

Amplitude and Phase Classification of ECG data



Chibueze Emmanuel Ogbonnaya
With: S. P. Preston, A. T. A. Wood and K. Bharath

University of Nottingham
pmxceog@nottingham.ac.uk

September 2, 2018

- 1 Electrocardiogram
- 2 Data Preprocessing and Registration
- 3 Fitting Parametric Models
- 4 Classification
- 5 Conclusion

- An Electrocardiogram (ECG) is used to record the electrical activity of the heart to identify and locate pathology.
- The ECG is essential for the diagnosis and management of abnormal cardiac rhythms.

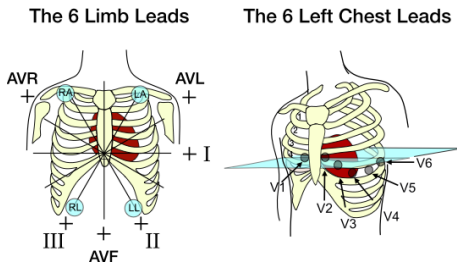
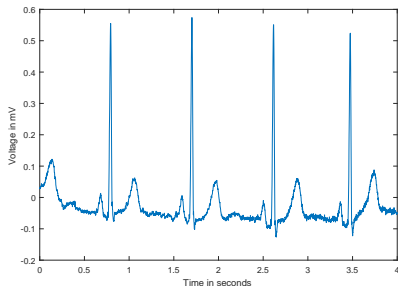
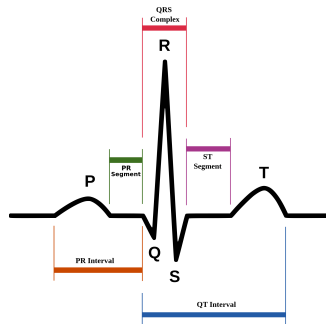


Figure: ECG 12-lead placement



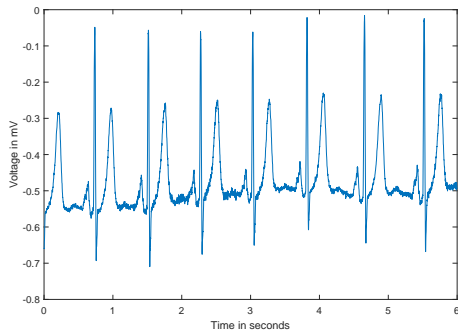
(a) ECG Signal for an individual from lead I



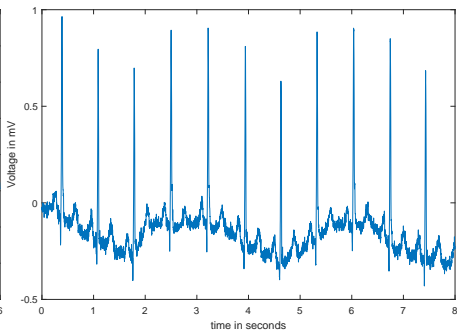
(b) Features of a Typical ECG Signal

Figure: Sample ECG Signal and Features

It is difficult to detect ECG features in a noisy ECG signal.



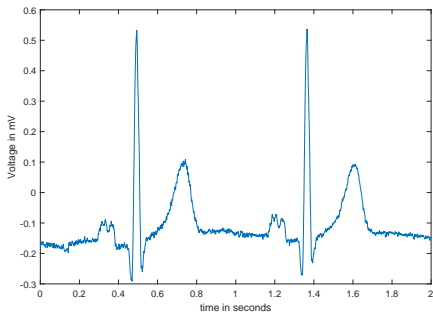
(a) Nice ECG signal



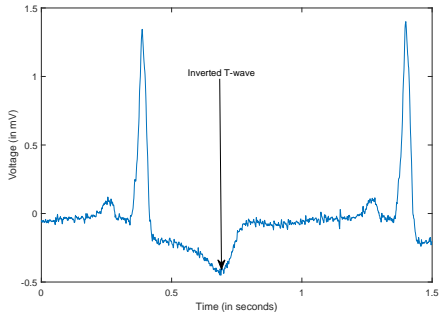
(b) Noisy ECG signal with baseline drift

Figure: ECG Signals of two different Healthy Controls showing effect of Noise

- Myocardial Infarction: ST elevation
- Cardiomyopathy: inverted T wave and prolonged QT interval.



(a) Normal ECG for healthy control



(b) ECG with inverted T wave

Figure: ECG changes caused by heart conditions

We propose using RR intervals for functional representation of an ECG signal.

- An RR interval corresponds to a heartbeat.
- The RR interval contains important features of interest: ST segment and T wave.

