Identifying Outliers in HRV–Seizure Signals using *p*-shift UFIR Baseline Estimates

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**Abstract:* Heart rate variability (HRV) is typically associated with neuroautonomic activity and viewed as a major non-invasive tool to detect seizures. The HRV has been assumed and analyzed as a stationary signal. However the presence of seizures can violate estimates of statistical parameters and conventional techniques intended to remove outliers can be inaccurate. A useful approach implies setting thresholds to compute the first and third quartiles from histogram data or residuals based on the estimated baseline. In this paper, we propose an accurate method to identify outliers in HRV measurements with partial epilepsy retaining relevant information. The baseline perturbed by the seizure in the HRV data is removed using the *p*-shift unbiased finite impulse response (UFIR) smoothing filter operating on optimal horizons. The residuals histogram is plotted and the upper bound (UB) and lower bound (LB) are computed as thresholds. A comparison is provided of a typical points detected in HRV/seizures based on several methods used to estimate the baseline. A time/frequency analysis is supplied to show the difference between the raw HRV and HRV without outliers. The method proposed is tested by partial seizures records taken from patients during continuous EEG/ECG and video monitoring.

Key-Words: Heart Rate Variability (HRV), outliers, seizures, p-shift (UFIR) smoothing filter.

1 Introduction

The term 'heart rate variability' (HRV) has been accepted to describe variations of both instantaneous heart rate and RR intervals, where R is a point corresponding to the peak of the QRS complex of the electrocardiogram (ECG) wave and RR is the interval between successive Rs [1]. Because the HRV is strongly coupled with the autonomic nervous system (ANS) [2], information from several physiological signals such as ECG is also used to detect seizures [3, 4]. Epileptic seizures are taken as indicators, which enable the diagnostic of ANS disorders. The HRV spectral content has frequency components, which are classified in three main peaks: very low frequency (VLF) < 0.04 Hz, low frequency (LF), 0.04 - 0.15Hz, and high frequency (HF) 0.15 - 0.4 Hz [5]. The LF and HF powers are associated with sympathetic nervous system (SNS) and parasympathetic nervous system (PNS) activity [6], respectively.

One of the main and oldest problems in biomedical signal processing is the noise reduction without disturbing the original signal, especially in ECG having a high content of vital information [7]. Special components of the disturbance, which immerse in noise signals, are outliers which affect significantly the performance of denoising algorithms. The outlier is defined as the measurement, which significantly deviates from the logical or normal pattern of sensed data [8]. Similarly, in statistics, an outlier is defined as a 'case that does not follow the same model as the rest of the data' [9].

There are many methods to remove outliers from data. The most simple and useful one is known as the Boxplot. This method displays batches of data [10] and only five values from a set are used: the extremes, the upper and lower hinges (quartiles), and the median [11]. However, this method is inefficient when data have non-stationary characteristics in some parts. Nevertheless, an increasingly common process to remove outliers is to estimate a model using several method, such as smoothing algorithms [12, 13, 14]. Complete the smoothing, the outliers are excluded at an arbitrary distance greater than $A\sigma$ from the model, where A can be specified by the user. Because there are many algorithms to smooth bio-signals, the selection of an ideal method faces difficult. Furthermore,

errors in the baseline optimal estimates of HRV signals strongly depend on the smoother degree.

It is now worth mentioning that an ideal baseline for HRV signals does not exist and it is far less available in the presence of partial epilepsy. Therefore, an accurate method is required to estimate the optimal baseline that helps identifying the outliers in HRV measurements. One approach used in medical applications is based on the *p*-shift unbiased finite impulse response (UFIR) filter, which minimizes the mean square error (MSE) on optimal averaging horizons N_{opt} . The *p*-shift UFIR filter does not require unavailable prior knowledge about the ECG signal statistics, but produces suboptimal estimates when $N_{\rm opt}$ is set properly. In [15], two ways have been proposed to find N_{opt} : a) by minimizing the trace of the error covariance matrix and b) by minimizing the derivative of the trace of the mean square value (MSV) associated with the raw signal and estimate as function of N. Because an exact mathematical model of the HRV signal is not available, the only option is to find N_{opt} using the second approach.

