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Automatic Epileptic Seizure Onset Detection Using Matching Pursuit: A Case Study

Thomas L. Sorensen †, Ulrich L. Olsen ‡, Isa Conradsen †, Jonas Henriksen †, Troels W. Kjaer *, Carsten E. Thomsen † and Helge B. D. Sorensen †

Abstract — An automatic alarm system for detecting epileptic seizure onsets could be of great assistance to patients and medical staff. A novel approach is proposed using the Matching Pursuit algorithm as a feature extractor combined with the Support Vector Machine (SVM) as a classifier for this purpose. The combination of Matching Pursuit and SVM for automatic seizure detection has never been tested before, making this a pilot study. Data from red different patients with 6 to 49 seizures are used to test our model. Three patients are recorded with scalp electroencephalography (sEEG) and three with intracranial electroencephalography (iEEG). A sensitivity of 78-100% and a detection latency of 5-18s has been achieved, while holding the false detection at 0.16-5.31/h. Our results show the potential of Matching Pursuit as a feature extractor for detection of epileptic seizures.

I. INTRODUCTION

About 1% of the world’s population suffers from epilepsy [1][2], making it one of the most frequent neurological disorders only outnumbered by stroke and headache [3]. About 75% of epilepsy patients can be seizure free on antiepileptic drugs, and some of the remaining 25% can be treated with other procedures, like surgical resection of the epileptic focus, a vagus nerve stimulator or a ketogenic diet [4].

The goal of this study is to build an automatic onset detection for epileptic seizures. Such an alarm would give patients suffering from epilepsy an opportunity to leave their homes knowing that family or medical personnel can come to their rescue if they encounter a seizure. Furthermore it is important to register the number of seizures the patient encounter in a given time frame. This can give medical doctors insight on how well a treatment is working. It can also be important to know when a patient has a seizure, in case of acute treatment, or if a tracer drug has to be administered for an ictal SPECT-scan. An automatic trigger for the vagus nerve stimulator is another possibility, since it has the greatest effect if it is activated early in the seizure [5].

An automated seizure detection system would also assist in detecting seizures in large encephalography (EEG) data sets, that often include recordings from several days.

Automatic seizure detection is not a new idea. Through the past couple of decades many attempts have been made, to find the optimal algorithm for classification, primarily using intracranial EEG (iEEG) or scalp EEG (sEEG) [6]. More recently other approaches have been attempted such as accelerometers, electromyography (EMG) and angular velocity recordings [1][4].

We have applied the Matching Pursuit algorithm on both iEEG and sEEG data providing features, which will be used with the Support Vector Machine (SVM) classifier. The algorithm was first used to study ictal EEGs by Jouyy et al. in 2003 [7]. However this is the first time SVM has been combined with Matching Pursuit for seizure onset detection.

II. METHOD

A. Clinical data

We have included six patients (pt.) with a total of 133 seizures in 305 hours of recordings (rec.) in this study. To investigate if the robustness of the algorithm depends on whether data is collected intracranially or extracranially, two of the patients are recorded with sEEG and two are recorded with iEEG. The EEG-data is recorded at the Epilepsy Monitoring Unit (EMU) at Rigshospitalet University Hospital, Copenhagen. The sEEG-data are recorded at a sampling frequency of 200 Hz from patients admitted for diagnostic workup, using Stellate™ Harmonie with 21-25 EEG channels, placed using the 10-20 system.

<table>
<thead>
<tr>
<th>Pt.</th>
<th>Sex</th>
<th>Age</th>
<th>Rec.</th>
<th>Modality</th>
<th>Type</th>
<th># of Seizures</th>
</tr>
</thead>
<tbody>
<tr>
<td>P1</td>
<td>M</td>
<td>6</td>
<td>49 h</td>
<td>sEEG</td>
<td>pGTCS</td>
<td>10</td>
</tr>
<tr>
<td>P2</td>
<td>M</td>
<td>63</td>
<td>8 h</td>
<td>sEEG</td>
<td>CPS</td>
<td>49</td>
</tr>
<tr>
<td>P3</td>
<td>F</td>
<td>33</td>
<td>44 h</td>
<td>sEEG</td>
<td>SPS</td>
<td>35</td>
</tr>
<tr>
<td>P4</td>
<td>M</td>
<td>45</td>
<td>95 h</td>
<td>iEEG</td>
<td>CPS</td>
<td>20</td>
</tr>
<tr>
<td>P5</td>
<td>F</td>
<td>28</td>
<td>66 h</td>
<td>iEEG</td>
<td>SPS/SPS</td>
<td>13</td>
</tr>
<tr>
<td>P6</td>
<td>M</td>
<td>45</td>
<td>43 h</td>
<td>iEEG</td>
<td>SPS/SPS</td>
<td>6</td>
</tr>
<tr>
<td>Sum</td>
<td></td>
<td></td>
<td>305 h</td>
<td></td>
<td></td>
<td>133</td>
</tr>
</tbody>
</table>

