SVM-Based Sleep Apnea Identification Using Optimal RR-Interval Features of the ECG Signal

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Abstract—Sleep apnea (AP) is the most commonly known sleeping disorder characterized by pauses of airflow to the lungs and often results in day and night time symptoms such as impaired concentration, depression, memory loss, snoring, nocturnal arousals, sweating and restless sleep. Obstructive Sleep Apnea (OSA), the most common SA, is a result of a collapsed upper respiratory airway, which is majorly undiagnosed due to the inconvenient Polysomnography (PSG) testing procedure at sleep labs. This paper introduces an automated approach towards identifying sleep apnea. The idea is based on efficient feature extraction of the electrocardiogram (ECG) signal by employing a hybrid of signal processing techniques and classification using a linear-kernel Support Vector Machine (SVM). The optimum set of RR-interval features of the ECG signal yields a high classification accuracy of 97.1% when tested on the Physionet Apnea-ECG recordings. The results provide motivating insights towards future developments of convenient and effective OSA screening setups.

Keywords: sleep apnea, PSG, ECG, RR interval, features extraction, SVMs.

I. INTRODUCTION
Sleep is the circadian rhythm which is among the most crucial needs in our day to day activities. On average, humans spend approximately one-third of their lifespan sleeping. Getting enough hours of sleep indicate repaired blood pressure, heart rate, and relaxed muscles and tissues [1].

A sleeping disorder occurs when one cannot sleep and has symptoms like excessive daytime sleepiness and fatigue. Sleep Apnea (SA) is among the very common respiratory sleeping disorders characterized by cessations of airflow to the lungs or having a very low breath. The cessations lasting in more than 10 seconds considered as apnea event might occur 5 to 30 times in an hour and up to 400 per night [2]. Clinically, sleep apnea is divided into Obstructive Sleep Apnea (OSA) and Central Sleep Apnea (CSA). OSA, being the most common SA, is generally caused by a collapse of the upper respiratory airway. CSA is a neurological condition where brain fails to appropriately control breathing [3] [4].

Statistics show that out of 18 million Americans suffering from OSA, around 10 million remain undiagnosed [5]. The undiagnosed cases are due to inconvenience, expenses and unavailability of testing. The Polysomnography (PSG) is the current and traditional testing process which is a standard procedure ordered for all sleep disorders. This testing records the breath airflow, respiratory movement, oxygen saturation, body saturation, body position, electroencephalogram (EEG), electrocuglagram (EOG), electromyogram (EMG), and electrocardiogram (ECG) to determine the sleep stages [6].

II. RELATED WORK
Over the past few years, most of the related research has focused on detecting OSA through statistical features of different signals such as thorax and abdomen effort signals, nasal air flow, oxygen saturation, electrical activity of the heart (ECG), and electrical activity of the brain (EEG).

Many studies perform detection of OSA through heart rate variability (HRV) and the ECG signal. Quiceno-Manrique et al. [8] proposed a simple diagnostic tool for OSA with a high accuracy (up to 92.67%) using time-frequency distributions and dynamic features in ECG signal. Moreover, based on spectral components of heart rate variability, frequency analysis was performed in [9] using Fourier and Wavelet Transformation with appropriate application of the Hilbert Transform, where the sensitivity was 90.8%. In addition, in [10] a bivariate autoregressive model was used to evaluate beat-by-beat power spectral density of HRV and R peak area, where the classification results showed accuracy higher than 85%.
During periods of prolonged OSA, cyclic increases and decreases of heart rate are typically associated with the apneic phase and the resumption of breathing [11]. Therefore, the technique in this work also relies on features of the ECG signal to detect and quantify these periods of OSA by the fully automated identification of these dynamic features in the RR-interbeat interval series based on the ones suggested by Chazal et al. [6], and Yilmaz et al. [12].

III. METHODOLOGY

In this work, the overall design involves acquiring the ECG signal. This signal is then processed to cancel the noise and detect RR-interval. Then, a combination of the most effective set of RR-interval based features of the ECG signal is calculated for classification. In what follows, our detection system design is described and shown in Figure 1.

A. Database

The database used in this study is available from the PhysioNet web site [13]. The Apnea-ECG Database contains 70 recordings, containing a single continuous ECG signal varying in length of approximately 8 hours duration. The sampling frequency of ECG signal is 100 Hz, with 16-bit resolution, with one sample bit representing 5µV [6].

