View Link





Matches World Wide Web Match View Link World Wide Web Match View Link **World Wide Web Match** View Link World Wide Web Match View Link **World Wide Web Match** View Link **World Wide Web Match** View Link World Wide Web Match View Link **World Wide Web Match** View Link **World Wide Web Match** View Link World Wide Web Match View Link World Wide Web Match View Link World Wide Web Match View Link **World Wide Web Match** View Link World Wide Web Match View Link World Wide Web Match View Link World Wide Web Match

World Wide Web Match
View Link
World Wide Web Match View Link
World Wide Web Match View Link
World Wide Web Match View Link
World Wide Web Match View Link
World Wide Web Match View Link
World Wide Web Match View Link
World Wide Web Match View Link
World Wide Web Match View Link
World Wide Web Match View Link
World Wide Web Match View Link
World Wide Web Match View Link
World Wide Web Match View Link
World Wide Web Match View Link
World Wide Web Match View Link
World Wide Web Match View Link
World Wide Web Match View Link
World Wide Web Match View Link
World Wide Web Match View Link

36	World Wide Web Match
30	<u>View Link</u>
37	World Wide Web Match View Link
38	World Wide Web Match View Link
39	World Wide Web Match View Link
40	World Wide Web Match View Link
41	World Wide Web Match View Link
42	World Wide Web Match View Link
43	World Wide Web Match View Link
44	World Wide Web Match View Link
45	World Wide Web Match View Link
46	World Wide Web Match View Link
47	World Wide Web Match View Link
48	World Wide Web Match View Link
49	World Wide Web Match View Link
50	World Wide Web Match View Link
51	World Wide Web Match View Link
52	World Wide Web Match View Link
53	World Wide Web Match View Link
54	World Wide Web Match View Link

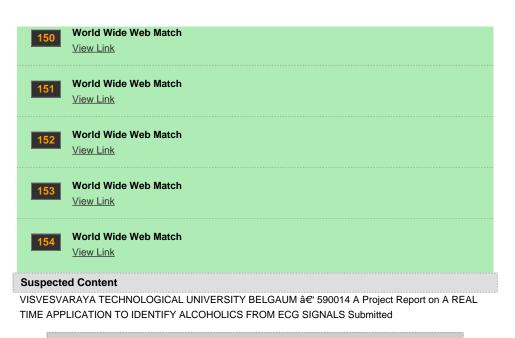
55	World Wide Web Match View Link
56	World Wide Web Match View Link
57	World Wide Web Match View Link
58	World Wide Web Match View Link
59	World Wide Web Match View Link
60	World Wide Web Match View Link
61	World Wide Web Match View Link
62	World Wide Web Match View Link
63	World Wide Web Match View Link
64	World Wide Web Match View Link
65	World Wide Web Match View Link
66	World Wide Web Match View Link
67	World Wide Web Match View Link
68	World Wide Web Match View Link
69	World Wide Web Match View Link
70	World Wide Web Match View Link
71	World Wide Web Match View Link
72	World Wide Web Match View Link
73	World Wide Web Match View Link

74	World Wide Web Match View Link
75	World Wide Web Match View Link
76	World Wide Web Match View Link
77	World Wide Web Match View Link
78	World Wide Web Match View Link
79	World Wide Web Match View Link
80	World Wide Web Match View Link
81	World Wide Web Match View Link
82	World Wide Web Match View Link
83	World Wide Web Match View Link
84	World Wide Web Match View Link
85	World Wide Web Match View Link
86	World Wide Web Match View Link
87	World Wide Web Match View Link
88	World Wide Web Match View Link
89	World Wide Web Match View Link
90	World Wide Web Match View Link
91	World Wide Web Match View Link
92	World Wide Web Match View Link

93	World Wide Web Match View Link
94	World Wide Web Match View Link
95	World Wide Web Match View Link
96	World Wide Web Match View Link
97	World Wide Web Match View Link
98	World Wide Web Match View Link
99	World Wide Web Match View Link
100	World Wide Web Match View Link
101	World Wide Web Match View Link
102	World Wide Web Match View Link
103	World Wide Web Match View Link
104	World Wide Web Match View Link
105	World Wide Web Match View Link
106	World Wide Web Match View Link
107	World Wide Web Match View Link
108	World Wide Web Match View Link
109	World Wide Web Match View Link
110	World Wide Web Match View Link
111	World Wide Web Match View Link
· <u></u>	

112	World Wide Web Match View Link
113	World Wide Web Match View Link
114	World Wide Web Match View Link
115	World Wide Web Match View Link
116	World Wide Web Match View Link
117	World Wide Web Match View Link
118	World Wide Web Match View Link
119	World Wide Web Match View Link
120	World Wide Web Match View Link
121	World Wide Web Match View Link
122	World Wide Web Match View Link
123	World Wide Web Match View Link
124	World Wide Web Match View Link
125	World Wide Web Match View Link
126	World Wide Web Match View Link
127	World Wide Web Match View Link
128	World Wide Web Match View Link
129	World Wide Web Match View Link
130	World Wide Web Match View Link

131	World Wide Web Match View Link
132	World Wide Web Match View Link
133	World Wide Web Match View Link
134	World Wide Web Match View Link
135	World Wide Web Match View Link
136	World Wide Web Match View Link
137	World Wide Web Match View Link
138	World Wide Web Match View Link
139	World Wide Web Match View Link
140	World Wide Web Match View Link
141	World Wide Web Match View Link
142	World Wide Web Match View Link
143	World Wide Web Match View Link
144	World Wide Web Match View Link
145	World Wide Web Match View Link
146	World Wide Web Match View Link
147	World Wide Web Match View Link
148	World Wide Web Match View Link
149	World Wide Web Match View Link



in partial fulfilment of the requirement for the award of degree of BACHELOR OF ENGINEERING IN ELECTRONICS AND COMMUNICATION By



Akarsh N Kolekar [1PI13EC009] Apoorv Vatsal [1PI13EC017] Rakshith Vishwanatha [1PI13EC075] Under the Guidance of: Dr. B. Niranjana Krupa Professor, Dept. of ECE,

PES Institute of Technology DEPARTMENT OF ELECTRONICS AND COMMUNICATION ENGINEERING



P.E.S. INSTITUTE OF TECHNOLOGY (An Autonomous Institute under VTU, Belgaum)



BENGALURU - 560085 DECLARATION We

hereby declare that the project report entitled



"A REAL TIME APPLICATION TO IDENTIFY ALCOHOLICS FROM ECG SIGNALS†is the bonafide record of the

project carried out at P.E.S. Institute of Technology in partial fulfilment of the requirements for the award of degree Bachelor of Engineering in Electronics and Communication Engineering of Visvesvaraya Technological University, Belgaum



during the academic year 2017. We further declare that the project report is not submitted to any other universities in

fulfilment of the requirements for the award of any degree.



By AKARSH N. KOLEKAR (1PI13EC009) APOORV VATSAL (1PI13EC017) RAKSHITH VISHWANATHA (1PI13EC075) VISVESVARAYA TECHNOLOGICAL UNIVERSITY BELGAUM ‹ 590014 PES

INSTITUTE OF TECHNOLOGY (An Autonomous Institute under VTU, Belgaum)



BENGALURU â€' 560085 CERTIFICATE Certified to the project entitled A REAL TIME APPLICATION TO IDENTIFY ALCOHOLICS FROM ECG SIGNALS

53

Kolekar, Apoorv Vatsal and Rakshith Vishwanatha bearing University Seat Number 1PI13EC009, 1PI13EC017 and 1PI13EC075 respectively

in partial fulfilment for the award of Bachelor of Engineering in Electronics and communication of the Visvesvaraya Technological University, Belgaum during the academic year 2017. It is certified that all correction /suggestions indicated for internal assessment have been incorporated in the report deposited in the department library. The project report has been approved as it satisfies the academic requirements with respect to the project work prescribed for the said degree. Head of

Department: Dr. Chandar T S

Dept. of ECE. PES Institute of Technology, Bengaluru ‑ 560085

112

External Viva: Guide: Dr. B. Niranjana Krupa Dept. of ECE PES Institute of Technology, Bengaluru â€' 560085 Principal Dr. K S Sridhar PESIT, Bengaluru â€' 560085

Name of the Examiner Signature with Date 1. 2. ACKNOWLEDGEMENT The

53

satisfaction and euphoria that accompany the successful completion of any task would be incomplete, without the mention of people who made it possible, whose constant guidance and encouragement crown all the efforts with success. Our most sincere and grateful acknowledgement to

the P.E.S Institute of Technology

for providing us with the opportunity to pursue our degree and

119

thus helping us in shaping our career. We would like to sincerely thank our project guide, Dr. B. Niranjana Krupa

Professor at the Department of Electronics and Communication Engineering

103

for her continuous valuable guidance, advice and persistent encouragement

throughout the project work. We would like to sincerely thank

34

Dr.T.S.Chandar,

Head of the Department Electronics and Communication, for his encouragement and support throughout the project work. We are thankful to Dr.Sridhar K.

34

S., Principal of

PES Institute of Technology, Bengaluru for his encouragement and support in

142

our endeavour. We would also like to thank

all the teaching and non-teaching staff and management of PES Institute of Technology, Bengaluru for their cooperation.

Last but not the least we wish to thank our family and friends for all their love support and

64

encouragement. ABSTRACT Several medical studies reveal alcohol consumption has pronounced effects on the physiology of the consumer. These physiological changes

can be seen in the heart rate variability (HRV) of the

consumer. In this article, electrocardiogram (ECG) samples of chronically alcoholic subjects and normative subjects are collected for HRV analysis and feature extraction. The features extracted are fed to machine learning algorithms to enable the algorithms to classify subjects into alcoholic or normative classes. For this classification problem, Support Vector Machines and Extreme Learning Machines have been trained, and their performance has been compared. While

time domain, frequency domain and non-linear features are

generally extracted from ECG signals for HRV analysis, in

this study a new set of features obtained from

Autoregressive Modelling (using Exogenous Inputs) have also been used to improve the accuracy of the algorithms being trained. Table of Contents CHAPTER

2.1 ECG Sensor Circuit Design 3.2.2 Heart Rate Monitor (AD8232) 3.2.3 Raspberry-Pi

3.2.4 Analog to Digital Converter (ADC)

Support Vector Machine (SVM) 3.4.2 Extreme Learning Machine (ELM)

REFERENCE 9. APPENDIX List of Figures Fig 3.2.1_1 Traditional placement of ECG probes â€â€â€â€â ۉ€â€â€â€â€â€â€â€. Fig 3.2.1_2_1 Circuit Design-Superposed Output â€' with Probe Connection to Limbs â€â€â€â€â€â€â€ Fig 3.2.1_4 50Hz Output without Probes Connected to Limbs Fig 3.2.1_5 Circuit Design-3 Fig 3.2.1_ 6 Soldered Circuit Fig 3.2.2_1 AD8232 Fig 3.2.2_2 Internal Pin Diagram of AD8232 Fig 3.2.3 _1 Raspberry Pi 2 model B Fig 3. 2.4_1 MCP 3008 ADC Fig 3.2.5_2 MCP 3008 interface to Raspberry Pi Fig 3.2.6_1 Synchronous Data Bus Fig 3.2.6_1 Sending and receiving data using SPI Fig 3.3.1_1 Baseline Wandering and Power-line noise in ECG Fig 3.3.2.1_1 PSD of a sample ECG signal with noise components Fig 3.3.2.1_2 IIR Filtered Signal without Baseline Wandering Fig 3.3.2.2_1 Diagrammatic Representation of Fourier Transform applied to a Signal Fig 3.3.2.2_2 Diagrammatic Representation of Short Fourier Transform applied to a Signal Fig 3.3.2.2_3 Fourier Transform applied Sine Signal Fig 3.3.2.2_4 Wavelet Decomposition of a Signal Fig 3. 3.2.2 4 Wavelet Decomposition of a Signal Fig 3. 3.2.2_6 Wavelet Decomposition Tree Fig 3.3.2.2_7 Mother wavelet Fig 3.3.2.2_8 Wavelet Decomposition in different modes Fig 3.3.2.2_9 Decomposition in Tree mode Fig 3.3.2.2_10 Complete Wavelet Decomposition Fig 3.3.2.2_11 Removal of Baseline Wandering by Wavelet Decomposition and RR peak detection Fig 3.3.3.2_1 Poincare Plot Fig 3.3.3.3_1 PSD of a sample from dataset used Fig 3.3.3.4_1 MATLAB Toolbox to select order of ARX polynomial Fig 3.3.3.4_3 ARX model structure selection Fig 3.3.3.4_3 Coefficients of ARX model fit to required order and input/output signals provided Fig 3.4.1_1 Importance of regularization Fig 3.4.2_2 Accuracy for a range of hidden neurons Fig 3.5_1 Real time capture of ECG signal Fig 3.5.2_1 GUI Fig 3.5.2_2 ECG signal loading process is started Fig 3.5.2_3 ECG signal is loaded Fig 3.5.2_4 Feature extraction completed Fig 3.5.2_5 Classification completed List of Tables Table 3.2.5_1 GPIO pins of Raspberry Pi Table 3.4.3.3_1 Confusion Matrix Table 4.1_1 Results of SVM for different feature sets Table 4.1_2 Confusion matrix for SVM with ARX features Table 4.2_1 Accuracies of ELM for different feature sets Table 4.2_2 Accuracy of ELM for Leave One Out Validation with ARX order 5 Table 4.2_3 Confusion matrix for ELM with ARX features order 5 Table 4.3_1 Comparative results of SVM and ELM Table 4.3_2 Comparative Sensitivity and Specificity of SVM and ELM CHAPTER â€' 1 INTRODUCTION/OVERVIEW 1.1 Introduction The electrical activity of the heart can be recorded by the