In this paper, we develop an approach to estimate the baseline model of HRV with partial seizures and identify the outliers by employing the *p*-shift UFIR filter. We also compare the number of atypical points based on the residuals calculated using several algorithms: LOESS, RLOESS, LOWESS, Moving Average, and Spline. To test the algorithms, we use measurement data from Post-Ictal Heart Rate Oscillations in Partial Epilepsy available at MIT-BIH [17]. The paper is structured as follows. In Section 2, a brief description is given for the HRV based on the electrocardiogram model and database features used in this work. In Section 3, we design the proposed algorithm to estimate the baseline of HRV signals and develop an algorithm to calculate N_{opt} . In Section 4, we find $N_{\rm opt}$ and identify the outliers using the *p*-shift UFIR filter and some other algorithms. Finally, conclusions and further work are discussed in Section 5.

2 HRV from Partial Seizures

In a typical ECG signal, a mixture of deflections Q, R, and S generated by the heart's muscles form a complex QRS. The RR interval, called tachogram, is obtained by computing the distance between the peaks R immerses in the complex QRS. The HRV is then represented with the instantaneous heart rate (IHR), which is the inverse of the RR interval [18].

This work is based on data of HRV measurements with partial seizures, which represent the shape alteration causing perturbations in the baseline. The HRV data are analyzed from partial seizures recorded in 5 female patients during continuous EEG/ECG and video monitoring considered as a *gold standard* [17]. The heart beat annotations were obtained using fully automated methods described in [20] and available in 'physionet'.

3 Baseline Estimate

The baseline estimate can be obtained using diverse FIR filtering algorithms [19]. However, smoothing allows for higher accuracy when the solution is not required in real time, as in our case. Therefore, we will base our investigation on the *p*-shift UFIER filter, which is briefly described below.

3.1 *p*-Shift UFIR Filtering.

Let us represent the ECG measurements as $s_n = x_n + \nu_n$, where *n* is the discrete time index, x_n is the ECG signal and ν_n is the random noise, which can be supposed to be white Gaussian. Assuming that x_n is the baseline and ν_n is the measurement residual, we can find $\nu_n = s_n - x_n$ based on an accurate approximation of x_n . The residual ν_n contains outliers in white Gaussian noise environment. Now, x_n can be expressed with a degree polynomial on a horizon [n - N + 1 - p, n - p] of N points, where $p \leq 0$ is a time-shift. The model can be estimated using the *p*-shift UFIR filtering algorithm proposed in [21]. In the convolution-based form, the estimate of x_n can be found via data taken from [n - N + 1 - p, n - p] as

$$\hat{x}_{n|n-p} = \sum_{i=p}^{N-1+p} h_{li}(p) s_{n-i} , \qquad (1)$$

where $h_{ln}(p) \triangleq h_{ln}(N, p)$ is the $\{N, p\}$ -variant impulse response of the *l*-degree discrete FIR filter.

Satisfied the unbiasedness condition of $E\{\hat{x}_{n|n-p}\} = E\{x_n\}$, the *p*-variant $h_{li}(p)$ can be represented as [22, 21]

$$h_{li}(p) = \sum_{j=0}^{l} a_{jl}(p)i^{j}, \qquad (2)$$

where $i \in [p, N - 1 + p]$ and coefficients $a_{jl}(p)$ are

$$a_{jl}(p) = (-1)^j \frac{M_{(j+1)1}(p)}{|\mathbf{D}(p)|},$$
 (3)

where $|\mathbf{D}(p)|$ is the determinant of matrix $\mathbf{D}(p) = \mathbf{V}^{\mathbf{T}}(p)\mathbf{V}(p)$, in which $\mathbf{V}(p)$ is the $(l+1)\times(l+1)$ Vandermonde matrix. The region of existence of $h_{li}(p)$ is

the following [21, 22],

$$h_{li}(N,p) = \begin{cases} \text{nontrivial}, & p \le i \le N - 1 + p \\ 0, & \text{otherwise} \end{cases}$$
(4)

Although the *p*-shift filtering estimate can be provided for any UFIR filter degree, the low-degree filters have found most applications owing to better noise reduction.

