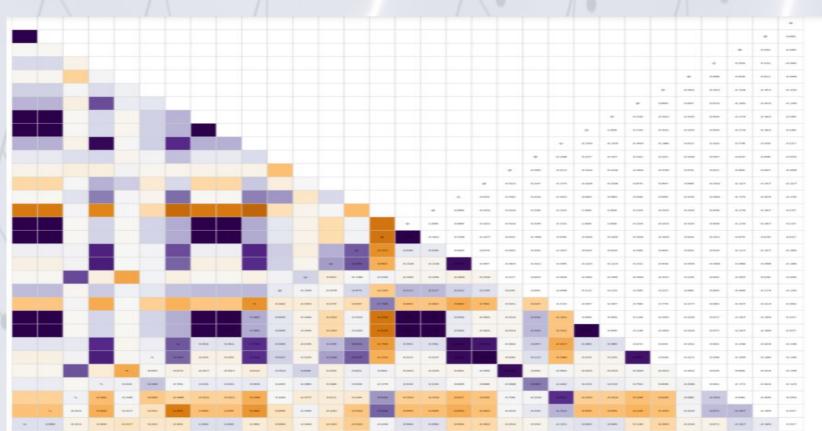
Here N is the number of values in the 5 second interval



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Clustering Algorithm

As it is unsupervised data, we do not know the actual clusters possible.

DBScan:

- Based on core point, border point and noise point
- No need to tell the number of clusters (like KMeans) + detects outliers too
- But challenge is to find right hyperparameters **eps** and **min_samples values**
 - Eps (epsilon distance): minimum distance between samples to be labeled as same cluster
 - Min_samples: minimum number of points to make a cluster to be considered one not outliers

Finding the optimal values:

- Now using K-means algorithm along with Silhouette distance to find the optimal number of clusters.
- Using that optimal cluster number to find minimum distance(eps) b\w points
- then using that distance and optimal cluster number to find **min_samples** parameter of DBScan

Source:

https://medium.com/@mohantysandip/a-step-by-step-approach-to-solve-dbscan-algorithms-by-tuning-its-hyper-parameters-93e693a91 289



















Results and Validation





Simulated data obtained from model of tremor that contains different transitions and have different dynamics.

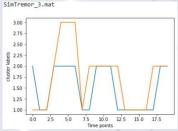
We use clustering to decode what the transitions are

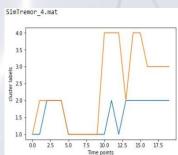
- Yellow = Actual
- Blue = Predicted

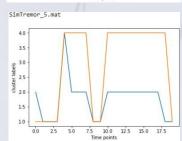
Source:

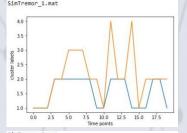
https://journals.plos.org/ploscompbiol/article?id=10.1371/journal.pcbi.1009281

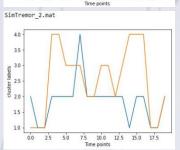
- States being close together and absorbing one another
- The difference between readings is not significant

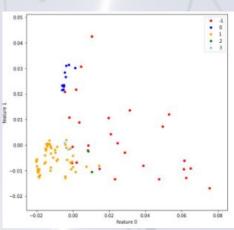












```
from sklearn.cluster import DBSCAN
clustering = DBSCAN(eps=0.011, min|_samples=2).fit(norm_matrix)
cluster=clustering.labels_
print(len(set(cluster)))
unique, counts = np.unique(cluster, return_counts=True)
dict(zip(unique, counts))
```

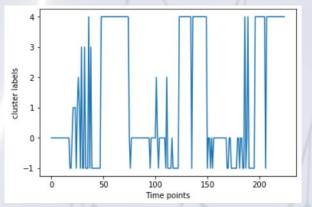


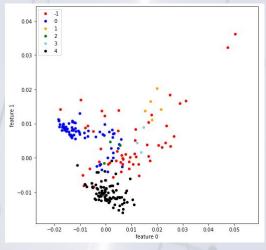
One Person All Axis (ACC+EMG) (without overlap)



Using one person, all axis.

- Optimal clusters = 4
- Min_points = 4
- Eps = 0.01031
- 18 = EMG variance
- 10 = ACC_env variance
- CPU time to compute -
 - ~1-2 mins





```
from sklearn.cluster import DBSCAN
clustering = DBSCAN(eps=0.01031, min_samples=4).fit(norm_matrix)
cluster=clustering.labels_
print(len(set(cluster)))
unique, counts = np.unique(cluster, return_counts=True)
dict(zip(unique, counts))
```

```
{-1: 56, 0: 74, 1: 5, 2: 3, 3: 3, 4: 84}
```

```
print(pca.explained_variance_ratio_)
# number of components
n_pcs= pca.components_.shape[0]
most_important = [np.abs(pca.components_[:][i]).argmax() for i in range(n_pcs)]
print(most_important)
```