process of Electrocardiography in which electrodes are placed on the skin and measure the potential difference between the points of measurement, which arise due to the electro-physiologic pattern of polarising and depolarising of the heart muscle. The popularity of ECG measurement stems from the fact that it is non-invasive, is very reliable, and

conveys a large amount of information about the structure of the heart, function of electrical conduction system,

condition of cells of the heart, chambers, among others, to clinicians and healthcare specialists. The effects of consumption of alcohol in bursts or over time in humans have been studied and documented extensively. Alcohol consumption affects the communication pathways in the brain, leading to changes in mood, behaviour, coordination, and ability to think clearly. Chronic alcohol consumption has also been shown to cause heart disorders such as cardiomyopathy, arrhythmias, stroke and high blood pressure, as well as liver ailments like alcoholic hepatitis, fibrosis, cirrhosis, and steatosis, as well as cause pancreatitis, weaken the immune system, and increase the risk of developing cancers of the mouth, esophagus, throat, liver and breast.

Heart Rate Variability (HRV) is the phenomenon of variation of the inter-beat interval or the time between

38

successive R-R peaks of the PQRST waveform on a standard ECG signal. This variation in the inter-beat interval is a physiological phenomenon brought about by different inputs to the Sino-Atrial (SA) Node of the heart, which is where the cluster of cells which produce electrical impulses are located inside the heart. These inputs to the

SA Node include the sympathetic nervous system (SNS) and parasympathetic nervous system (PSNS)

38

as well as humeral factors. The SNS and PSNS are the 2 divisions under the autonomic nervous system, which performs unconsciously and is responsible for regulating bodily functions such as heart rate, et cetera. The SNS is known to prepare the body for intense physical activities, that is, activate the fight-or-flight response. This includes changes such as dilation of pupils and increase in the heart rate. The preganglionic neurons originating from the spinal cord release acetylcholine, a neurotransmitter, at the synapse with the ganglia, which then activates the receptors on the postganglionic neurons, which in turn releases norepinephrine, which stimulates the adrenergic receptors on target tissues, and this is a response which primarily acts on the cardiovascular system, increases Heart Rate, and hence affects HRV. The PSNS relaxes the body and slows down or inhibits many high energy expending functions, and hence lowers the heart rate and blood pressure among other physiological changes. Hence, increased SNS activity and decreased PSNS activity results in decreased HRV.

High Frequency (HF) activity (0.15 Hz to 0.4 Hz) is known to

83

be associated with PSNS, whereas Low Frequency (LF) activity (0.04 Hz

to 0.15 Hz) is accepted to be a combination of both SNS and PSNS activity. Consumption of alcohol has been known to act as a depressant on the brain and nervous tissue. Several studies have linked chronic consumption of alcohol to changes in HRV, and these have also detailed the correlation between amount of alcohol consumed as well as gender to extent of change in HRV. Hence, conclusions could be drawn about differences in alcoholic and normative test subjects based purely on HRV. HRV analysis consists of several methods, which are grouped under time-domain, frequency domain and non-linear.

Time domain analysis focuses on the heart rhythm and its variations.

Changes in time domain features gives an indication of magnitude of change in autonomic tone.

Spectral analysis methods are now more sensitive

or extracting more information regarding sympathetic and parasympathetic tone. For example, the

power value of the HF content is considered as a measure of parasympathetic activity, while the power value of the LF content is reflective of both sympathetic and parasympathetic tone.

The ratio of the LF component to that of the HF component

137

is

an index of sympathetic activity as well as balance between the sympathetic and parasympathetic nerves. The

12

non-linear methods, which include the Poincare plot, reflect the

autonomic function changes associated with long term consumption of



alcohol. The plot

not only delivers an outline, but also a detailed picture about the beat-to-beat



behaviour of cardio- physiology. Studies have shown that chronic alcohol intake caused the decrease in Poincare plot indices accompanied with a decrease in area of the plot. Auto Regressive modeling of the system also provides coefficients which can be used as features. These set of compiled features are used to train the classifiers, which will be used to detect if a test subject is alcoholic or not. The classifiers chosen for comparison included

Support Vector Machines (SVM) and Extreme Learning Machines (ELM). The



Kernel trick was also applied on the dataset using

Radial Basis Function (RBF) kernel. The uniqueness of



this methodology lies in the usage of ARX coefficients as features, along with features provided from HRV analysis, and application of ELM to classification of test subjects as alcoholic or not based solely on ECG signals, it's performance compared to SVM, and construction of a simple, precise, hand held device to do everything from signal acquisition, pre-processing, feature extraction and classification, in real-time. Table 1.1_1 Diagrammatic relation of alcohol consumption, its effect on physiology and HRV, to feature extraction using the HRV analysis 1.2 Problem Statement Technology has advanced in various fields at rapid rates, however in an area that concerns the common well-being of humans, technology has remained dormant. Identifying accurately if a person is intoxicated is utmost important to keep public harm and nuisance at bay. The most common device used, the breathalyser has drawbacks that we aim to rectify. The common drawbacks of a breathalyser are: i) Contamination of SiO2 sensor requires frequent recalibration and replacement ii) Breathalysers being a medium for propagation of contagious diseases iii) Interfering components (like acetone) being higher in the breath of dieters and diabetics make them more prone to being falsely detected iv) Infrared sensors detect the absorbance of the compound as a function of the wavelength of the beam when the infrared beam is passed through the sample breath chamber. The chamber is prone to environmental pollutants and aerosols leading to errors. These disadvantages mentioned above are addressed with the use of infallible computers and well trained machine learning algorithms. Heart rate variability [HRV] obtained from Electrocardiogram [ECG] is a useful biomarker and is used extensively in our paradigm to extract features. The features extracted are then used to train the system to classify patients as chronic alcoholic or otherwise. This may be useful in discriminating individuals based on their habits while preventing other external environmental conditions from altering or corrupts the readings.

1.3 Objective The aim of this project is to: 1) Develop



a prototype to read ECG signals 2) Extract features to perform HRV analysis 3) Classify the person under test as an alcoholic or otherwise, in real time. 1.4 Proposed Methodology i. The first step is to study the advantages and disadvantages of HRV analysis over other markers for cardiovascular health and the effect of alcohol consumption on ECG signals. ii. The next step is the familiarisation with the form and structure of the ECG signals, and understanding distortions (artefacts) that can occur in the signal. Two common

artefacts seen in ECG signals are the wandering baseline and fuzzy (60 Hz) distortions caused due to probe movement, uneven conductive gel, muscle movement, etc. iii. A survey of available devices for measuring and recording ECG signals is done. The device made by a previous student connects to a phone via bluetooth and the data is then routed to a server from the phone. This device was found to be noisy and required extra software like Audacity for filtering purposes and hence was not real time. It was proposed to interface the sensor directly with the Raspberry Pi 2 and do all the processing on it, making the overall system more compact and real time. iv. Parallel to working on the sensor hardware, once the ECG waveforms are understood, work with the training

data set is started to decide the feature extraction techniques

46

and classifiers that could be implemented. Survey of tools like KubiosHRV to understand time domain and frequency domain feature extraction for HRV analysis is done. The training data set is obtained from Autonomic Lab, Department of Neurophysiology, NIMHANS and consists of 38 samples of alcoholic and 29 samples of normative subjects. v. To this data, preprocessing is performed to remove artefacts, and then

time domain, frequency domain, and non-linear methods are

60

used for feature extraction from the ECG signals. On extracting the

29

desired features, ELM and SVM are classifiers that are chosen to be trained using the dataset. Kernel functions are also tried in order to check if their usage might improve accuracy. vi. Accuracy of the classifiers is checked using

leave-one-out cross validation and k-fold cross validation,

144

and classifiers providing highest consistent average accuracy in these validation methods is selected. vii. Based on the algorithms finalised on, the controller board on which the feature extraction and classification algorithms will be implemented are chosen. The target device is the Raspberry Pi 2 currently, as

it has a 900 MHz quad-core CPU that is necessary for

16

machine learning applications. viii. Training of the classifiers to obtain weights is done on MATLAB and the trained parameters are used on the Raspberry Pi 2. ix. Finally, the sensor is interfaced with the Raspberry Pi 2, which is uploaded with trained parameters, ready to classify real time ECG data. CHAPTER †2 LITERATURE SURVEY

This chapter covers literature survey performed

in order to understand the effect of alcohol consumption on HRV, the

100

methods involved in HRV analysis, the features that can be extracted using these methods and the classifiers which can be used for the dataset obtained. It also covers the literature studied for similar implementations, to find out which features were extracted, which classifiers were used and what the accuracies obtained were. Several papers were studied for design of the sensors used in order to record ECG signals and documentation of the micro controllers that could be used to implement the real time device. 2.1 Electrocardiogram, Noise, Artefacts and Signal Acquisition The Electrocardiogram provides the healthcare specialist a lot of information about the heart, understand and come to conclusions about any underlying factors which result in variations observed in the waveforms. The ECG signal is captured

by placing several electrodes on the skin of the test subject at the

13

limbs and on the chest and measuring the magnitude of the potential at these points. There can be a standard 10 electrode ECG as well as a 3 electrode ECG in which the three electrodes are placed at the points shown by Einthoven's triangle. The 3 electrode ECG measures the potential difference between the different electrodes.

a few hundred microvolts to a few millivolts, and hence needs to be amplified greatly using amplifiers. The signal is also affected by noise, from sources like power sources, environment and static electric charges, and artefacts, caused due to muscle movement and breathing. These need to be filtered out using hardware and software filters like

high pass filters, low pass filters, notch filters and IIR filters.

102

Several designs for ECG signal acquisition were studied and adopted in designing our circuit, and these can be found in [14], [11], [13], and [12]. 2.2 Effect of Alcohol on Heart Rate Variability The ECG waveform consists of PQRST complexes, which represent the polarization and depolarization of the chambers of the heart. The R peak in the waveform represents the

depolarization of the right and left ventricles of the heart. This is the

85

portion of the PQRST complex that is most striking, and hence can be detected easily. The

time interval between successive RR peaks is known as inter-beat-interval.

49

HRV analysis involves finding variations in these inter- beat-intervals. Studies have shown that alcohol consumption affects

autonomic nervous system activity, which includes the

17

sympathetic and parasympathetic nervous system, which in turn affect the activity of

17

the heart. HRV analysis has been performed on alcoholics for different amounts of alcohol consumption and for time intervals after the subsequent consumption, for males and females, and it was concluded in [1] that HRV decreases for test subjects after ingestion of alcohol, and the effect of decreased HRV was less pronounced in females than it was in males. HRV and its various applications in assessing activities of autonomic nervous system are comprehensively detailed in [2]. In [1], electrocardiograms of 1,000 chronic alcoholic patients were taken and analyzed to find evidence that excessive consumption of alcohol may produce changes in the electrocardiogram. The predominant abnormalities that were observed by the authors were sinus tachycardia and nonspecific T-wave changes. Studies conducted as shown in [22] is one of the first few studies that emphasized the need to study the relation between not the just alcohol consumption and the liver but also consider the effect of alcohol consumption on the heart. Various changes were seen in the ECG of alcoholic patients like the dimple T wave, spinous T wave, cloven T waves, etc. This study clearly establishes the fact that alcohol consumption leads to variation in heart beat. A direct correlation between moderate to heavy alcohol consumption over time and heart rate variability, difference in HRV between the genders, effect on test subjects during exercise, effect of alcohol consumption

on the autonomic nervous system and hence the effect

143

on HRV and ECG signals obtained has been observed and documented in [1], [3], [4], [23], and [5]. 2.3 HRV Analysis and Features Extracted Now that sufficient evidence had been obtained about the effect of alcohol consumption on HRV, the focus shifts towards extracting features from the ECG signal. Several softwares that performed HRV analysis were looked into, which included KubiosHRV, HRVAS (Heart Rate Variability Analysis Software), NerveExpress and IntelleWave, out of which KubiosHRV was chosen for further study since it was available as freeware, was lucid and elaborative. The documentation for KubiosHRV [6] listed out all the methods used for HRV analysis, which are broadly grouped

into time domain methods, frequency domain methods and non-linear methods.