3.1.1 Optimimal Horizon

The most important parameter of the UFIR filter is the optimal horizon N_{opt} . Because the mathematical model of HRV and baseline signal is not available, we find N_{opt} follow the method proposed in [15, 16]. The horizon N_{opt} can be estimated from the measurement residual $s_n - \hat{x}_{n|n-p}(N)$ by minimizing the derivative $\partial \eta(N) / \partial N$, where η is the MSV $\eta(N) = E\{[s_n - \hat{x}_{n|n-p}(N)]^2\}$, and solving the optimization problem [15]

$$\hat{N}_{\text{opt}} = \arg\min_{N} \frac{\partial \eta(N)}{\partial N} + 1.$$
(5)

Provided N_{opt} , the baseline of HRV van be estimates using the *p*-shift UFIR filtering approach.

4 Baseline Estimates and Outliers Detection

In order to estimate the baseline of HRV with partial seizures, N_{opt} is required. Provided N_{opt} , the shift p can be specified for each degree as shown in [21]. Following the suggestion given in [16], we select l = 2 and specify p by $p = -(N_{\text{opt}} - 1)/2$. Using the physionet database, we next compute the MSE of HRVseizures (s01, s02, ..., s07) for the estimate $x_{n|n-p}(N)$ as function of N[1:2:105]. The results are plotted in Fig. 1a.

Next, we solve the optimization problem (5) and find $N_{\rm opt} = 61$ as shown in Fig.1b. Typical values are labeled when the point is larger than the upper bound $UB = Q_3 + w(Q_3 - Q_1)$ or smaller than the lower bound $LB = Q_1 - w(Q_3 - Q_1)$, where Q_1 and Q_3 are specified in the 25% and 75% sense, respectively. The value w = 1.5 corresponds to $\approx \pm 2.7\sigma$ and the probability of 0.993 for normal distributed data.

The baseline for each measurement can now be obtained using N_{opt} to calculate the residuals and detect outliers for the given upper and lower bounds. In Fig. 2a we show a signal s07 (doted) and the baseline estimated (solid) along with the the data residual (dashed). The normalized histogram (H1) of sam-

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Figure 1: Estimation of optimal horizon as $N_{\text{opt}} = 61$: a) MSE of HRV/seizures data (s01, s02,..., s07) for the estimate $x_{n|n-p}(N)$ and b) minimization of $\partial \eta(N)/\partial N$ to solve the optimization problem (4).

ple s07 and the outliers identified using the boxplot method are plotted in Fig. 2b and Fig. 2c, respectively. Build similarly, the histogram (H2) of the residuals and its typical values are sketched in Fig. 2d and Fig. 2e.

An comparison is next provided of the baseline estimates obtained by the UFIR smoothing filter and by several other techniques such as the moving average filter (MAF), Splines, LOESS, LOWESS, and RLOESS. The LOWESS and LOESS methods employ the locally weighted linear regression to smooth data using linear polynomial and quadratic polynomial respectively. A robust version of the LOESS is called RLOESS. Taking as a benchmark the baseline estimates sketched in Fig. 3a and Fig. 3b, we show the outliers identified during the seizure of signal s07. As can be observed, the loss of information using LOESS, RLOESS, and LOWESS is greater than by others methods.

Finally, Table 1 generalizes the results. The MAF ansd SPLINE algorithms identify the minimal number of outliers, while the LOESS, RLOESS, and LOWES ones the maximum number. Unlike in other methods, the number of points detected by the *p*-shift UFIR smoothing filter is located in the middle of data. Herewith, the identification based on the boxplot of raw



Figure 2: Baseline estimation and outliers detection using the *p*-shift UFIR filter with $N_{opt} = 61$ (solid) from HRV seizures measurements s07 (doted) and residuals (dashed): (a) baseline estimates; (b)–(e) histograms (H1) and typical points for raw HRV with an extracted average.

