Improving accuracy of derived 12-lead electrocardiography by waveform segmentation

Indra Hardian Mulyadi1 , Nelmiawati2 , Eko Supriyanto3

^{1,3}School of Biomedical Engineering and Health Sciences, Universiti Teknologi Malaysia, Malaysia ²Department of Electrical Engineering, Politeknik Negeri Batam, Indonesia

Corresponding Author:

Indra Hardian Mulyadi, School of Biomedical Engineering and Health Sciences, Universiti Teknologi Malaysia, Johor, Malaysia. Email: indra.mulyadi@fkegraduate.utm.my

1. INTRODUCTION

Facts mentioned that around 2.6 million people above 15 years old in Indonesia suffered from coronary heart disease [1]. Around 17.3 million World population in 2013 died caused by cardiovascular diseases [2], including coronary heart disease, heart failure, hypertension, and stroke. Advanced technologies have been implemented to reduce this number and – at the same time - to increase life expectancy. The focuses is prevention [3], including prediction and early diagnosis [4], for instance personalized cardiovascular disease monitoring devices [5].

For diagnosis, cardiologists analyze morphology of the ECG waveform. Each segment in the waveform contains information of specific heart activities [6]. Optimal electrodes location for each segment is different, as investigated by Finlat et. al. [7].They introduced Eigenleads which is useful for pre-diagnosing heart pathologies and for wearable ECG, which requires high signal to noise ratio (SNR). However, for wearable ECG, practical aspects must be considered [8].

The standard 12-lead ECG with ten electrodes has been established as diagnostic reference in hospitals. However, it is impractical for 24-hours monitoring, wearable, and ambulatory applications due to difficulty to attach electrodes and sensitivity to wiring noise and motion artifacts [9]. On the other hand, nonclinical users are not trained to find the proper electrodes location, whereas misplacement of the electrodes

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may lead to misdiagnosis [10]. For these reasons, a number of studies have been conducted to employ a limited set of electrodes and transform the acquired signal to produce 12-lead ECG, which is called derived 12-lead ECG. The leads should be nearly orthogonal, for instance lead I, II, and V2 [11].

A promising technique-called EASI-was demonstrated by Dower et al. [12] and has been implemented in hospital for years. One of its benefits is easiness to find body landmarks to place electrodes Figure 1. Three approaches have been investigated to calculate EASI coefficients: generic, patient-specific, and populationspecific [13]. To obtain maximum accuracy, the coefficients should be patient-specific (personalized coefficients) [14]. To improve EASI coefficients, several techniques have been presented such as Dower's method, Linear Regression (LR), Polynomial Regression (PR), Support Vector Regression (SVR), and Artificial Neural Network (ANN) [9]. In those techniques, the coefficients were calculated by utiizing full cycle (FC) of ECG signal, i.e. all segments.

In this study, we propose a new approach by utilizing different EASI coefficients for each ECG segment; therefore, it is segment-specific (SS). We segmented the waveform into three segments: 1) PR-Interval, i.e. P wave and PR segment, 2) QRS complex, and 3) ST interval, i.e. ST segment and T wave. It was hypothesized that the result might minimize the transformation error. The proposed approach, i.e. SS, was then compared with the conventional one, i.e. FC.

2. RESEARCH METHOD

In this study, we used EASI lead system to reconstruct 12-lead ECG. Instead of calculating for full cycle ECG, we proposed to segment the ECG waveform into three segments: PR, QRS, and ST as described in Figure 2. So, we have different transformation coefficients for each segment. At the end, the reconstructed lead of segment PR, QRS, and ST were then combined to be a full ECG signal. For instance, $I_{\text{Perived-PR}}$, $I_{\text{Perived,ORS}}$, and $I_{\text{Perived,ST}}$ were combined into $I_{\text{Perived,SS}}$ as formulated in (1). $V_{\text{Perived,SS}}$ represents voltage of lead I to V6.

Figure 1. EASI lead system: Electrodes placement
Figure 2. Segmenting one cycle of ECG signal

into three segments: PR interval, QRS complex, and ST interval.