The database was scored by clinical experts by dividing the recordings into a set of one-minute segments. The segments were classified as “apnea”, if at any time during that minute there was evidence of SA on the basis of respiration and oxygen saturation. Otherwise, it was classified as “normal” [6].

B. Noise cancellation and R Wave detection

SA episodes consist of bradycardia during apnea followed by tachycardia upon its cessation, which represent cyclic variations in the duration of a heartbeat, also known as RR intervals of ECG signal [6]. Generally, the ECG sleep apnea recognition techniques used have two parts: characteristics (or features) extraction, and waveform classification and recognition [14].

The characteristics extraction includes noise cancellation and QRS complex wave detection. The R-wave which has the highest (or lowest) value in the QRS complex wave is the outstanding characteristic of the ECG signal.

The R-wave detection technique used in this paper is a modified version of the traditional “Pan and Tompkins” algorithm [15] with the use of adaptive filters in noise cancellation. The whole algorithm is divided into five-steps; noise cancellation using adaptive filtering, signal slope detection, squaring, windowing, and RR-wave interval calculation.

1) Adaptive Filtering: The Least Mean Square adaptive algorithm is one of the most robust techniques used to reduce any random noise signal interfaced to the ECG. A step size of $0.8 \times 10^{-5}$ and filter length of $10^6$ can be used to cancel any noise added to the recorded ECG.

2) Signal Slope Detection: A differentiator is used to compute the QRS complex slope waveform information. A five point derivative is used with the following transfer function:

$$H(z) = (1/8T)(-2z^{-2} - 2z^{-1} + 2z^1 + Z^2)$$ (1)

The difference equation is:

$$y(nT) = (1/8T)[-x(nT - 2T) - 2x(nT - T) + 2x(nT + T) + x(nT + 2T)]$$ (2)

3) Squaring: The point by point squaring function is described by the following equation:

$$y(nT) = [x(nT)]^2$$ (3)

This results in positive data points and also performs nonlinear amplification of the differentiated ECG frequencies.

4) Windowing: Additional waveform features are calculated by a moving-window integration equation given by:

$$y(nT) = (1/N)[x(nT - (N - 1)T) + x(nT - (N - 2)T) + ... + x(nT)]$$ (4)

Where $N$ is the number of samples in the width of the integration window. For a sample rate of 200, window of 30 samples wide (150 ms) is used.

5) RR wave interval calculation: Automatically adjustable thresholds to float over the noise are applied to the integrated waveform. The applied set of thresholds are calculated from:

$$SPK = 0.125PEAK + 0.875SPK$$ (5)

(if $PEAK$ is the signal peak)

$$NPK = 0.125PEAK + 0.875NPK$$ (6)

(if $PEAK$ is the noise peak)

$$THRESHOLD = NPK + 0.25(SPKE - NPK)$$ (7)
\[ \text{THRESHOLD}_2 = 0.5 \text{THRESHOLD}_1 \] \hfill (8)

In the above equations, all the variables refer to the integration waveform: PEAK is the overall peak, SPK is the running estimate of the signal peak, NPK is the running estimate of the noise peak, \text{THRESHOLD}_1 is the first threshold applies, and \text{THRESHOLD}_2 is the second threshold applied. Every time a peak is recognized, a QRS complex is identified in the filtered and integrated waveform. The RR average is then given by taking the mean of the eight most recent consecutive RR intervals.

\[ RR_{\text{AVERAGE}} = 0.125 \left( RR_{n-7} + RR_{n-6} + \ldots + RR_n \right) \] \hfill (9)

C. Features Extraction

Our technique relies on a large set of an effective combination of ECG signal features. These features could potentially be used for classification. The features considered are a novel hybrid of features extracted from [6] and [12]. The following are the most effective set of RR-interval based features of the ECG signal for apnea detection:

- Mean epoch and recording RR-interval.
- Standard deviation of the epoch and recording RR-interval.
- The NN50 measure (variant 1), defined as the number of pairs of adjacent RR-intervals where the first RR-interval exceeds the second RR-interval by more than 50 ms.
- The NN50 measure (variant 2), defined as the number of pairs of adjacent RR-intervals where the second RR-interval exceeds the first RR interval by more than 50 ms.
- Two pNN50 measures, defined as each NN50 measure divided by the total number of RR-intervals.
- The SDSD measures, defined as the standard deviation of the differences between adjacent RR-intervals.
- The RMSSD measures, defined as the square root of the mean of the sum of the squares of differences between adjacent RR-intervals.
- Median of RR-intervals.
- Inter-quartile range, defined as difference between 75th and 25th percentiles of the RR-interval value distribution.
- Mean absolute deviation values, defined as mean of absolute values obtained by the subtraction of the mean RR-interval values from all the RR-interval values in an epoch.