110

The documentation also detailed the features that were extracted and methods of extracting these

methods first involve obtaining the R-R intervals. The frequency domain feature extraction methods involve calculating the power spectral density. Non-linear feature extraction methods utilises Poincare plots, et cetera. Along with these features, in order to obtain better classification accuracies, exogenous autoregressive modelling was also looked into in order to obtain more features, and the system identification toolbox was identified to be an aid in this process, as is described in [16], [11], [17]. 2.4 Classifiers and Similar Implementations In [24] the authors collected ECGs from 50 volunteers. They applied preprocessing for noise suppression and signal segregation

into a number of samples, where each sample represented the heart activity for one heartbeat. To identify characteristics of an alcoholic, features such as Pmax, Pd, means and variances of P, R, S waves and R-R intervals are extracted. The



paper used SVM as a classifier and 10-fold cross validation. This provided sufficient encouragement for us to go ahead with using our dataset and try different classifiers for the same, try to obtain higher accuracies, and implement the real time device. [25] gives a comprehensive view of various supervised machine learning classification techniques and provides interesting domains where machine learning can be applied. In [7], the author describes the architecture of ELM, how it is different from other

neural networks which use back propagation to iteratively tune parameters,

44

and compare it with other classifiers like SVM and Least Squares SVM (LS-SVM). In [18], the authors describe an

algorithm to build the structure of the ELM neural network during the

129

training phase, instead of fixing the architecture a- priori. This would optimise the

number of hidden layer neurons in

40

an intelligent way, instead of using brute force to

find the number of hidden layer neurons that

40

gives best accuracy without over fitting. [8], [9] provide us with an understanding of the Support Vector Machine learning algorithm, and the also talks about Kernels, which allow us to vary the dimensionality of the feature dataset, and the SMO algorithm, which gives us an efficient implementation of SVMs.

CHAPTER â€' 3 METHODOLOGY

88

3.1 INTRODUCTION This section describes the

overall working of the ECG classification project. The

chapter is divided into two sections (Section 3.2, Section 3.3, and Section 3.4),

66

and gives details about the hardware, software, and real time portions of the project. 3.2 HARDWARE This section of the chapter covers topics about the hardware aspects of the project. The idea was to make a portable handheld device that can record a person's ECG signal and detect if the person is a chronic alcoholic. To have sufficient computation power and still be portable, a Raspberry Pi has been used. To the Raspberry Pi, a sensor to amplify and filter the ECG signal is attached. This chapter covers details about the Raspberry Pi, the self-designed ECG sensor and the readymade ECG sensor (AD8232). 3.2.1 ECG Sensor Circuit Design To begin the design of the ECG sensor, first, details about the human physiology were studied. This was a necessary step to figure out where to place the ECG probes and to estimate what amplitude of signal could be expected from the probes. There are several places on the body where the probes can be placed as shown in Fig 3.2.1_1. The requirement of this project was portability and ease of

use. Thus the differential voltage across the left and right index finders was calculated. Once the probe placement was studied and differential signal voltage was estimated, the possible sources of noise in ECG signals were studied. This was done to ensure that the correct filtering specifications were known before starting the actual design of the sensor and filters. Finally, the block diagram level of the sensor design was understood and the complete ECG sensor and filters were designed. Fig 3.2.1_1 Traditional placement of ECG probes From [14] it was noted that ECG signals can be detected only in the milli-volts at the surface of the skin. This meant that an amplifier would be required to bring the signal level to the order of a few volts, so that signal filtering and meaningful signal processing could be performed on the signals for feature extraction. The ECG signal being a low amplitude signal, required an very accurate, low noise amplifier. A suitable such amplifier which was designed for biomedical applications was the AD620. An ECG signal sensor circuit was provided in the datasheet of the AD620. This same circuit was tested on multisim and on obtaining good amplification this amplifier was procured and the hardware circuit was realized on a breadboard. The circuit built is given below (Fig. 3.2.1_2): Fig 3.2.1_2_1 Circuit Design-1 The two stage amplification circuit that was made on the breadboard did not seem to work as a complete unit. The virtual ground and first stage amplification sections of the circuit worked fine, however signals got at the output of the second amplifier were erroneous in shape and amplitude. The HPF present at the junction of the two amplifier was dropped in hopes that proper two stage amplification could be achieved, but that did not solve the issue either. In the process of experimenting with the two stage amplifier, it was seen that the AD620 provided high levels of accurate amplification of signals and a single stage amplifier would be sufficient. Thus, by changing the Rg resistors being supplied to the IC, a single stage amplifier was designed. Rg of 470 ohms was used to obtain a gain of 1000. A gain of 1000 was aimed for, to enable voltages in the mV 3.2.1_1) Some reading up on the purpose and need for virtual ground showed that virtual grounds are used when only a single polarity of voltage supply is available but two polarities of voltage supply is required. The circuit that was planned to be developed needed to be portable and a 9V battery was sufficient to power the amplifiers. It was decided to simplify the circuit further by removing the virtual ground circuit and use two 9V batteries to create a two-polarity source. Now the two-stage amplifier circuit had been reduced to a single stage amplifier and the virtual ground circuit had also been removed. This posed a problem to the filtering that was being done to the signal as the HPF that was present at the junction of the two amplifiers was removed. This required the use of filters at the output of the single stage amplifier. Sources [12] and [13] showed that a HPF and an LPF of 100Hz and 0.05Hz values respectively would be needed at the output of the amplifier. Fig 3.2.1_2_1 Circuit Design-2 The circuit above (fig 3.2.1_2_1) gave output a signal as shown figure (Fig 3.2.1_4) when the probes were connected to the limbs. The signal did not seem to have meaning. However, when the probes were detached, there was a clean signal as seen in Fig 3.2.1_5. It was noticed that this signal had exactly 50Hz of frequency, which is the same frequency as AC sources in Asia. This realization showed that Fig 3.2.1_4 was actually amplified 50Hz power-line noise with the ECG signal being superimposed on it. Fig 3.2.1_3 Superposed Output â€" with Probe Connection to Limbs Fig 3.2.1_4 50Hz Output without Probes Connected to Limbs The goal for the next circuit was to remove the 50Hz signal components and retain on the ECG signal. To do that, a twin T notch filter was designed with variable Q factor. References [19] and [20] were used to design the notch filter. The circuit designed is as given in (Fig 3.2.1_5) and the calculations made to obtain a 50Hz band rejection were as given by (eqn 3.2.1_2). The Q factor of the notch filter was set to around 0.75 by varying the values of R6 and R7 according to (eqn 3.2.1_3). δ "' δ 'oe δ 'j δ 'ce δ 'j δ ' \hat{a} , \hat{Z} = $2\hat{l}$ δ '... δ ¶ 1 (eqn 3.2.1_2) $1\hat{a}$ " $1 = \delta$ '... 7 δ '... $7+\delta$ '...6 (eqn 3.2.1_3) 4ð ',, The Rg used was 100Ω in order to obtain an amplification of about 1000. The ð '... and ð ¶ were set to 3.3ð '`Î⊚ and 1𠜇ð ¹ to obtain a cutoff frequency of 50Hz. Q was set to 0.75 by using R6 and R7 as 1ð "Ω and 2ð "Ω respectively. All modifications resulted in the circuit (Fig 3.2.1_3) below: Fig 3.2.1_5 Circuit Design-3 The above circuit was tested on a breadboard and a pulsating signal of approximately 1Hz was obtained on the CRO. However as the circuit was made on the breadboard, slight disturbances to the wires caused a loss in the output signal. To counter that, the same circuit was soldered onto a breakout breadboard/PCB. Figure (Fig 3.2.1_6) shows the soldered notch filter circuit. Fig 3.2.1_6 Soldered Circuit The entire circuit containing the AD620 and the HPF/LPF could not be soldered within the course of this project. However, in order the serve the greater overall need of the project which was to acquired ECG signals of subjects and analyse the signals for classification purpose, a market ready ECG sensor, the AD8232 was procured and used. Details about the AD8232 and its integration with the Raspberry Pi are given in the section (Section 3.2.5) that follow in this chapter. 3.2.2 Heart Rate Monitor – AD 8232 While a large amount of research had been done on developing a sensor from scratch, it was not a part of the initial scope of the project. While the notch filter showed promising signals on the breadboard for a short duration of time, the soldered version could not be tested fully. To ensure the completion and validation of the alcoholic classification project on the whole, a readymade ECG sensor was procured. Details about the sensor hare mentioned in this section. The Heart Rate Monitor is

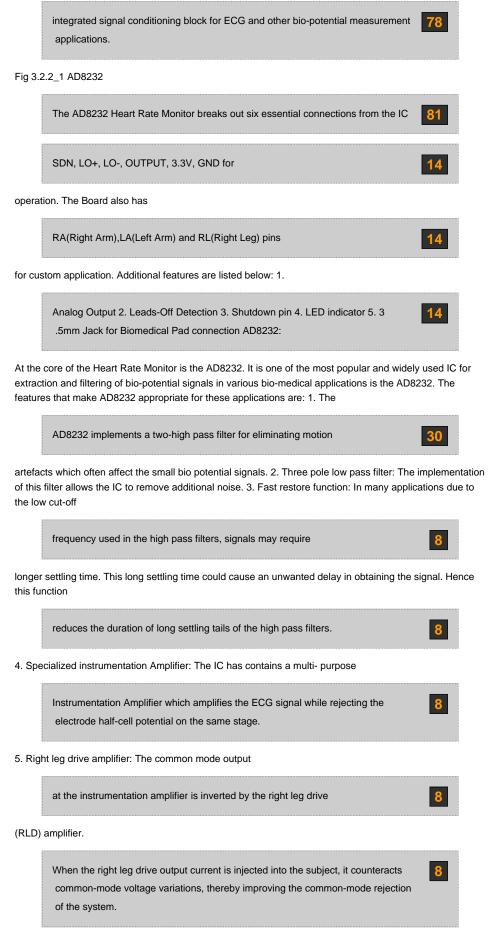
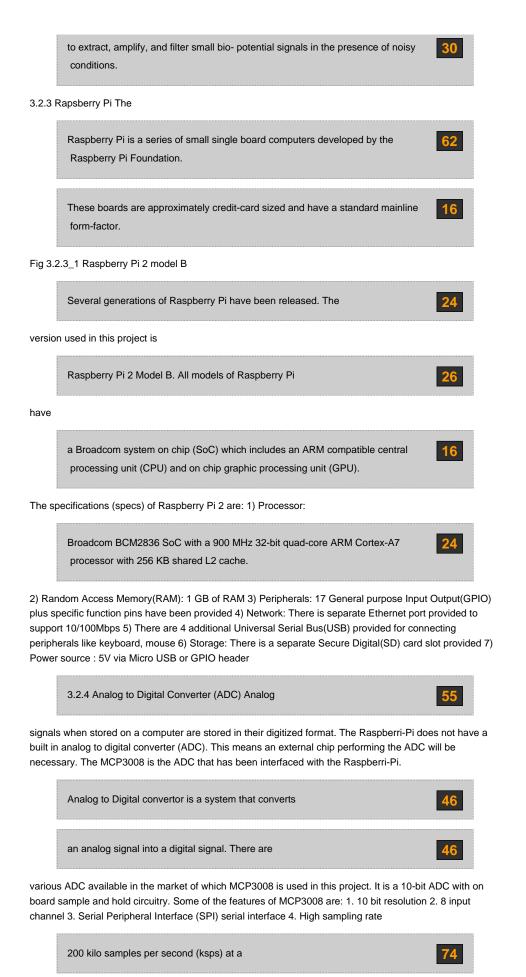