- pGTCS = primary Generalized Tonic Clonic Seizures
- CPS = Complex Partial Seizures
- SPS = Simple Partial Seizures

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The iEEG-data are recorded from patients monitored prior to epilepsy surgery. It is recorded at a sampling frequency of 200 Hz using grid and strip electrodes. The seizures are outlined by a specialist in clinical neurophysiology (Troels W, Kjaer, MD, Ph.D.), to have a frame of reference for training the detection algorithm, and for the calculation of latency between the true EEG onset and the onset estimated by the algorithm. Furthermore three channels that clearly show epileptic activity are chosen by the specialist, which greatly minimizes the amount of data. Table I shows information for the four patients and their seizure types.

B. Matching Pursuit

The Matching Pursuit algorithm was developed in the early 1990’s by Mallat and Zhang [9]. The fundamental concept is to approximate a signal by the sum of functions found in a dictionary. The functions found in the dictionary, referred to as atoms. Often Gabor functions are used, which are sinusoids multiplied by a gaussian function [9]. This study uses the Gabor function for the decomposition. The software used to compute the Matching Pursuit algorithm, is the original software by Mallat and Zhang, rewritten by Dr. C. Jouny and Dr. P. Franaszczuk from the Epilepsy Research Laboratory at Johns Hopkins Medical Institutions. It can be downloaded at: http://erl.neuro.jhmi.edu/mpsoft/. Dr. Supratim Ray from Harvard Medical School has adapted the code to run from within MATLAB.

To find the Gabor function that best describes the signal, the inner product between the Gabor function and the signal is calculated. The Gabor function that leads to the highest inner product is used as the amplitude for the Gabor function and the Gabor function is subtracted from the signal. An iterative process is run until the energy is below a specified threshold. The Matching Pursuit algorithm is described by equation (1) where a finite number of Gabor functions, \( m \), are used to decompose the signal [9].

\[
f = \sum_{n=0}^{m-1} \langle R_n f, g_{\gamma_n} \rangle g_{\gamma_n} + R^m f \tag{1}
\]

\( R^m f \) is the residual of the \( m \)th iteration and \( g_{\gamma_n} \) is the \( n \)th Gabor function.

Because the Gabor functions are sinusoids limited in time by a gaussian function, a non-complex Gabor function is given by:

\[
g_{\gamma}(t) = K(\gamma)e^{-\pi \left( \frac{t-u}{\Delta t} \right)^2} \cos(\omega(t-u)+\phi), \tag{2}
\]

where \( \gamma = \{u, s, \omega, \phi\} \). \( u \) shifts the atom in time, \( s \) defines the width of the atom, \( \omega \) is the frequency of the atom in rad/s and \( \phi \) is the phase of the atom. \( K(\gamma) \) is a scaling factor making \( |g_{\gamma}(t)| = 1 \). Matching Pursuit is an iterative process and therefore a stop criteria is needed. Jouny et. al [7] introduced a stop criteria that is simply an energy threshold of the last atom. The decomposition will continue to run until the energy of the last atom is lower than the specified energy threshold.

C. Features

The Matching Pursuit algorithm returns the number of Gabor functions needed to reconstruct the signal. In [7], focus is on the number of atoms \( m \) needed to describe the signal. They state that the complexity of ictal EEG is higher than the complexity of interictal EEG, thus more atoms are needed to describe ictal EEG signals than is needed for interictal EEG. Therefore we apply the density of Gabor functions normalized with the window length, Gabor Atom Density (GAD) [7][8], as a feature for seizure detection. The equation for GAD is:

\[
GAD = \frac{m}{\Delta t \cdot \Delta f} = \frac{2 \cdot m}{N}, \tag{3}
\]

where \( \Delta t \) is the window size in seconds and \( \Delta f \) is the size of the frequency scale; making \( N \) the window size in samples.
We found that the frequencies increase significantly in the EEG during a seizure. This makes the mean of all the gabor functions within a window, \( \text{Mean Atom Frequency} \) (MAF), another possible feature enabling the algorithm to accurately distinguish between seizure and non-seizure. MAF can be described by:

\[
\text{MAF} = \frac{1}{m} \sum_{n=1}^{m} \omega_{g_{n}},
\]

where \( \omega_{g_{n}} \) is the frequency in rad/s of the atom \( g_{n} \).

It has been proved that working on data from multiple channels increases the accuracy, why three focal channels are used for this study. Taking the GAD and MAF for each of the three channels, gives a total of six features. The features are fed as input to the SVM classifier.