The classification results confirm the improved accuracy compared to the [6] and [12] techniques alone.

D. Data Randomization

Since apnea is defined as a pause in breathing, and can last from a few seconds to minutes (almost \(\geq\) 10 sec), we investigate and analyze three cases of data partitioning to determine the best accuracy that can be achieved. The apnea and regular data are partitioned into 10 sec, 15 sec, and 30 sec pieces. In this step, we separate the training data and testing data with 80\% for training and 20\% for testing. A MATLAB built-in function (rand) is used to determine whether a feature set in 10 sec, 15 sec or 30 sec of data belongs to test group or rule creation group. After the signals are separated, we perform the training for SVMs.

E. Support Vector Machines (SVMs)

In order to investigate apneic epoch detection, we use SVMs as a classification method. SVM is one of a powerful machine learning technique from supervised learning category, which used as a training algorithm for analyzing data and recognizing patterns. When we have two classes of data, SVM performs classification by building a maximal margin hyperplane that optimally separates the data into two groups. In general, the larger the margin is, the lower the generalization error of the classifier. SVM handles the separation by using a kernel function to map the data into a different space with a hyperplane. There are many kernels available for SVM, which provides flexibility for the constructed hyperplane to partition the data [4].

In our implementation, we use a linear kernel function to map the training data into kernel space. In the optimization process, we use a method called sequential minimal optimization to find the separating hyperplane.

IV. RESULTS

To build our model, we used MATLAB toolset. The data records were imported as MATLAB matrices (.mat) from physionet web site. We evaluated the effectiveness of our model on the Apnea ECG database, using different records available in that database. We ran two scenarios in our experiment: the whole combination of the features, and a combination of every two separate features.

A. All features

To evaluate the performance of the classification system, two statistical indicators, Sensitivity (Se) and Specificity (Sp) in addition to the Accuracy (Acc) have been used. The sensitivity of a test is the percentage of patients in the OSA positive group correctly diagnosed, whereas the specificity is the percentage of subjects in the OSA negative group correctly classified by the test.

<table>
<thead>
<tr>
<th>Input/Output</th>
<th>Regular</th>
<th>Apnea</th>
</tr>
</thead>
<tbody>
<tr>
<td>10 sec. (Accuracy is 86.1%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Regular</td>
<td>97.2%</td>
<td>2.78%</td>
</tr>
<tr>
<td>Apnea</td>
<td>25%</td>
<td>75%</td>
</tr>
<tr>
<td>15 sec. (Accuracy is 96.5%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Regular</td>
<td>100%</td>
<td>0%</td>
</tr>
<tr>
<td>Apnea</td>
<td>7.1%</td>
<td>92.9%</td>
</tr>
<tr>
<td>30 sec. (Accuracy is 95%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Regular</td>
<td>100%</td>
<td>0%</td>
</tr>
<tr>
<td>Apnea</td>
<td>10%</td>
<td>90%</td>
</tr>
</tbody>
</table>
Our model was based on a linear kernel and classification process for 10 times to take the average value.

From Table II, SVM with linear kernel using 15 second epochs shows the best classification accuracy with high successful rate of correct prediction.

### V. CONCLUSIONS AND FUTURE WORKS

In this work, we studied the possibility of the detection of SA events from the ECG signal variation patterns. We evaluated the effectiveness of our model on the Apnea ECG database, using different records available in that database. Our model was based on a selective set of RR-interval based features that were given to SVM for classification. We evaluated our model on three different epoch lengths and different combination of two-features set scenarios. From the experimental results, we conclude that SVM with linear kernel shows the best accuracy with 15 second epoch length. Two optimum features were also derived from statistical analysis. A future direction to this work would be incorporating this work into a real time monitoring system that acquires and analyzes the ECG signal of subjects during sleep.

### REFERENCES