Fig 3.2.2_2 Internal Pin Diagram of AD8232 All the above feature allows AD8232



supply voltage of 3.3V Fig 3.2.4_1 MCP 3008 ADC 3.2.5 Integration of hardware: The previous section gave specifications about the different components used in the hardware portion of the project. This section describes how all of the hardware was integrated. 3.2.5.1 Setting up Raspberry Pi (a) SD card setup Raspberry Pi 2.0 does not come with a pre-built operating system. Instead the operating system has to be flashed on the SD card which is then inserted into the Raspberry Pi. The operating system on SD card is installed by using the following steps: i. ii. SD card is inserted into the SD card reader. Win32DiskImager utility is downloaded from the Sourceforge Project page as an installer file, and the software is installed by running it. iii. The Operating System (OS) of choice is downloaded from official raspberry site. For this project Raspbian Wheezy was used. iv. v. vi. The image of the OS is then extracted from the downloaded file. The Win32DiskImager utility software installed is then opened. The extracted image is then selected and â€Write' button is pressed to write the OS to the SD card. vii. On successful completion of the process the SD card is ejected and inserted

into the Raspberry Pi. a. The Raspberry Pi is

ready for use with Raspbian Wheezy installed. b. ##The Raspberry Pi is then connected to the Internet using Ethernet cable. (b) Internet Protocol (IP) address of Raspberry Pi (R-Pi) In order to

work with the Raspberry Pi the IP address of

Pi is needed. There are two ways to obtain this: i. Working with Rpi using Television (TV) mode: In this method the Rpi is connected to a TV screen using HDMI cable. The Rpi is connected to a router using Ethernet connection for Internet connection. ii. The IP address is then obtained by typing in "ipconfig⠀ in the terminal window of the Rpi. iii. Working with Rpi in "headless†mode: In this case a static IP address is assigned by the user. This is the approach used in this project as it is not always convenient to have a TV screen. The steps to set the static IP address is as follows: (a) SD card is removed from Rpi and then inserted into the SD card reader. (b) The SD card contents are then opened. (c) A file named â €œcmdline.txt†is opened using Notepad++ (d) At the end of the file the following line is appended: Syntax:

ip=<client-ip>:<server-ip>:<gw- ip>:<netmask>:<hostname>:<device>:<autoconf> 79

Example: ip=

169.254.3.14::169.254.56.85:255.255.0.0:rpi:eth0:off (e) The file is then saved and SD card is safely ejected from the system and inserted back into the Raspberry Pi. The Rpi is then connected to a laptop using Ethernet (c) Communicating with the Rpi SSH protocol is used for establishing the communication between the Rpi and the laptop. For this a free and open-source terminal emulator called PuTTY is used.

PuTTY supports many variations on the secure remote terminal, and provides user control over the SSH encryption key and protocol version.

42

3.2.5.2 Connecting Heart Rate Monitor to RPi using ADC MCP3008 The following is the circuit diagram required for the setup: Fig 3.2.5_2 MCP 3008 interface to Raspberry Pi First a 3v3 pin is connected to the positive rail on the breadboard and a

ground pin to the ground rail on the breadboard. The

following connections are made: MCP3008 Rpi VDD (Pin 16) 3.3V VREF (Pin 15) 3.3V AGND (Pin 14) GROUND CLK (Pin 13) GPIO11 (Pin 23/SCLK) DOUT (Pin 12) GPIO9 (Pin 21/MISO) DIN (Pin 11) GPIO10 (Pin 19/MOSI) CS (Pin 10) GPIO8 (Pin 24/CE0) DGND (Pin 9) GROUND Table 3.2.5_1 Connections between MCP3008 and Raspberry Pi 3.2.6 Communication protocols With so many components being used for the hardware, on integrating all of it, some form of communication between all the components was required. Following are the two communication protocols used in this project: 1. Serial-Peripheral Interface (SPI) protocol: This communication protocol was used between the Raspberry-Pi and the MCP3008. SPI is a synchronous data bus



(d) Secure file transfer 3.3 SOFTWARE In this section details about the dataset used, pre-processing performed on the dataset, features extracted from the dataset, feature reduction performed and classification algorithms trained has been provided. While pre-processing and feature reduction were important steps in obtaining a well trained classifier, the bulk of the software end of the project focusses on explaining feature extraction and classification algorithms. Details about the various types of features extracted has been explained in section 3.3.3, and some background information about the two classifiers used on the dataset has been provided in sections 3.3.5 and 3.3.6. 3.3.1 Dataset Description In order to classify test subjects as alcoholics or normative, with a reasonable accuracy, the classifier has to be trained with several samples of the test subjects, and subsequently tested with more samples. These samples along with the appropriate labelling constitute the dataset. The data in the dataset is an array of values which represent the ECG signal of the test subject. The ECG data used has been recorded at the Autonomic Lab, Department of Neurophysiology, NIMHANS, Bengaluru. The data was recorded after taking informed consent adhering to Helsinki's declaration. At the Autonomic Lab , HRV is done using proprietary hardware and software setup by AD instruments, Australia. The product used in the lab premises for observing and recording HRV is PowerLab. This device uses high sample frequency (of the order of 1kHz) to record electronic activity of the heart and other signals pertaining to other functions such as respiratory functions et cetera. The raw ECG data was extracted as five minute samples in European Data Format [EDF]. The dataset comprises of 67 samples, out of which 38 are ECG recordings of alcoholic test subjects and 29 are of normative test subjects. 3.3.2 Pre-processing ECG data acquired from a patient contains various types of disturbances and noise, which makes it difficult to extract features from the signal. The most common types of disturbances are [21]: a. Power-line Interference: ECG signals are measured as the differential voltage that exists between two points on the body and turn out to have very small voltage amplitudes (in the mV range). The small voltage amplitude of ECG signals makes it susceptible to interference from AC signals present at any point in the circuit. The primary source of power-line interference is the AC power-line used as the power source for the ECG recording devices and display monitors or CROs. The frequency of the signal in the power-line in India is 50Hz, leading disturbances on the ECG signals also with the same frequency, b. Baseline Wandering: Gradual changes in the skin impedance of the patient and the patient's breathing lead to low frequency baseline wandering. Baseline wandering is seen as an overall rise and fall of the PQRST complexes of the ECG signal. c. Motion Artefacts: Movements by the patient or the electrode cause mild to severe disturbances in the baseline of the ECG signal. While mild movements like breathing or slight movement of limbs not connected with the probes do not completely â€submerge' the ECG signal, abrupt movements by the patient can completely mask the QRS complex with meaningless noise. In the dataset used, all three forms of artefacts are observable and have been dealt with to obtain cleaner waveforms for feature extraction. Fig 3.3.2_1 Baseline Wandering and Powerline noise in ECG 3.3.2.1 Infinite Impule Respose (IIR) Filtering Different filtering techniques have been used to obtain suitable signals

for time domain, frequency domain and non-linear frequency

109

extractions. Time domain and non-linear feature extraction rely primarily on the RR intervals of the ECG signal. Thus, we focus on filtering out artefacts and other sections of the ECG sequence while maintaining a high amplitude for the QRS complex. Such results have been obtained using

a first order, low pass butterworth filter, having a pass band attenuation of

121

0.2 and cut-off frequencies of 5Hz and 7Hz. The filter was designed for values mentioned previously since, [6] indicates motion artefacts to be present in the 0Hz to 5Hz range. The same filter also helped rid the signal of baseline wandering. Removing the motion artefacts and the baseline wandering essentially ensured that the required R peaks remained the prominent portion of the signal. Finally, the RR intervals were obtained by setting a threshold and measuring the time difference between the occurrences of maximas in those sections of the signal that crossed the threshold. A simple first order IIR filter sufficiently filtered the signal to make the R peaks prominent and allow the calculation of RR intervals. Fig 3.3.2.1_1 PSD of a sample ECG signal with noise components Fig 3.3.2.1_2 IIR Filtered Signal without Baseline Wandering The red line in the above graph represents the threshold that was used to detect the peaks. The threshold value was set at 70% of the total voltage swing occurring in the ECG signal. The same IIR filtered signal was not sufficient clean to accurately extract frequency domain features, as it still contained some low amplitude noise at its baseline. Frequency domain features are obtained primarily by taking the

a clean signal is required. Wavelet Transform has been used to achieve high precision filtering. 3.3.2.2 Wavelet Decomposition One of the most well know signal analysis tools to obtain frequency information of the signal, is the Fourier Analysis. Fourier Analysis breaks the signal down into sinusoids of different frequencies. Fig 3.3.2.2_1 Diagrammatic Representation of Fourier Transform applied to a Signal However, Fourier Analysis has a serious drawback that when a signal is transformed from time domain to frequency domain, the time information is lost. This drawback is not important for stationary signals. But most real world signals contain numerous non-stationary or transitory characteristics: drift, trends, abrupt changes, and beginnings and ends of events. Another analysis method that overcomes the drawback of Fourier Transform is the Short Fourier Transform (STFT). The STFT is a compromise between the time and frequency-based views of a signal. A signal is mapped into two dimensional function of time and frequency. Fig 3.3.2.2_2 Diagrammatic Representation of Short Fourier Transform applied to a Signal The drawbacks of STFT are: (a) Limited precision depending on the size of window (b) Once a particular size for the time window is selected, that window is the same for all frequencies Hence there is a need for an analysis technique that represents a windowing technique with variable-sized regions. Wavelet analysis allows the use of long time intervals where more precise low frequency information is needed, and shorter regions where high frequency information is needed. Wavelet analysis is the breaking up of a signal into shifted and scaled versions of the original wavelet. A wavelet is a waveform of limited duration that has an average value of zero.

over different frequency ranges. To obtain consistent PSD values and accurate frequency domain features,

Fig 3.3.2.2_3 Fourier Transform applied Sine Signal

Depending on the application, different types of wavelet transform tools are used. They are: 1. Continuous Wavelet Transform (CWT) The continuous wavelet transform (CWT) is defined as the sum over all time of the signal multiplied by scaled, shifted versions of the wavelet function \ddot{l} : The output of the CWT are many wavelet coefficients C, which are a function of scale and position. These coefficients on multiplication by scaled and shifted wavelets yield the constituent wavelets of the original signal. Fig 3 .3. 2.2_4 Wavelet Decomposition of a Signal 2. Discrete Wavelet Transform (DWT) The Discrete Wavelet Transform(DWT) is wavelet analysis technique in which the scale and position of the wavelets are varied in powers of two which is called dyadic scales and positions. For calculation of DWT, series of filters are used which give the approximation and detail coefficients which are explained in the next section. The wavelet transform method used in this project is DWT as the results can be more accurately determined by processing less number of data sets. For most real world signal the low frequency content forms the most important part. For example consider human voice. If high frequency component is removed the voice sounds different what is being said can still be understood. Now if low frequency components is removed then only gibberish is heard. Hence it can be said that the high frequency component imparts the flavor to the signal whereas the low frequency component forms the identity of the signal. The DWT of a signal is calculated with the use of series of filters. The approximations (A) are the high-scale, low-frequency components of the

signal. Fig 3.3.2.2_5 Wavelet decomposition into high and low frequencies The

the

signal whereas the details (D) are the low-scale, high-frequency components of

original signal, S, passes through two complementary filters and emerges as two signals. The decomposition process is iterated, with successive approximations being decomposed in turn. In this way the original signal is broken down into many lower-resolution components. This thus leads to a tree formation as shown in the figure below which is called the wavelet decomposition tree. Fig 3. 3.2.2_6 Wavelet Decomposition Tree The detail coefficients (D) consist mainly of the high-frequency noise, while the approximation coefficients (A) contains much less noise than does the original signal. Hence in this way the decomposition coefficients of a signal are obtaine Wavelet transforms also has another concept of the wavelet family and the choice of the mother wavelet is very important. There are various Wavelet Families from which the mother wavelet for analysis is chosen. Some of them are 1. Haar 2. Daubechies 3. Biorthogonal 4. Morlet Fig 3.3.2.2_7 Mother wavelet When an ECG is recorded many kinds on unwanted noise is also recorded with it. These noises cause an alternate shift in baseline of the ECG signal. A process of removing the baseline drift of a signal is called as de-trending. Wavelet transform is used in this project to de-trend the ECG signal that is obtained from the sensor. The Daubechies wavelet of the DWT wavelet family is selected because the shape of the ECG signal and that of db5 is same. Also Daubechies wavelet families are similar in shape to QRS complex and their energy spectrums are concentrated around lower frequencies. Finally, the signal is loaded into Wavelet Analysis and Design Toolbox available in Matlab and the following results were obtained. Fig 3.3.2.2_8 Wavelet Decomposition in different modes Fig 3.3.2.2_9 Decomposition in Tree mode Fig 3.3.2.2_10 Complete Wavelet Decomposition Fig 3.3.2.2_8 shows the decomposition of the signal in separate mode. The right hand side depicts the high frequency decomposition components of the signal whereas the left hand side depicts the low frequency decomposition components of the original signal. The

low frequency approximation of the signal is shown on the right. The

124

complete decomposition of the signal with all high frequency components and the last approximation is seen in fig 3.3.2.2_10. An overall stepwise approach to get complete wavelet decomposition to ECG signals is elaborated from here on.