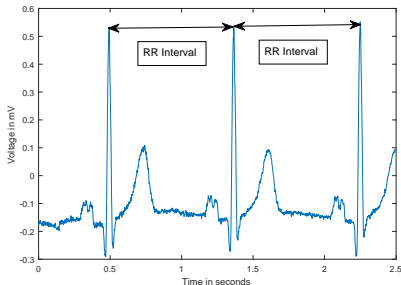
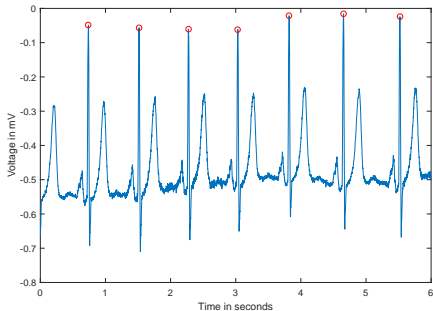
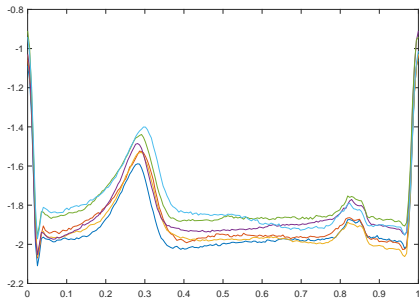


Figure: An ECG signal showing RR intervals



(a) ECG with R peaks shown in Red



(b) Chopped-up RR functions

Figure: ECG signal and chopped-up functions

Data is taken from the PTB ECG database and contains 10 seconds recordings. We consider the conventional 12-leads.

For our data, we have ECG for

- Healthy controls
- Myocardial infarction
- Cardiomyopathy.

Registration and Fitting

- Remove noise in ECG signals through amplitude registration.
- Estimate amplitude and phase components of registered ECGs using parametric models.

Classification

Classification of ECGs using estimated amplitude and phase components.

Model

$$y_i(t) = b_i x_i(t) + \sum_{j=1}^q a_{ji} u_j(t)$$

where $t \in [0, 1]$. We define the following:

- $x_i(t)$ are the observed RR functions.
- $y_i(t)$ are the registered RR functions
- $u_j(t)$ form an orthonormal basis function for noise.

Registration implies estimating a_{ji} and b_i with template $f(t)$

$$\underset{a_{ji}, b_i \in \mathbb{R}}{\text{minimise}} \quad \sum_{i=1}^n \|y_i - f\|^2.$$

Example Solution for $f(t) = 0$

- Use the zero function as template

Solutions:

$$\hat{a}_{ji} = -b_i \langle x_i, u_j \rangle,$$

$$y_i(t) = b_i \left(x_i(t) - \sum_{j=1}^q \langle x_i, u_j \rangle u_j(t) \right).$$

- Estimate b_i : Constraint $\sum_{i=1}^n \log b_i = 0$.
- $\hat{b}_i = c_i^{-1/2} (\prod_{i=1}^n c_i^{-1/2n})$.
- $c_i = \|x_i\|^2 - \sum_{j=1}^q \langle x_i, u_j \rangle^2$.

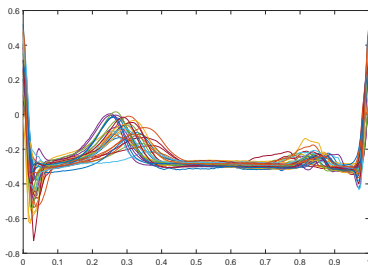
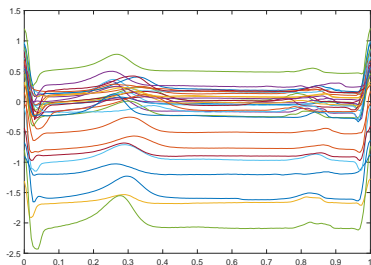


Figure: Left: Observed RR functions. Right: Amplitude registration.

- We use sum of Gaussian functions and B-splines to fit the RR functions
- The Gaussian models have been used previously to generate synthetic ECGs in Clifford (2006).

For the Gaussian mixture model with phase $t \in [0, 1]$, we have

$$z(t, \alpha_i, \theta_i, \beta_i) = \sum_{j=1}^k \alpha_{ij} \exp \left[- \frac{(t - \theta_{ij})^2}{2\beta_{ij}^2} \right] \quad (1)$$

where $\theta_{i1} = 0 \leq \theta_{i2} \leq \dots \leq \theta_{ik} = 1$.

To fit this model to actual ECG signal $y_i(t)$, we will need to solve the non-linear optimisation problem

$$\min_{\alpha_i, \theta_i, \beta_i} \int_0^1 (y_i(t) - z(t, \alpha_i, \theta_i, \beta_i))^2 dt$$

subject to $\theta_{i2} - \theta_{i3} < 0, \theta_{i3} - \theta_{i4} < 0, \dots, \theta_{i(k-2)} - \theta_{i(k-1)} < 0.$

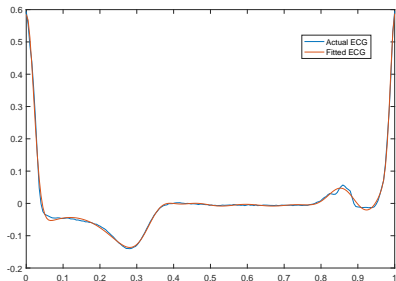
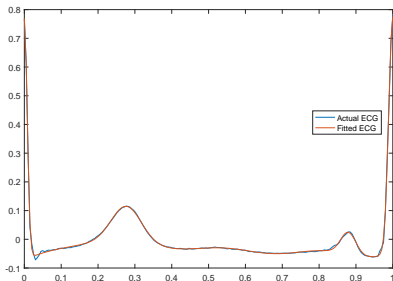


Figure: Fitting gaussian parametric models to actual ECGs ($k = 11$). Left: healthy control ECG. Right: Cardiomyopathy ECG.