Data sets used in this study were taken from Physionet [15]. Calculation was performed using MATLAB (MathWorks, Natick). To evaluate our new segment-specific (SS) approach, we applied it into six existing methods and compared the result with conventional full cycle (FC) approach. The six existing methods are Dower's method with improved generic coefficients, Dower's method with individual (patient-specific) coefficients, Linear Regression (LR), $2nd$ -degree Polynomial Regression (PR), $3rd$ -degree PR, and Artificial Neural Network (ANN). To simplify, these methods are called method A, B, C, D, E, and F. Method A to E were also presented in our previous work [16].

a. Method A: Dower's method with improved generic coefficients [17] This method has been applied in commercial EASI 12-lead ECG machines. Generic EASI coefficients were pre-determined statistically from a data set of 983 adult subjects. The (2) was used to calculate derived ECG lead voltage ($V_{\text{Derived},A}$) from three bipolar leads: ES (V_{ES}), AS (V_{AS}), and AI (V_{AI}) with coefficients of $\beta_{0,A}, \beta_{1,A}$, and $\beta_{2,A}$. As the coefficients are pre-determined, SS and FC result in a same result.

$$
V_{\text{Derived},A} = \beta_{0,A} V_{ES} + \beta_{1,A} V_{AS} + \beta_{2,A} V_{AI} \tag{2}
$$

- b. Method B: Dower's method with individual (patient-specific) coefficients Basically, it is similar to method A, but it utilizes individual (patient-specific) coefficients instead of using pre-determined ones.
- c. Method C: Linear Regression (LR)

The (3) is a formula used to calculate derived ECG lead voltage ($V_{\text{Derived},c}$) from unipolar EASI lead: E (V_E) , A (V_A) , S (V_S) , and I (V_I) with coefficients of $\beta_{0,C}$, $\beta_{1,C}$, $\beta_{2,C}$, $\beta_{3,C}$, and $\beta_{4,C}$.

$$
V_{\text{Derived},C} = \beta_{0,C} + \beta_{1,C} V_E + \beta_{2,C} V_A + \beta_{3,C} V_S + \beta_{4,C} V_I
$$
\n(3)

d. Method D: 2nd-degree Polynomial Regression (2nd PR)

The (4) is a formula used to calculate derived ECG lead voltage ($V_{\text{Perived},D}$) from unipolar EASI leads: E (V_E) , A (V_A) , S (V_S) , and I (V_I) with coefficients of $\beta_{0,D}, \beta_{1,D}, \dots, \beta_{14,D}$.

$$
V_{Derived,D} = \beta_{0,D} + \beta_{1,D} V_E + \beta_{2,D} V_A + \beta_{3,D} V_S + \beta_{4,D} V_I + \beta_{5,D} V_E V_A + \beta_{6,D} V_E V_S + \beta_{7,D} V_E V_I + \beta_{8,D} V_A V_S + \beta_{9,D} V_A V_I + \beta_{10,D} V_S V_I + \beta_{11,D} V_E^2 + \beta_{12,D} V_A^2 + \beta_{13,D} V_S^2 + \beta_{14,D} V_I^2
$$
\n(4)

- e. Method E: 3^{rd} -degree Polynomial Regression $(3^{rd} PR)$
	- The (5) is a formula used to calculate derived ECG lead voltage ($V_{\text{Derived},E}$) from unipolar EASI leads: E (V_E) , A (V_A) , S (V_S) , and I (V_I) with coefficients of $\beta_{0,E}, \beta_{1,E}, \dots, \beta_{33,E}$.