From the decomposition of ECG signal it is seen that the low frequency component is the cause for baseline



wandering. Hence these components have to be removed from the original signal to get a clean signal which is free from baseline wandering. From the above figures we can see that the low frequency component of the decomposed signal is A8. Therefore this component is subtracted from the original signal to get a de-trended signal. De-trended Signal = Original Signal â€' (A8) Thus in this way

Wavelet Transform is used to remove the baseline wandering



present in the ECG signal. Fig 3.3.2.2_11 Removal of Baseline Wandering by Wavelet Decomposition and RR peak detection The above figure shows the complete steps involved in removing the base line wandering

of the signal using Wavelet transform and detecting the R-peaks. Part



(a) of the figure shows the original ECG signal. Part (b) shows the level8 approximation obtained after Wavelet analysis on the ECG signal. The de- trended signal is shown in part (c). The RR-peaks is detected after detrending the signal as shown by part (d). Another manner in which the results of the wavelet transformed signal was used, was to validate the results of the IIR filtered signals. The peaks of the IIR filtered signal were detected using threshold technique with the threshold set at seventy percent of the total ECG swing. The peaks of the wavelet decomposed ECG signal were also detected, using MATLAB's built in findpeaks() function. Since peak values obtained through IIR filtering and wavelet decomposition matched, each one served as a method to validate the results of the other. 3.3.3

Feature Extraction Feature extraction is a method of



extracting useful information from an otherwise meaningless ECG signal dataset. These features (ie. â €'useful informationâ€[™]) that are extracted from data samples are used by the classifer algorithms to learn how to categorize and classify samples belonging to different classes. Four

types of features have been extracted from each of the



data samples, and they are: a. Time Domain features: This kind of feature extraction works on calculations made on the RR interval sequence. Seven time domain features have been extracted. b. Non-Linear features: This kind of feature extraction also works on calculations made on the RR interval sequence. Here however, the mathematical equations that are used are obtained via graphical analysis. Three time domain features have been extracted. c. Frequency Domain features: This kind of feature extraction applies Power Spectral Density (PSD) on different sections of the signal, to come up with features. Thirteen frequency domain features have been extracted. d. Exogenous Input Auto Regressive (ARX) Model Coefficients: This kind of feature extraction involves breaking up the signal into two halves and using one half of the signal as input and other half of the signal as output to model a system. The coefficient that is calculated for the system can directly be used as features. All the feature extraction operations performed make use of either the IIR filtered signal or the wavelet decomposed signal and not the original ECG. Time domain and non-linear features used IIR filtered signals, whereas the frequency domain and ARX modelled features used the wavelet transformed signal. 3.3.3

.1 Time Domain Time domain feature extraction is the simplest of

59

Standard Deviation of RR intervals (RR_

152

std) This feature is

the standard deviation of the RR interval series and

131

was calculated using $\Tilde{\delta}$ '... $\Tilde{$

Standard Deviation of Heart Rate (HR_std) The standard deviation of the

49

heart rate was calculated in a similar manner as

the mean heart rate. First the standard deviation of

52

the of the frequency of occurrence of the RR intervals per minute was calculated. Then, the inverse was taken to obtain the standard deviation of the heart rate. δ » δ '... δ ' δ ' δ " " δ " δ " " δ " δ " "00 (eqn 3.3.3.1_4) e. Root Mean Square of RR intervals (RR_rms) The

square root of the mean of the sum of the squares of all the

69

entries in the

RR interval series results in the RMS of the

150

RR interval series. The formula for the same is as follows: ð '...ð 'Ÿð 'šð ' = â^šð 'bâ^1 â^'ð 'b'ð '-=â^'01(ð '...ð '...ð '-+1 â^' ð '...ð '-..ð '-)2 1 (eqn 3.3.3.1_5) f. Number of Intervals Varying by Larger than a Threshold (RR_50) This feature is slightly different from the rest, in the sense that it involves an additional step of taking differences. While obtaining the RR interval series required taking successive differences of the time instants at which R peaks occurred, here, successive differences are taken for the values in the RR interval series itself. On this new series of difference, the number of time differences that are larger than 50ms are counted to yield the RR_50 feature. g. Relative Number of Intervals Varying Larger than a Threshold (RR_r50) The previous feature obtained divided by the total length of the the RR interval series gives rise to the final time domain feature. This can be represented by the following equation: \eth '... \eth '... \eth ' $\ddot{\forall}$ 50 = \eth '... \eth ' $\dot{\eth}$ '... \eth '510 \ddot{A} — 100% (eqn 3.3.3.1_6) 3.3.3.2 Non-the following equation: \eth '... \eth '... \eth ' $\ddot{\forall}$ 50 = \eth '... \eth Linear Three non-linear features have been extracted for each of the samples in the dataset. There were two types of analysis done, one of which yielded two features while the other gave rise to a single feature. These two methods are given below: i. Poincare Plot This feature is extracted â€graphically'. A graph of Poincare points is plotted, where the horizontal axis value of all the points is some i-th value in the RR interval series, while the vertical axis value of the point is the (i+1)-th value of the RR interval series. Then, the standard deviations of the points along two different axes are calculated to yield two of three non-linear features denoted by SD1 and SD2 respectively. The axes along which the standard deviations

17

A sample of the Poincare plot obtained for the first alcoholic sample in the dataset has been shown below. Fig 3.3.3.2_1 Poincare Plot ii. Approximate Entropy This feature is essentially a measure of how much irregularity exists within the RR interval series. The following steps explain how the feature value was calculated: a. From the RR interval sequence, a set of

ð ' + ð 'š â^' 1 vectors of length ð 'š are formed. 154
ð ' is the total length of the RR interval series. b. Then the maximum of the

element-wise differences of all the pairs of vectors is found and stored. c. Then a quantity δ \P is calculated that counts the number of distance metrics that were lesser than a threshold δ ' Υ . d. The natural logarithm of the quantity δ \P is calculated e. The sum of all such logarithmic values corresponding to all the vectors is obtained. f. Similar steps are performed

for vectors of length ð 'š + 1

g. The difference between the results for

vectors of length ð 'š and ð 'š + 1

gives the approximate entropy. Exact equations to implement the above steps are mentioned in [6] 3.3

.3.3 Frequency Domain The frequency domain features calculated are obtained from the

order to obtain good PSDs that accurately map to the power contained in the ECG signal, the signal needs

power spectral density (PSD) of the ECG signal. In

to be free of as much noise as possible. This is the reason

wavelet transform is used to filter the signal while obtaining the

frequency domain features. Fig 3.3.3.3_1 PSD of a sample from dataset used Some features are calculated for the whole signal while the rest are calculated only within some frequency sections of the signal. The three main frequency ranges where features are calculated are the

very low frequency (VLF) range, the low frequency (LF) range and the high frequency range (HF). VLF has a range

from 0Hz to 0.04Hz, LF has a range from 0.04Hz to 0.15Hz, and HF is the range of all frequencies above 0.40Hz. The frequency domain features that have been extracted are: a. Peak Frequency in VLF, LF, HF (pk_freq_vlf, pk_freq_lf, pk_freq_hf): The amplitude of the peak frequency values in

very low, low, and high frequency ranges of the PSD of the signal

form the first three frequency domain features. b. Absolute Power in VLF, LF, HF (ab_pow_vlf, ab_pow_lf, ab_pow_hf): The total power contained in the

very low, low and high frequency ranges of the PSD of the signal

form the next three frequency domain features. c. Total Power of the Signal (pw_ttl): The seventh frequency

feature used is the total power contained in the PSD of the signal. d. Relative Power in VLF, LF, HF (rp_lf, rp_lf, rp_hf): The relative power of a certain band is calculated as the power in that band divided by the total power present in the signal. The folmula for VLF is given below (eqn 3.3.3.1_7). The formulae for LF and HF are similar. \eth ' $\check{Z}\eth$ ' $-\check{D}$ ' \eth 'od 'od ' $\check{Z}\eth$ (eqn 3.3.3.1_7) e. Normalized Power in

LF, HF (norm _lf, norm _hf): The normalized power of the LF and HF range is the

absolute power present in that range divided by the difference in the total

power of the signal and the power contained in the

VLF band. The equation for the normalized power in the LF range is given by (eqn 3.3.3.1_8) and that for the HF band is also similar. \eth '' \eth 'œ \eth '' \eth 'oe \eth '' \eth 'os 'oe \eth '' \eth 'os 'oe \eth '' \eth 'os 'oe \eth 'in 'oe \eth '' \eth ''os 'oe \eth 'in 'oe \eth '' \eth ''os 'oe \eth '' \eth ''oe \eth '' \eth ''oe \eth '' \eth ''oe \bullet ''oe

can be represented as: ŏ '¦(ŏ '›) = ŏ 'Ž_ŏ '¦(ŏ '› â^' 1) + ŏ 'Ž_ŏ '¦(ŏ '› â^' 2)

+ ŏ 'Ž_ŏ '¦(ŏ '› â^' ŏ ') + ŏ œ€(ŏ '›)

where ð 'Ž_(ð '~ = 1,2, â€, ð ') are

the model coefficients and $\hat{l}\mu(\,n)$ is a white noise series. In the

condensed form it can be written as: δ ' δ ' $i = \hat{a} \hat{a} \hat{b}$ ' δ '-=1 δ ' ' δ ' "-=1 δ ' ' δ ' " δ " "-=1 δ " ' δ " " δ

δ 'Žδ '-.δ ''δ 'iâ''δ '-+â''δ ' δ '-.δ 'Œδ 'iân'δ '-+ \hat{I} μδ '-=1 δ '-=1

where ð 'Œð '¡ is the input ECG signal After the

ECG signal is free from base line wandering,

the signal is loaded into the System Identification Toolbox in MATLAB to calculate the ARX coefficient. The following steps are used to obtain ARX coefficients for one ECG sample. 1. The System Identification toolbox is opened from the list of apps available or

by typing in the following command "systemIdentification†in the command window

of Matlab.

2. In the System Identification app window, select Import data and then Time domain data.

The

"insert†button is then clicked which adds a new

iii.

can then be viewed by double clicking the model from the model board. Fig 3.3.3.4_3 Coefficients of ARX model fit to required order and input/output signals provided All the information related to the arx model is listed in this window. The model can also be seen on the command line of Matlab by clicking the â €∞Present†button. In this way the ARX coefficients of the system is calculated. These ARX coefficients is used as feature for the classifiers. In conclusion, seven time domain, three non-linear, 13 frequency domain and four or six ARX features (depending on the order of ARX polynomial coefficients) were obtained from each sample. These twenty six features are then fed into the classifiers to train them and obtain the optimal set of weights for classification of any new input data. 3.4 Classifiers Classification is the process of identifying which sub category (or class) a certain sample belongs to. Several classification algorithms have been developed and tested for a number of datasets, for example, Support Vector Machines, Naive Bayes Classifier and Neural Networks. Some classifiers perform better than others for a certain application, or for a specific dataset. There are a number of parameters that are looked into which selecting a certain classifier for a certain application, like accuracy, training time, testing time, etc. Two classifier algorithms have been implemented. One of the classifiers is

the Support Vector Machine (SVM), while the

other is the

Extreme Learning Machine (ELM). This section covers the basic

concepts and ideas behind each of the algorithms used. 3.4

.1 Support Vector Machine A support vector machine (SVM) is a classifier that works on finding

70

a decision boundary that can separate the classes of a dataset. Similar to a two dimensional problem where lines and polynomial curves are used to separate the data belonging to different classes, SVMs use separating surfaces/planes of higher dimensions called hyperplanes. SVMs have a primary hyperplane that behaves as the actual decision boundary for classification, but also uses two other hyperplanes in order to achieve the optimized primary hyperplane. Optimization of the primary hyperplane is done generally solving a Lagrangian dual to the actual geometric equation that needs to be solved. As the number of features that have been extracted for each samples in the dataset is twenty-six, all the data points and hyperplanes exist in the twenty- sixth dimension. This means that twenty-six coordinates exist to describe each data point. A hyperplane in such a dimension is given by (eqn 3.6.1_2), where ð '¤, ð '¥ and ð 'have 26 coordinates (eqn 3.6.1_1) ð '¤1 ð '¥1 ð ' 1 ð '¤2 ð '¥2 ð '¤3 ð '¥3 ð ' 2 â ® â ® ð ' 3 (ð '¤26) (ð '¥26)

ð '¤ð '‡ð '¥+ð ' =0ð '¤ð '‡ð '¥+ð ' =1ð '¤ð '‡ð '¥+ð '

= â^'1

(eqn 3.4.1_1) (eqn 3.4.1_2) (eqn 3.4.1_3) (eqn 3.4.1_4)