- 1 Determine Template $\hat{\mu}(t)$.
- 2 Using template, fix $\beta_i = \beta$, $\theta_i = \theta$, $i = 1, \dots, n$

$$\min_{\alpha, \beta, \theta} \int_0^1 (\hat{\mu}(t) - z(t, \alpha, \beta, \theta))^2.$$

- 3 New Model: $z(t, \alpha_i) = \sum_{j=1}^k \alpha_{ij} \exp \left[-\frac{(t-\theta_j)^2}{2\beta_j^2} \right]$.
- 4 For each i , estimate α_i .

- 1 Determine Template $\hat{\mu}(t)$.
- 2 Using template, fix $\beta_i = \beta$, $\alpha_i = \alpha$, $i = 1, \dots, n$

$$\min_{\alpha, \beta, \theta} \int_0^1 (\hat{\mu}(t) - z(t, \alpha, \beta, \theta))^2 dt.$$

- 3 New Model: $z(t, \theta_i) = \sum_{j=1}^k \alpha_j \exp \left[- \frac{(t - \theta_{ij})^2}{2\beta_j^2} \right]$
- 4 For each i , estimate $\theta_{i2}, \dots, \theta_{i(k-1)}$.

- 1 We conduct a two-sample Hotelling's t-test.
- 2 Healthy vs Cardiomyopathy
- 3 NULL HYPOTHESIS: No difference in amplitude

Result

- $F = 19.3507$, $p\text{-value} = 1.4433 \times 10^{-15}$.
- Strong evidence of difference in amplitude.

Classification is done using the estimated components, for Lead I.

Accuracy		
Method	LDA	SVM
Gaussian	0.9714	0.9429
BSpline	0.9714	0.8714

Table: Amplitude Classification of Cardiomyopathy

Accuracy		
Method	LDA	SVM
Gaussian	0.9286	0.9000
BSpline	0.9571	0.9571

Table: Phase Classification of Cardiomyopathy

Classification results from Lead I.

Accuracy		
Method	LDA	SVM
Gaussian	0.8477	0.8376
BSpline	0.8325	0.8426

Table: Amplitude Classification of MI

Combining multiple leads by concatenation, this improves to

Accuracy		
Method	LDA	SVM
Gaussian	0.8782	0.9086
BSpline	0.8731	0.8832

Table: Amplitude Classification of MI (Multiple Leads)

- We have proposed amplitude registration models for ECG signals.
- Parametric models are a good alternative to dimension reduction techniques like FPCA.
- Variable selection possible using estimated amplitude components.
- Automation greatly improves ECG diagnosis when compared to clinicians.
- Applicable to analysis of gait data for diagnosis of Parkinson's.

Reference	Method	Result
McCabe et al. (2013)[2]	Physicians	Sensitivity: 65%, Specificity: 79%
Sun et al. (2012)[3]	ST segments using 5-order polynomial	Sensitivity: 92.3%, Specificity: 88.1%
Kurtek et al. (2013)[1]	NN (SRVF)	Accuracy: 90%
Previous work	Functional PCA	Accuracy: 92.86%
Proposed	Gaussian Model	Accuracy: 90.86%

Table: Comparison of methods for detection of myocardial infarction

Reference	Method	Result
Tucker et al. (2013)	Horizontal FPCA (SRVF)	0.9429
Tang and Müller (2008)	Pairwise Synchronisation (PACE)	0.7857
Proposed	B-Spline Model	0.9571

Table: Comparison of methods for detection of Cardiomyopathy

Thank You



Kurtek, Sebastian and Wu, Wei and Christensen, Gary E and Srivastava, Anuj (2013)

Segmentation, alignment and statistical analysis of biosignals with application to disease classification

Journal of Applied Statistics .



McCabe, James M and Armstrong, Ehrin J and Ku, Ivy and Kulkarni, Ameya and Hoffmayer, Kurt S and Bhawe, Prashant D and Waldo, Stephen W and Hsue, Priscilla and Stein, John C and Marcus, Gregory M and others (2013)

Physician accuracy in interpreting potential ST-segment elevation myocardial infarction electrocardiograms

Journal of the American Heart Association .



Li Sun and Yanping Lu and Kaitao Yang and Shaozi Li (2012)

ECG Analysis Using Multiple Instance Learning for Myocardial Infarction Detection

IEEE Transactions on Biomedical Engineering .