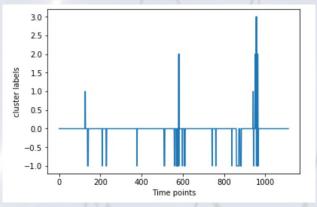
```
[0.32240092 0.22684091 0.19854599 0.06516051 0.03687473 0.03654934 0.02119217 0.01952025 0.01768636 0.01105906] [18, 18, 10, 3, 13, 13, 23, 14, 23, 6]
```



One Person All Axis (ACC+EMG) (with overlap)

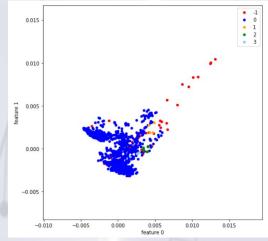
Using one person, all axis.

- Optimal clusters = 4
- Min_points = 4
- Eps =0.0023000000000000004
- 18 = EMG variance
- 10 = ACC_env variance



0.03058722 0.02575668 0.01964407 0.01244224]

[18, 18, 10, 30, 3, 13, 23, 6, 25, 14]



```
print(pca.explained_variance_ratio_)
# number of components
n_pcs= pca.components_.shape[0]
most_important = [np.abs(pca.components_[:][i]).argmax() for i in range(n_pcs)]
print(most_important)

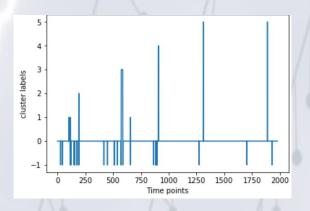
[0.32259502 0.22853658 0.16328989 0.07447037 0.04350702 0.03677964
```

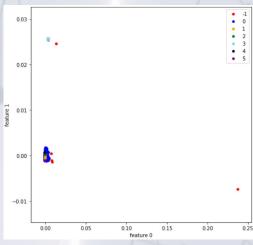


All Person All Axis (ACC+EMG) Beatriz Dataset

without overlap

- Optimal clusters = 2
- But here I put 5 to show that if we change number of clusters then results change
- 18 = EMG variance
- 24 = EMG_envelope power





```
print(pca.explained_variance_ratio_)
# number of components
n_pcs= pca.components_.shape[0]
most_important = [np.abs(pca.components_[:][i]).argmax() for i in range(n_pcs)]
print(most_important)

[0.76555548 0.15087007 0.01736329 0.01710454]
```

[18, 24, 10, 21]

Modular code

- Modification of code to make it more user friendly
- For using all these functions:
 - o python main.py
- For loading data:
 - import preprocessing helper
 - Data = preprocessing_helper.ThreeAxisACC(directory)
- Similarly can run for 1-axis, 3-axis Acc or EMG with or without sliding window of any length

```
feature_extraction.py

import matplotlib.pyplot as plt

# ^^^ pyforest auto-imports - don't write above this line
import numpy as np
import scipy.io
from scipy.fftpack import fft
from scipy import signal
from scipy.signal import hilbert

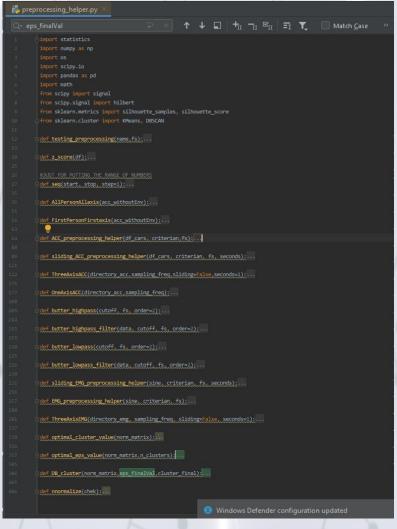
from matplotlib import rcParams

def TSI feature(acc instFreq):...

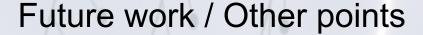
def dom freq with range(power spectrum1, frequency):...

def featureACC(data, cover, sampling rate):...

def featureEMG(data, cover, sampling rate):...
```









Todo tasks:

- Clustering on Beatriz dataset with overlap
- Clustering on Carolina's tremor data posture and spiral drawing known "transitions"
- Validation on simulated data 1(with overlap) and 2(with and without overlap)
- See the feature importance using Univariate ANOVA
- Refine the code to be a pipeline usable for everyone (GITHUB)

Feature extraction using Deep learning:

- Regarding transfer learning:
 - o Pretrained CNNs (that have been trained on real images) will be much confused by your 2D reshaped data
 - o Pretrained timenet (RNN) wouldn't have worked either because of the difference in domain
- Solution is to learn the representation from scratch, given "enough" data
- For starting we can use 1D convolutional autoencoders

Thank you for the opportunity!



Any Questions?