If (eqn 3.4.1_2) is considered to be primary hyperplane that takes on the role of being the decision boundary, there exist two other hyperplanes lying on either side of the primary hyperplane given by (eqn 3.4.1 3) and(eqn 3.4.1 4). The purpose of the two adjacent decision boundaries is to aid in arriving at a geometric optimization problem. What needs to be ensured is that the adjacent hyperplanes lie as far away from each other as possible without misclassifying any of the data samples. In other words, the margin between the primary hyperplane and the adjacent hyperplanes needs to be maximized. The margin/distance that exists between the adjacent hyperplanes can be given by (egn 3.4.1_5), and this is the value that needs to be maximized. \eth 'š \eth 'Ž \eth 'Y \eth '" \eth '- \eth '> = 2 || \eth ' \blacksquare | (eqn 3.4.1_5) \eth 'š \eth '- \eth '> || ð '¤||2 2 (eqn 3.4.1_6) Maximizing (eqn 3.4.1_5) is the same as minimizing (eqn 3.4.1_6). Equation (eqn 3.4.1_7) can be solved directly or, as per [8] the Lagrangian's dual can be solved. The Lagrangian dual for (eqn 3.4.1_6) is given by (eqn 3.4.1_7). $\check{\delta}$ $\dot{\xi}(\check{\delta}$ ' \dot{x} , $\check{\delta}$ ' \dot{x}) = $||\check{\delta}$ ' $\dot{x}||^2$ \hat{a} ' \hat{a} ' \hat{a} ' \hat{a} ' \hat{b} ' \hat{x} ' \hat{b} ' ' $\hat{b$ $\hat{a}^{+}\hat{o}^{-} \hat{o}^{-} \hat{s} = 0 \; \hat{o}^{-} \hat{o}^{+} \hat{o}^{-} \hat{o}^{-}$ â^'ð 'šð '-=0 ð >¼ð '-ð 'lð '-ð '¥ð '- (eqn 3.4.1_10) These yields the final Lagrangian optimization equation: $\check{\sigma}$ ' $\check{s}\check{\sigma}$ ' $\check{Z}\check{\sigma}$ ' $\check{Y}\check{\sigma}$ ' $\check{Y}\check{\sigma}$ ' $\check{S}(\check{\sigma}$ ' $\check{Y}\check{\sigma}$) = \hat{a} ' $\check{\sigma}$ ' $\check{s}\check{\sigma}$ '—=0 $\check{\sigma}$) $\check{Y}\check{\sigma}$ '— \hat{a} ' 12 \hat{a} ' $\check{\sigma}$ ' $\check{s}\check{\sigma}$ '—, $\check{\sigma}$ '—=0 ð '¦ð '-ð '¦ð '-ð '¼ð '-ð '¼ð '-ð '¾ð '-ð '¾(ð '¥ð '-, ð '¥ð '-)

(eqn 3.4.1_ 11) In equation (eqn 3.4.1_ 11), ŏ 'š represents the

total number of training samples supplied to the classifier, ŏ '¥ŏ '-/ŏ '
130

stands for the i-th or j-th training sample and ð 'lð '-

/ð '— stands for the class label of the i -th

or j-th sample. \eth 1 /4 \eth 4 is the Lagrangian multiplier for the i-th sample. Lagrangian multiplier values are assigned based on whether they are support vectors or not, and \eth 3 /4 stands for the kernel function that is applied on the input dataset. More information on the application of and need for kernels is given later on in this section. The equation for the decision boundary that is finally obtained is given by (eqn 3.4.1_12) \eth 4 4 4 4 5 4 5 4 5 5 4 5

* SVs = Support vectors are those points of the dataset through which the adjacent hyperplanes pass. The Lagrangian coefficient obtained for scuh data points is nonzero helping increase the training accuracy of the algorithm. Some of the common kernels that are used are the polynomial kernel, Gaussian (also called the

Radial Basis Function (RBF)) kernel and the wavelet kernel.

123

Multiple kernels have been used with the SVM classifier and based on the training accuracy obtained, the Gaussian kernel (given by eqn 3.4.1_13) has been selected. ð ¾(ð '¥ð '-, ð '¥ð '--) = exp (â' || ð '¥ð '-2â''𠜎ð '¥2ð '--||2) (eqn 3.4.1_13) Another important concept about decision boundary type of classifiers like SVMs is regularization. It is common for a dataset to have a few outliers and anomalies. Since the algorithm attempts to classify all the samples correctly, sometimes the algorithm can overfit or go out of the way to include an anomalous sample to the class it had been assigned. As a result, the decision boundary that the algorithm comes up with does not remain generic to new test data that is fed in. Regularization is a method by which some equation parameter/coefficient can be tweaked and tuned to control the algorithm's sensitivity to outliers and anomalies. This control over how closely the algorithm should fit to the training data supplied to the algorithm is very important for real world applications, and has been implemented in the SVM classifier as the parameter †C'. The use of a regularization parameter puts an additional constraint on the Lagrangian coefficients ð 1/4. That is, apart from the constraints given by (eqn 3.4.1_8 to eqn 3.4.1_10), there is an additional constraint as given by (eqn 3.4.1_14) 0<ð →¼<ð ¶ (eqn 3.4.1_14) Fig 3.6.1_1 Importance of regularization A final thought on SVMs is how the SVM, once trained with the weights of the decision boundary, is used to classify a new data sample. This process of classification of some i-th sample after obtaining a trained model is very straightforward. The (eqn 3.4.1_15) is calculated to obtain a number that represents which class the sample belongs to. If the number calculated is positive, then the sample belongs to the first class. In case the number is negative, it is classified to the other class. \check{o} ' \check{o} '£ = \hat{a} ' \check{o} '† \check{o} '-, \check{o} '% \check{o} '—= \check{o} ' 0 \check{o} '† \check{o} '- \check{o} '% \check{o} '— \check{o} '* \check{o} (ð '¥ð '-â€', ð '¥ð '-â€') + ð ' (eqn 3.4.1_15) where ð ' ð '£ is the prediction value and ð '¥ð '-â € and ð '¥ð '—†is the test point needing to be classified into one of the two classes. If the outcome of (eqn 3.4.1_15) is positive, the

test sample is classified as belonging to one class,

28

but if the outcome is negative, the

test sample is classified to the other class.

ð '-ð "" ð ' ð '£ < 0 (egn 3.4.1_15) To summarize all that was explained above and piece together how all the steps fall into place to train an SVM, the following steps have been provided: 1. The input matrix dataset is fed to the algorithm, and all the features are normalized to ensure that all the features contribute equally to the learning algorithm. 2. A combination of regularization parameter ð ¶ and Gaussian variance value 𠜎 are chosen. 3. To this normalized data set, the desired kernel function is applied. If the kernel being used is a Gaussian kernel, then the 𠜎 chosen in the previous step is used. If a different kernel is used that requires no parameter to be tuned, then only regularization parameter ŏ ¶ is used. 4. Now, the SVM is actually â€trained' using some optimization algorithm like Sequential Minimal Optimization (SMO) to solve the Lagrangian dual and obtain a set of weights for the hyperplane. Further reference for the working of the SMO algorithm is provided in [9]. 5. Once the weights for the hyperplane is obtained, the same kernel is applied to the normalized validation dataset which is sent to the trained classifier to calculate the accuracy. 6. Accuracy calculation may be performed either on a single validation dataset, or averaged over multiple folds of data using a technique like k- fold cross validation. For each fold the algorithm is retrained and the weights are recalculated, and validation is performed using the new weights 7. All the above steps are performed for new pairs of values of the regularization parameter ð and Gaussian variance 𠜎, and the pair that yields the best accuracy is chosen to train the final SVM model. Once a trained SVM is present (ie. The weights for the hyperplane giving high training accuracy without being biased and without overfitting have been obtained), classification of a new sample is done as follows: 1. The sample to be classified is normalized using the same method that was used to normalize the original dataset 2. Features are then extracted from the normalized sample and fed to the classifier 3. The classifier then passes this sample through the Gaussian kernel and uses (eqn 3.4.1_16) to get a positive or $\tilde{o} \text{ '} + \tilde{o} \text{ '} - \text{'} + \tilde{o} \text{ '} \text{ (eqn 3.4.1_16) 4. If the expression}$

on the right hand side of (eqn 3. 4.1_16) is

positive it belongs to a particular class, and if negative, it belongs to the other class. As mentioned in the steps on how to train the SVM algorithm, the regularization parameter $\check{\delta}$ ¶ and Gaussian kernel variance $\check{\delta}$ œŽ need to be fixed at the right values to obtain optimal accuracy. There is no real fixed or preferred method that is used to fix these parameters. The method used to fix the parameters in this project has been described here. 1. Two arrays containing the values of $\check{\delta}$ ¶ and $\check{\delta}$ œŽ are initialized. 2. For each value of $\check{\delta}$ ¶ considered, a loop is run 200 (or 400) times 3. In each iteration of the loop, the training dataset is labelled into k-sets using the crossvalind() function. 4. One out of the \hat{a} € k†folds is selected

as the validation set, while the rest are taken as the training data.

5. The model is trained for such a dataset and validated using the validation samples as given by crossvalind(). 6. Similar validation is performed for all of the k-folds to obtain the k-fold cross validation accuracy. 7. An array is maintained to keep track of which 𠜎 values yielded the most accuracy in each of the 200 iterations. 8. Similar steps are performed for all ð ¶ values, and each array is appended to form a matrix. 9. The selection of ð ¶ and 𠜎 that are picked the most in the 200 iteration loop are selected to be the values for which the final model is trained. Crossvalind was found for 25 loops of crossvalind to average the randomness. Fig 3.4.1_2 A count of the C and sigma used for different loops in the case of ARX order 5 features is given. It is clear that C = 0.3 and sigma = 1 is chosen most. 3.4.2

Extreme Learning Machine [ELM] A neural network 45

is a processing unit consisting of sub units called neurons, which are interconnected to each other. These interconnects are assigned weights, representing the acquired knowledge, which may or may not be changed as the classifier is trained using the training samples. A

neural network consists of an input and output layer, along with one or more hidden layers.

In machine learning, if the classifier is being trained using labelled data, that is, the corresponding target output is given for a certain input, it is known as supervised learning. If the classifier is being trained using unlabeled data, and clustering algorithms are required, it is known as unsupervised learning. A feed forward network is one in which the values at the input are propagated towards the output through the hidden layers without being looped back to any preceding layers as an input. Back propagation consists of values that are fed forward through to the output, the error is calculated, and fed back to the preceding layers in order to

test data is used to test the accuracy of the classifier. The percentage of training and testing data is varied in order to prevent under-learning and over-learning and obtain best accuracy. The Input Weights and Biases of Hidden Neurons are generated and assigned randomly. These values are never changed. Let the Input Weights be designated as

W and the Bias as B. The

dimensions of W is

(n,k), where n is the number of hidden layer neurons and k is the dimension of

72

the feature set of sample after passing it through a kernel. The dimensions of B

is (n,p), where p is the number of

140

input training samples given as a batch. If so, each column vector of matrix B will be identical. \check{o} ' α 11 \hat{a} \check{o} ' α 11 \hat{a} \hat{o} ' α 11 \hat{a} \hat{o} ' α 12 \hat{a} \hat{o} ' α 12 \hat{a} \hat{o} ' α 13 \hat{a} \hat{o} ' α 14 \hat{a} \hat{o} ' α 15 \hat{a} ' α 16 \hat{o} ' α 16 \hat{o} ' α 17 \hat{o} ' α 17 \hat{o} ' α 18 \hat{o} ' α 18 \hat{o} ' α 18 \hat{o} ' α 19 \hat{o} ' α 11 \hat{o} α 19 \hat{o} ' α 10 \hat{o} 19 \hat{o} 29 \hat{o} 30 \hat{o} 30 \hat{o} 30 \hat{o} 30 \hat{o} 30 \hat{o} 30 \hat{o}

calculate the output of each neuron in the hidden layer.