$$
V_{Derived,E} = \beta_{0,E} + \beta_{1,E} V_E + \beta_{2,E} V_A + \beta_{3,E} V_S + \beta_{4,E} V_I + \beta_{5,E} V_E V_A + \beta_{6,E} V_E V_S + \n\beta_{7,E} V_E V_I + \beta_{8,E} V_A V_S + \beta_{9,E} V_A V_I + \beta_{10,E} V_S V_I + \beta_{11,E} V_E^2 + \beta_{12,E} V_A^2 + \beta_{13,E} V_S^2 + \n\beta_{14,E} V_I^2 + \beta_{15,E} V_E^3 + \beta_{16,E} V_A^3 + \beta_{17,E} V_S^3 + \beta_{18,E} V_I^3 + \beta_{19,E} V_E^2 V_A + \beta_{20,E} V_E^2 V_S + \n\beta_{21,E} V_E^2 V_I + \beta_{22,E} V_A^2 V_E + \beta_{23,E} V_A^2 V_S + \beta_{24,E} V_A^2 V_I + \beta_{25,E} V_S^2 V_E + \beta_{26,E} V_S^2 V_A + \n\beta_{27,E} V_S^2 V_I + \beta_{28,E} V_I^2 V_E + \beta_{29,E} V_I^2 V_A + \beta_{30,E} V_I^2 V_S + \beta_{31,E} V_E V_A V_S + \n\beta_{32,E} V_E V_A V_I + \beta_{33,E} V_E V_S V_I
$$

f. Method F: Artificial Neural Network (ANN) with Backpropagation algorithm utilizing Levenberg-Marquardt optimization. The network consists of 88 nodes: one input layer (4 nodes), two hidden layers (72 nodes) with hyperbolic tangent sigmoid activation function, and one output layer (12 nodes) with linear activation function. The network is described in Figure 3. To speed up the calculation, it was divided into 12 networks, one network for each lead. The trained networks produced $V_{\text{Perived},F}$.

Transformation coefficients of method B to F were calculated from segment-specific (PR, QRS, ST) as well as full-cycle approaches. The results from both were then compared to evaluate their performance. Root mean squared error (RMSE) (6) and correlation coefficient (r) (7) were used as a metric to measure the performances, where N is number of signal samples, $V_{Measured,i}$ is voltage of measured (original) ECG of i -th sample, and $V_{\text{Derived},i}$ is derived ECG of *i*-th sample.

$$
RMSE = \sqrt{\frac{1}{N} \sum_{i=1}^{N} (V_{Measured,i} - V_{\text{Period},i})^2}
$$
(6)

$$
r = \frac{N(\sum_{i=1}^{N} V_{Measured,i} \times V_{\text{Derived},i}) - (\sum_{i=1}^{N} V_{Measured,i})(\sum_{i=1}^{N} V_{\text{Derived},i})}{\sqrt{(7)}}
$$

$$
-\frac{1}{\sqrt{(N\sum_{i=1}^{N}V_{Measured,i}^{2}-(\sum_{i=1}^{N}V_{Measured,i})^{2})}\times (N\sum_{i=1}^{N}V_{Derived,i}^{2}-(\sum_{i=1}^{N}V_{Derived,i})^{2})}
$$
(7)

Figure 3. Reconstruction of derived 12-lead ECG from lead E, A, S, and I using 88 nodes ANN (Method F)

3. RESULTS AND ANALYSIS

After obtaining segment-specific coefficients for PR, QRS, and ST, we combined coefficients from those three segments to build derived ECG for a whole segment ($V_{\text{Perived,SS}}$) using (1). RMSEs comparison of FC and SS is presented in Figure 4. For method B, C, D, and E the new combined segment-specific (SS) outperformed the conventional one (FC). The exception applies to method A, where the coefficients are fixed and pre-determined statistically for the entire segments; hence SS and FC result in a same result [17].

Figure 3 illustrates ECG waveforms from 12 leads (I, II, III, aVR, aVL, aVF, V1, V2, V3, V4, V5, and V6) obtained from measured ECG, derived ECG from conventional FC using method A, and derived ECG from new SS using method E. Table 1 details the comparison of SS and FC in term of RMSE, while Table 2 presents the correlation. From the figure and tables, we conclude that the new segment-specific approach outperformed conventional full-cycle in term of accuracy and correlation, i.e. reducing error significantly up to 74.50%, and improving correlation up to 0.66%. However, the correlation improvement, which indicates similarity degree of ECG morphology, was not significant.