HRV seizures has demonstrated an erratic behavior. It is also worth mentioning that the MSEs produced by these models is proportional to the number of the identified outliers.

4.1 Time/Frequency Comparison

We finish our investigation by using the time/frequency (TF) technique to analyze a signal in the time and frequency domains simultaneously. We provide it for the normalized sample frequency Fs = 1 and a 'flattop' window using the command 'tftb_window' from Matlab. A TF comparison of HRV in Pre and Post -Seizure is showed in Fig. 4a and Fig. 4b. In this example, data correspond to a 48-year-old woman with partial epilepsy. The Pre-Seizure (< 0min) demonstrates a relative HF of 0.15 - 0.4 Hz, while in Post-Seizure (> 3min), we have indicated a frequency transition between 3 and 4.5 min. Removed the outliers, the mean difference between the raw HRV and HRV is in the lapse seizure and we notice the frequency weakness from Fig. 4a to Fig. 4b. However, the principal peaks power remain almost intact.



Figure 3: Atypical points identified at the beginning of epileptic attack in s07 data and outliers found with estimated baseline using: (a) MA (cross), Spline (point) and *p*-shift UFIR (circle) and (b) LOESS (cross), RLOESS (point) and LOWESS (circle) algorithms.

Table 1: Outliers identified in HRV data in the presence of seizures from s01 to s07 using different methods estimating the baseline. The 'Normal' column means atypical values localized with the boxplot technique applied to raw HRV with an extracted average.

OUTLIERS BY METHOD						
Normal	UFIR	MA	SPLINE	LOESS	RLOESS	LOWESS
58	689	414	428	740	1049	796
1261	426	185	405	828	970	890
1578	866	508	580	1580	1965	1721
403	300	244	250	389	359	403
637	450	493	572	686	726	909
518	424	111	231	493	568	497
484	356	159	251	711	529	870
	Normal 58 1261 1578 403 637 518 484	Normal UFIR 58 689 1261 426 1578 866 403 300 637 450 518 424 484 356	Normal UFIR MA 58 689 414 1261 426 185 1578 866 508 403 300 244 637 450 493 518 424 111 484 356 159	OUTLIERS B Normal UFIR MA SPLINE 58 689 414 428 1261 426 185 405 1578 866 508 580 403 300 244 250 637 450 493 572 518 424 111 231 484 356 159 251	OUTLIERS BY METHO Normal UFIR MA SPLINE LOESS 58 689 414 428 740 1261 426 185 405 828 1578 866 508 580 1580 403 300 244 250 389 637 450 493 572 686 518 424 111 231 493 484 356 159 251 711	OUTLIERS BY METHOD Normal UFIR MA SPLINE LOESS RLOESS 58 689 414 428 740 1049 1261 426 185 405 828 970 1578 866 508 580 1580 1965 403 300 244 250 389 359 637 450 493 572 686 726 518 424 111 231 493 568 484 356 159 251 711 529

5 Conclusions

The *p*-shift UFIR smoothing filter operating on optimal horizon has demonstrated an ability to estimate the baseline in spite of statistical changes caused by the seizures. The residual distribution at the UFIR smoothing filter output is more normalized than by other techniques. Therefore, the discrimination of data has appeared to be most accurate. Because the l = 2 degree UFIR smoothing filter is invariant to sharp changes, the proposed solution allows erasing atypical values with a minimal loss of information. Finally, the time/frequency analysis has shown that the proposed method does not alter the ECG principal frequencies.

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Figure 4: Time/frequency test of the raw HRV with an extracted average: (a) a pre and post epylepsy seizure and (b) TF analysis of HRV without atypical points, which are removed based on the smooth UFIR method in the same lapse.

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