108

Examples of Activation functions

include Sigmoid, Sinusoidal, Hard Limit, Triangular Basis function, Radial Basis functions. Htemp is the



input argument to the activation function, which results in H, as shown in (eqn $3.4.2_{-2}$) δ » = δ δ ' δ '

Moore- Penrose pseudo -inverse of H with the targets of training

45

is the index of the maximum value in the output vector

39

given in Fig 3.4.2_2 3.4.3 Validation Validation is an incredibly important part of testing how well a classifier works. Having trained the classifier for a certain portion of the dataset, it is essential to find out how well the classifier performs when exposed to a set of test data it has not seen before. Sometimes, the data that a classifier is trained for might result in the classifier being biased towards a certain class and might perform poorly when given test data that belongs to a different class. So it is important to test the classifier using different test data sets or combinations of data for a classifier which has been trained for a corresponding different combination of training data set, so that upon averaging out the accuracies obtained from each train-test combination, we obtain an overall generalised picture of how the classifier performs in the real world and allows us to prevent over-fitting of the classifier. In cross-validation, the entire dataset is split into subsets of training and testing data, which are complementary to each other, and several rounds of training and testing are done, and results averaged over all the rounds of validation. There are two types of cross validation, namely Exhaustive Validation and Non- Exhaustive Validation. Exhaustive cross-validation consists of Leave-P-Out

Cross-Validation and Leave-One-Out Cross-Validation. In exhaustive cross validation,

all possible combination of samples of data set are chosen and used to train and test the classifier. Non-Exhaustive cross-validation consists of K- Fold Cross-Validation, Hold-Out Method and Repeated Random Sub-Sampling Validation. Non-exhaustive cross validation methods do not include all possible combinations of samples from the dataset for training and testing the classifier. These methods are an approximation of the leave-p-out cross validation method that falls under exhaustive

cross-validation. 3 .4.3. 1 Leave- One -Out Cross-Validation

This is a type of exhaustive cross-validation and an explicit example of leave-p- out cross-validation for p=1, in which one sample from the dataset is chosen

for testing and all other samples are chosen to train the classifier

per iteration, and this procedure is iterated for as many times as there are samples in the dataset, only that the sample chosen for testing is different every single iteration. The accuracies over every single iteration are averaged over and an average accuracy is obtained. This method is better than leave-p-out cross-validation in that the number of combinations of train-test data are numerically

equal to the number of samples in the dataset. For

example, in a dataset having 100 samples, in each iteration, 1 sample is

chosen as the test data while the other 99 samples are chosen as the

training data. 3

.4.3.2 K-Fold Cross-Validation This is a type of

non-exhaustive cross-validation in which the

dataset is randomly partitioned into k number of equally sized subsets,

and in each of k number of iterations, one of the subset is chosen as the test data set while the rest of the k-1 subsets constitute the training data set, and the subset chosen to test and the subsets chosen to train the classifier are unique in every iteration of training and testing. The accuracies obtained in each kth "foldâ er averaged out to obtain an average accuracy that gives us an idea about the performance of the classifier. The advantage of this method of cross validation is that every single subset is used for both training and testing and a certain subset is used for testing only one time. The value of k determines the percentage of the dataset that is used for training and testing. For example, if k=4, for a dataset having 100 samples, the dataset is split into 4 subsets of 25 samples each. In each iteration, 75



This procedure happens 3 more times for the other 3 subsets of 25 samples, while the rest of the 75 samples in each fold participate as the training set. The accuracies over the 4 folds are averaged and presented as the accuracy of the classifier. When k equals the number of samples in the dataset, it becomes nothing but leave-one-out cross-validation. 3

.4.3. 3 Confusion Matrix A confusion matrix is a table of



values which provides us with a way to visualise the performance of an algorithm, or in the case of pattern classification problems, the performance of a classifier. The rows represent instances of the

actual class while the columns represent instances of the predicted class.

141

From this matrix, we can get to know how many times the classifier is classifying the sample correctly and and how many times a certain class of sample is misclassified as another class, and the number for each class. The table of confusion (also called the confusion matrix) is one consisting of 2 rows and 2 columns. The 4 elements of this matrix give us

the number of True Positives, False Negatives, False Positives and True Negatives.



True positives are those samples which were classified correctly for a specific class. False negatives are those in which the actual label was that of the class we are calculating the matrix for, and the classifier predicted wrongly. False positives are those in which the predicted label was that of the class for which we were finding the matrix for, but the actual label was that of another class. True negatives are those in which all the other classes apart from the class we were calculating the matrix for, were classified correctly. Total Population Prediction Positive Prediction

Negative Condition Positive True Positive False Negative Condition Negative False Positive True Negative



Table 3.4.3.3_1 Confusion Matrix Sensitivity is a measure of the

proportion of positives that are correctly classified as

153

positives. It is also known as recall, hit rate or true positive rate. δ '† δ " δ ' δ ' δ ' δ ' δ '† δ '† δ '† δ '† δ ' δ ' δ ' δ '† δ ' δ '† δ '† δ ' δ ' δ '† δ '† δ ' δ " (eqn 3.4.3.3_1) δ '‡ δ '† δ '¢ δ " (eqn 3.4.3.3_1) δ '‡ δ '† δ '¢ δ " (eqn 3.4.3.3_1) δ '‡ δ '† δ '¢ δ " (eqn 3.4.3.3_1) δ '‡ δ '† δ '† δ '° δ " (eqn 3.4.3.3_1) δ '‡ δ '† δ '† δ '† δ " δ " (eqn 3.4.3.3_1) δ '‡ δ '† δ '† δ '† δ " (eqn 3.4.3.3_1) δ '† δ " δ " (eqn 3.4.3.3_2) δ '† δ " δ " (eqn 3.4.3.3_2) δ '† δ " (eqn 3.4.3.3_2) δ

the confusion matrix and from that, the sensitivity and specificity

29

is that it allows us to find biases in the system if there exist any. For example, if it seen that the sensitivity of a system is low but the specificity is high, it would mean that the decision hyperplane (in the case of SVM) or the neural connections (in the case of ELM) might be biased toward the negative class. Getting to know such information would help tweak parameters of the system to correct it from being biased toward the negative class. 3.5 REAL TIME APPLICATION: As the title suggest "A real time application to classify alcoholics from ECG Signals†, the ECG signal was captured in real time using the AD8232 heart rate monitor and Raspberry pi, features were extracted and the signal was classified. The issue of absence of an analog pin on Raspberry pi was removed by the use of an ADC which is the MACP3008. The problem of mismatch of the sampling rate of the ADC and the Raspberry Pi was removed by the use of time library in Python which ensured that the Raspberry Pi read values every 1millisecond. The below image shows the ECG signal in real time on the CRO. Fig 3.5_1 Real time capture of ECG

easy demonstration of the real time application built here. The module used for GUI creation in python is know as Tkinter. All the above mentioned processing part of feature extraction and classification was implemented on Raspberry Pi. 3.5.1 Tkinter: Tkinter is the Python's application programming interface(API) for Tk GUI toolkit library. Tk is a free and open source library used for creating a GUI across many programming languages. 3.5.2 Graphical User Interface (GUI) The image below shows a GUI that is built for this project. Fig 3.5.2_1 GUI The first row of the GUI is to implement the offline samples available. It has buttons to load the signal, extract features and also to classify it based on the features extracted. The second row of the GUI to implement a real time process wherein the user will click the "Start Signal Capture†and the signal will be captured for a given interval of time. On clicking "Extract features†button the features will be extracted from the signal and will be further used for classification. All the required codes for signal capture, feature extraction and classifier are written in Python so that it can be easily integrated with GUI which is also written in Python. The snapshots below show the entire process which is Signal capture, feature extraction and classification: Fig 3.5.2_2 ECG signal loading process is started Fig 3.5.2_3 ECG signal is loaded Fig 3.5.2_4 Feature extraction completed Fig 3.5.2_5

This chapter is dedicated to describing the results obtained by the analyses performed in the project. For all the results, 28 alcoholic and 28 normative samples were used to train the system. Different features sets were used to train the classifier. A novel set of features that was used 4.1 Results of SVM The process of selecting the ð ¶ and 𠜎 pair, training the SVM, obtaining the k- fold validation accuracy have been mentioned in Chapter 3. Here, the results obtained for specific groups of feature sets used and the corresponding confusion matrices has been provided. The accuracies and optimal ð ¶ and 𠜎 pair obtained for the different types of feature sets are as given the table below. Features Used Time, non-linear, frequency Optimal (ð ¹a, ð ˆ) Pair (0.1, 0.3) 8-Fold Accuracy 80% Time, non-linear, frequency, ARX order 3 (0.3, 1) 82% Time, non-linear, frequency, ARX order 5 (0.3, 1) 86% Table 4.1_1 Results of SVM for different feature sets

From the table above, it is seen clearly that the accuracy of the

48

system improves when ARX coefficients are used along with

the time domain, non-linear and frequency domain

61

features. It is also important to understand how many samples of each class were classified correctly. For this the confusion matrix was obtained and the sensitivity and specificity (eqn 3.4.3.3_1 and eqn 3.4.3.3_2) were calculated for each of the cases given above in (Table 4.1_2). Features Used Time, non-linear, frequency, ARX order 5 Confusion Matrix 25 3 (Sensitivity, Specificity) (89%, 82%) 5 23 Table 4.1_2 Confusion matrix for SVM with ARX features The table above shows that the system is very slightly biased toward the positive class since a few more samples in the negative class are being misclassified. However, this is not of much concern as both sensitivity and specificity are well above 80% accuracy. 4.2 Results of ELM The process of selecting the

number of neurons in the hidden layer and 𠜎, training the ELM, obtaining the

10

k-fold validation accuracy have been mentioned in Chapter 3. Here, the results obtained for specific groups of feature sets used and the corresponding confusion matrices has been provided. The accuracies and optimal

number of hidden layer neurons and 𠜎 obtained for the different types of

127

feature sets are as given the table below. Features Used Time, non-linear, frequency Optimal (Number 7-Fold Accuracy of Hidden Layer Neurons, ŏ ^) (6, 1.4) 89.29% Time, non-linear, frequency, ARX order 3 (7, 2) 92.86% Time, non-linear, frequency, ARX order 5 (15, 1.8) 94.64% Table 4.2_1 Accuracies of ELM for different feature sets Features Used Optimal (Number Leave-One-Out of Hidden Layer Validation Accuracy Neurons, ŏ ^) Time, non-linear, (6, 2) 92.86% frequency, ARX order 5 Table 4.2_2 Accuracy of ELM for Leave One Out Validation with ARX order 5

system improves when ARX coefficients are used along with

the time domain, non-linear and frequency domain

61

features. It is also important to understand how many samples of each class were classified correctly. For this the confusion matrix was obtained and the sensitivity and specificity (eqn 3.4.3.3_1 and eqn 3.4.3.3_2) were calculated for each of the cases given above in (Table 4.2_3). Features Used Time, non-linear, frequency, ARX order 5 Confusion Matrix 26 2 (Sensitivity, Specificity) (92.86%, 85.71%) 4 24 Table 4.2_3 Confusion matrix for ELM with ARX features order 5 The time above shows that the system is very slightly biased toward the positive class since a few more samples in the negative class are being misclassified. However, this is not of much concern as both sensitivity and specificity are well above 85% accuracy. 4.3 Comparative Results and Points of Discussion Accuracies obtained for k-fold cross validation for SVM and ELM are compared in Table 4.3_1 and the sensitivity and specificity obtained for SVM and ELM are given in Table 4.3_2. Without ARX Coefficients SVM 80% ELM 89% With ARX Coefficients of order 5 86% 94% Table 4.3_1 Comparative results of SVM and ELM Sensitivity SVM 89% ELM 92.86% Specificity 82% 85.71% Table 4.3_2 Comparative Sensitivity and Specificity of SVM and ELM Points of Discussion? Classifiers can be applied to features extracted via HRV analysis and good accuracies are obtained. ? ELM classifier was found to have a better accuracy than SVM for the same feature set. ? ARX features improve the accuracy of both ELM and SVM. CONCLUSION AND FUTURE WORK

Summary: There was a lot of learning and knowledge gained by the end of this project. First, the ECG signal and it's capture by building a hardware circuit was explored and understood. Then, study of papers led to exposure toward the physiological effects of alcohol on the body and the change in heart rate variability due to alcoholic consumption. After this, various techniques to extract features were explored and implemented. Feature extraction also required reading up on FIR filter, IIR filters and Wavelet Transforms. This was followed with exploring classifiers like the

Support Vector Machine and Extreme Learning Machine which were to be applied to the

82

features extracted to classify samples. Finally, signal sampling, ADC interface and sensor interface to the Raspberry Pi was implemented to create a Real Time system with a GUI. At the end of this project it could be seen that meaningful features could be extracted from HRV analysis applied to the ECG signals, and that these features provided good accuracies for both classifiers. It was also seen that ELM performed better than SVM by around five percent of training accuracy. Another crucial point that could be drawn from this project is the use of ARX coefficient as features that improved the accuracy of both the algorithms substantially. Future Work: The real time application developed was tested for signals recorded for normative subjects only. To verify the complete working of the real time system, the ECG signal of an alcoholic subject needs to be obtained and classified. Currently, the ELM algorithm is optimized by †brute force' through multiple trial and error loops. The plan ahead, is to prevent this randomness and utilize a technique called Meta-Cognitive Learning to ELM to make it learn more intelligently and accurately the first time around itself. Another point for future work to further improve the accuracy of the classifiers, is to use non- linear ARX model coefficients. The final goal of the project is to be able to come up with an algorithm that is able to classify subjects into multiple classes based on the level of alcohol intake. REFERENCES