Segmenting ECG waveform into three segments and then calculating the transformation coefficients for each segment results in more accurate and higher correlation derived 12-lead ECG signal. Mathematically, the segmenting can be any, i.e. not necessarily PR, QRS, and ST. However, by segmenting it into the three well-known segments, we might have benefits from the medical perspective, since each segment indicates different information of heart activities [6]. These segments are utilized for different pathological heart diagnosis; for instance, myocardial ischemia can be diagnosed by observing ST segment and T-wave. Segmenting into these three segments is also very useful for pre-diagnosed patients.

Potential drawback of the segment-specific approach would be difficulty of detecting P, Q, R, S, and T wave automatically; however, several methods to detect these waves have been introduced [18-20]. Besides segment-specific, this study also assumed patient-specific for calculation. Although difficult to implement, patient-specific approach is the most accurate than generic or population-specific. The difficulty can be handled by current technologies; for instance, the individual coefficients can be stored in a memory card [11] or Cloud database.

Derivation technique in this paper used data based on Mason-Likar electrode placement. However, for clinical application, a more accurate derivation might be generated from standard 12-lead placement. Mason Likar lead placement may cause misdiagnosis, for instance misdiagnosis and inefficient ablation in predicting outflow tract premature ventricular contraction (OT-PVC) origin [21].

Figure 4. RMSEs of conventional, i.e. FC and our new approach, i.e. SS. SS outperformed FC in term of accuracy

Table 1. Generated RMSEs (μ V) of 12-lead derived ECG from method A to F. The new segment-specific (SS) outperformed full-cycle (FC) in term of accuracy

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Leads	А	B		C		D		E		F	
	FC & SS	FC	SS	FC	SS	FC	SS	FC	SS	FC	SS
	34.91	22.19	14.02	14.16	10.39	9.27	7.06	7.23	2.55	8.46	8.13
П	113.29	38.01	15.03	28.98	9.54	18.59	3.96	10.10	1.77	26.19	7.87
Ш	135.79	34.73	19.39	29.14	13.95	17.76	7.56	10.58	2.52	18.95	3.99
aVR	48.99	25.83	10.83	17.55	7.13	11.70	4.30	7.01	1.79	10.00	4.86
aVL	81.27	22.09	15.16	17.74	11.34	10.69	7.04	7.53	2.37	12.10	5.10
aVF	123.86	34.67	15.87	28.19	10.76	17.58	4.89	9.69	1.76	21.45	5.53
V1	91.98	11.75	7.50	10.04	6.42	6.86	3.10	4.55	1.17	8.25	4.01
V ₂	108.53	48.97	26.26	41.97	20.91	22.35	9.91	12.90	2.98	16.49	4.83
V ₃	57.22	36.54	21.34	31.93	15.98	20.53	8.07	11.26	2.83	16.23	5.51
V4	89.23	72.10	26.38	43.35	18.13	25.86	9.79	15.98	4.66	26.64	5.04
V ₅	60.46	33.96	12.33	17.71	8.23	9.02	4.80	5.97	1.80	11.45	3.62
V6	36.69	11.15	5.14	10.37	4.25	6.51	2.29	3.56	0.91	6.43	5.13
Average	81.85	32.67	15.77	24.26	11.42	14.73	6.06	8.86	2.26	15.22	5.30
Error Reduction		50.37%		52.93%		58.83%		74.50%		65.17%	

Table 2. Correlation coefficient of 12-lead derived ECG from method A to F. The new segment-specific (SS) outperformed full-cycle (FC) in term of waveform correlation.

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Figure 3. ECG signal: Measured (original), derived from full-cycle (FC) using method A, and derived from segment-specific (SS) using method E. The figure shows that the SS approach results in more accurate derived ECG

4. CONCLUSION

Segment-specific approach for deriving 12-lead ECG from a limited set of electrodes has been demonstrated. This approach outperformed the conventional full cycle calculation when compared using several methods (Dower's method with individual (patient-specific) coefficients, LR, 2nd PR, 3rd PR, and ANN). It was able to reduce error significantly up to 74.50% as well as improve the correlation up to 0.66%. This new approach is promising to provide accurate derived 12-lead ECG for personalized (24-hours monitoring), wearable, and ambulatory ECG. Future works would be implementing this approach for real clinical use.

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