For the scalp EEG the features are extracted from the signal by applying a 512 sample window to a 200 Hz signal, hence 2.56 seconds. For iEEG the window size is 2048 samples, also sampled at 200 Hz. The reason for the different window sizes is to optimize the difference between interictal and ictal data, and is a result of testing the performance with different sizes. An overlap of 75% is used on both modalities resulting in a temporal resolution of 0.64 and 2.56 seconds respectively. For the stop criteria an energy threshold of 200 \( \mu \text{V}^2 \) was used for sEEG, as was found in [7] to be the optimum. For the iEEG data an energy threshold of 500 \( \mu \text{V}^2 \) was used as it results in the largest difference between the interictal and ictal data. On Fig. 2 the two features calculated for a 230 second period is shown. During the seizure, a clear rise in both GAD and MAF can be observed.

### III. Results

Fig. 3 shows the two features, GAD and MAF, for one channel recorded using iEEG. It can be seen that the seizure epochs are well separated from the background data and confined to a relatively small area. This leads to a low \( \text{False Detection Rate} \) (FDR), which is particularly true for the iEEG data, as can be seen in table II.

The sEEG data have higher FDRs and by looking at the feature plots for these data (not depicted) it can be seen that, though they are separated from the background data, they are not as clearly separated as the iEEG data are.

For some patients it was found that both GAD and MAF peaked early in the seizure and then decayed steadily to the end of the seizure. This indicates that these features are best suited for seizure onset detection, and might not be able to

### Table II

<table>
<thead>
<tr>
<th>Patient</th>
<th># of Seizures</th>
<th>Sensitivity (%)</th>
<th>FDR (1/h)</th>
<th>Latency (s)</th>
</tr>
</thead>
<tbody>
<tr>
<td>P1</td>
<td>10</td>
<td>100</td>
<td>0.59</td>
<td>18.3</td>
</tr>
<tr>
<td>P2</td>
<td>49</td>
<td>91.4</td>
<td>5.31</td>
<td>9.1</td>
</tr>
<tr>
<td>P3</td>
<td>35</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>P4</td>
<td>20</td>
<td>95.0</td>
<td>0.20</td>
<td>6.06</td>
</tr>
<tr>
<td>P5</td>
<td>13</td>
<td>77.8</td>
<td>0.16</td>
<td>5.67</td>
</tr>
<tr>
<td>P6</td>
<td>6</td>
<td>100</td>
<td>1.8</td>
<td>7.44</td>
</tr>
</tbody>
</table>
detect seizure duration. This has not been investigated in this study, but is a subject for our future work.

The algorithm is patient-specific. This means that the SVM is trained for each patient, and then tested on the rest of the seizures and background activity for the same patient.

From Table II it can be seen that the algorithm in this initial case study is capable of detecting 78-100% of all seizures, while keeping the FDR between 0.16-5.31/h. The detection latency was found to be 5-8s for iEEG and 9-18s for sEEG. We were unable to detect any seizures of P3 with the features used in this study.

IV. DISCUSSION & CONCLUSION

The reason for the high FDR in patient P2, is due to a large amount of artifacts. The algorithm is sensitive to the artifacts found in this particular patient. A solution to avoid the artifact could be to pre-filter the data before feeding it to the algorithm. The seizure dynamics for P3 are different from the other patients. This results in a poor detection and a high FDR. For P3 different features are needed for effective seizure detection.

Another issue is the number of electrodes. We used three focal electrodes for the detection. In patient P4 and P5 the epileptic focus is mainly in one hemisphere, but the patients do encounter some seizures that have focus in the opposite hemisphere. These seizures are not detected, since no electrodes from the focus area was used in the algorithm. A better way would be to include all the available electrodes. This could also introduce the idea of focus localization.

This is a case study, so the full potential of the method is not fully investigated. The goal is to improve performance, especially on the onset detection latency which does seem plausible. With an artifact removal algorithm, some of the artifacts introduced when the medical personnel are interacting with the patient etc. could be avoided. This would remove the need for the epoch constraint, and thereby result in a better latency performance. Furthermore a larger patient database would show the robustness of the method better, and perhaps introduce the idea of a non-patient specific approach based on different seizure types. Also a larger patient database could show if there is a difference in modality, i.e. iEEG or sEEG, and if there is significant differences between the different types of seizures. Results show a tendency for iEEG to be the more robust modality for this algorithm, but further work will have to be conducted to confirm this findings.

The idea of using Matching Pursuit for seizure detection seems very promising, and we will investigate it further to achieve even better results than presented in this paper.

V. ACKNOWLEDGMENTS

The authors would like to thank neurophysiology assistant Lennart Dermt from Rigshospitalet University Hospital, Copenhagen, for invaluable help collecting, preprocessing and reformating the EEG-data. Also a special thanks to Dr. Supratim Ray from Harvard Medical School for his help with implementing the Matching Pursuit algorithm into MATLAB.

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