[1] Ping Shi, Ying Chen,

Ming-Ming Guo and Hong-Liu Yu, "Acute Effects Of Alcohol On Heart Rate Variability: Time-Related Changes And Gender Difference†, Biomedical Engineering: Applications, Basis and Communications,Vol. 26, No. 3 (2014) 1450048 (10 pages) [2] U. Rajendra Acharya, K. Paul Joseph, N. Kannathal, Choo Min Lim, Jasjit S. Suri, "Heart rate variability: a review†, Med Bio Eng Comput (2006) 44:1031â€'1051 [3] Kusuma Ramanna1, Fazal M Gahlot2, Nagaraja Puranik1, â €œElectrocardiogram changes and heart rate variability during moderate exercise in chronic alcoholics†, International Journal of Medical Science and Public Health Vol 4, Issue 4 (2015) pp. 492-495 [4] Jon T. Ingjaldsson, Jon C. Laberg, and Julian F. Thayer, "Reduced Heart Rate Variability in Chronic Alcohol Abuse: Relationship with Negative Mood, Chronic Thought Suppression, and Compulsive Drinking†, Society of Biological Psychiatry, (2002), pp. 1427-1436 [5] Katsuyuki Murata 1,2 Philip J. Landrigan 2 and Shunichi Araki, "Effects of age, heart rate, gender, tobacco and alcohol ingestion on R-R interval variability in human ECG†, Journal of the Autonomic Nervous System, 37 (1992) pp.199-206 [6] Mika P. Tarvainen and Juha-Pekka Niskanen, "Kubios HRV Analysis version 2.0 beta USER'S GUIDE†, Biosignal Analysis and Medical Imaging Group, Department of Physics, University of Kuopio, Finland [7] G.-B. Huang, "What are Extreme Learning Machines? Filling the Gap between Frank Rosenblatt's Dream

and John von Neumann's Puzzle,†Cognitive Computation, vol. 7, pp. 263-278, 2015. [8] Andrew Ng, â €œSupport Vector Machinesâ€, 2011. [Online] Available: http://cs229.stanford.edu/notes/cs229-notes3.pdf Accessed: 10-Feb-2016 [9] Andrew Ng, "The Simplified SMO Algorithmâ€, 2012. [Online] Available: http://cs229.stanford.edu/materials/smo.pdf Accessed: 10-Feb-2016 [10] Branislav Vuksanovic & Mustafa Alhamdi, "AR-based Method for ECG Classification and Patient Recognition,†International Journal of Biometrics and Bioinformatics (IJBB), vol. 7 ,Issue 2, 2013 [11] Dingfei Ge, "Cardiac arrhythmia classification using autoregressive modelling†. [Online] Available: http://biomedical-engineeringonline.biomedcentral.com/articles/10.1186/1475-925X-1-5 [12] Pico Technoology, "Electrocardiogram (ECG) circuit for use with oscilloscopesâ€. [Online] Available: https://www.picotech.com/library/application-note/electrocardiogram-ecg- circuit-for-use-with-oscilloscopes [13] Lv Jinhua and Xu Yanyi, "Circuit Design for Front-End Electrocardiographâ€, International Journal of Multimedia and Ubiquitous Engineering Vol.11, No.5 (2016), pp.345-354 [14] Ajay Bharadwaj and Umanath Kamath, Cypress Semiconductor Corp., "Techniques for accurate ECG signal processingâ€ [Online] Available: http://www.eetimes.com/document.asp?doc_id=1278571 [15] NTHU, "ECG Circuits, Signal Sampling and Digitalization†[Online] Available: https://www.google.co.in/url? sa=t&rct=j&q=&esrc=s&source=web&cd=1&cad =rja&uact=8&ved=0ahUKEwins5jwg43TAhWCq48KHcxPAysQFgggMAA&u rl=http://lms.nthu.edu.tw/sys/read_attach.php?id=69 2351&usg=AFQjCNFJqOwwhxZZG0FP1yQNMuNpHBV73Q&sig2=e_- aLgXrLbFMIB-XhTd2Kg [16] L. Ljung,"System identification toolbox,†The Matlab user's guide, 2012. [Online] Available: http://radio.feld.cvut.cz/matlab/pdf_doc/ident/ident.pdf [17] Hong He, Xiaowen Yan and Wei Wei, â €ceMeridian ECG Information Transmission System Modeling Using NARX Neural Network†,IEEE/ACIS 15th International Conference on Computer and Information Science (ICIS), 2016. [Online] Available:http://ieeexplore.ieee.org/document/7550775/ [18] R. Savitha, S. Suresh, H.J. Kim, "A Meta-Cognitive Learning Algorithm for an Extreme Learning Machine Classifier†Cognitive Computation, vol. 6, pp. 253-263, 2014. [19] Ian Poole, "Opamp Notch Filter Circuit†[Online] Available: http://www.radioelectronics.com/info/circuits/opamp_notch_filter/opamp_notch_filter.php [20] "Band Stop Filterâ€ [Online] Available: http://www.electronics- tutorials.ws/filter/band-stop-filter.html [21] Mr. Hrishikesh Limaye1, Mrs. V.V. Deshmukh2, "ECG Noise Sources and Various Noise Removal Techniques: A Survey†, International Journal of Application or Innovation in Engineering & Management, Volume 5, Issue 2, (2016) [22] William Evans, "The Electrocardiogram of Alcoholic Cardiomyopathyâ€, British Heart Journal, 21(4), (Oct. 1959): pp.445-456 [23] Phyllis K. Stein, et. al., "Heart Rate Variability and Measure of Autonomic Tone†, American Heart Journal, vol. 127 no. 5 (Sept. 1993) pp. 1376-1381 [24] Chung Kit Wu, et. al. "A Precise Drunk Driving Detection Using Weighted Kernel based on Electrocardiogramâ€, Sensors. [Online]. 16(5), pp659. Available: http://www.mdpi.com/1424-8220/16/5/659/htm [25] Kotsianntis, Sotiris B., et. al. "Supervised Machine Learning: A review of classification techniques†3-24, (2007) APPENDIX â€' A Datasheets: APPENDIX â € B Akarsh N Kolekar (1PI13EC009) Hardware: 1. Selection of low noise, high gain, Instrumentation Amplifier IC 2. Study of Circuit â€' 1 (Two stage Op-Amp Circuit) 3. Re-soldering of Notch Circuit Design â€' 3 Software: 1. Implementation of ELM Classifier (MATLAB and Python) 2. Study of Meta Cognitive ELM 3. Python GUI for Real Time System APPENDIX â€" C Apoorv Vatsal (1PI13EC017) _ Hardware: 1. Set-Up of Raspberry Pi 2. Interface of ADC MCP 3008 3. Interface of AD8232 with Raspberry Pi 4. Implemented signal capture code to save as CSV file on Raspberry Pi Software: 1. Implementation of Wavelet Decomposition for pre-processing 2. Implementation of Frequency Feature Extraction (MATLAB and Python) 3. Implementation of Autoregressive Modelling with Exogenous Inputs 4. Python GUI for Real Time System APPENDIX â€" D Rakshith Vishwanatha (1PI13EC075) Hardware: 1. Study of LPF, HPF filter ranges and Q-Factor for band reject filters 2. Design of Circuit â€" 3 (Notch Filter Circuit) 3. Soldering of Notch Circuit Design †3 and MCP 3008 4. Solved 1kHz sampling issue on Raspberry Pi Software: 1. Implementation of IIR filtering for pre-processing 2. Implementation of Time Domain Feature Extraction (MATLAB and Python) 3. Implementation of Non-Linear Feature Extraction (MATLAB) 4. Implementation of SVM classifier A Real Time Application to Identify Alcoholics from ECG Signals Dept. of ECE A Real Time Application to Identify

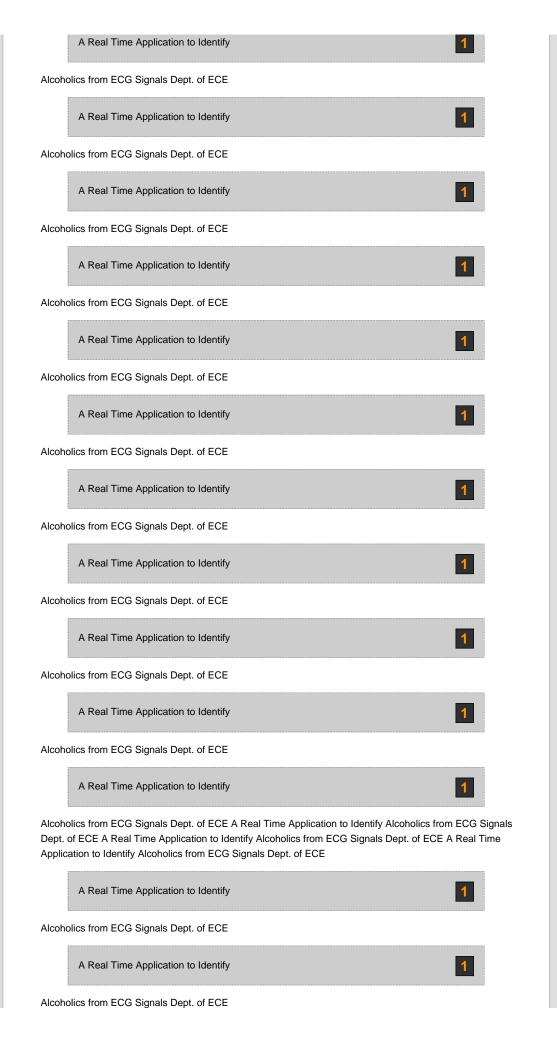
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE

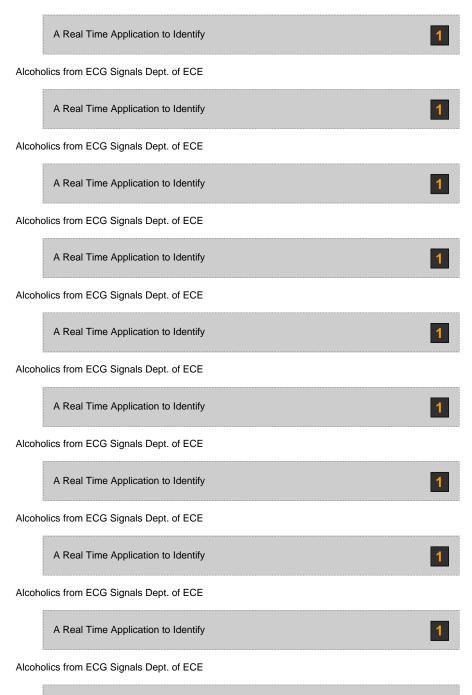
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	

A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	1
Alcoholics from ECG Signals Dept. of ECE	

A Real Time Application to Identify	
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	
Alcoholics from ECG Signals Dept. of ECE	!
A Real Time Application to Identify	
Alcoholics from ECG Signals Dept. of ECE	!
A Real Time Application to Identify	
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	
Alcoholics from ECG Signals Dept. of ECE	;
A Real Time Application to Identify	
Alcoholics from ECG Signals Dept. of ECE	;
A Real Time Application to Identify	
Alcoholics from ECG Signals Dept. of ECE	;
A Real Time Application to Identify	
Alcoholics from ECG Signals Dept. of ECE	;
A Real Time Application to Identify	
Alcoholics from ECG Signals Dept. of ECE	
A Real Time Application to Identify	
Alcoholics from ECG Signals Dept. of ECE	

A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE
A Real Time Application to Identify
Alcoholics from ECG Signals Dept. of ECE





Page 1 Page 2 Page 3 Page 4 Page 5 Page 6 Page 7 Page 8 Page 9 Page 10

Page 11 Page 12 Page 13 Page 14 Page 15 Page 16 Page 17 Page 18 Page 19

Page 20 Page 21 Page 22 Page 23 Page 24 Page 25 Page 26 Page 27 Page 28 Page 29 Page 30 Page 31 Page 32 Page 33 Page 34 Page 35 Page 36 Page 37 Page 38

Page 39 Page 40 Page 41 Page 42 Page 43 Page 44 Page 45 Page 46 Page 47 Page 48 Page 49 Page 50 Page 51 Page 52 Page 53 Page 54 Page 55 Page 56 Page 57

Page 58 Page 59 Page 60 Page 61 Page 62 Page 63 Page 64 Page 65 Page 66 Page 67 Page 68 Page 69 Page 70 Page 71 Page 72 Page 73 Page 74 Page 75 Page 76

Page 77 Page 78 Page 79 Page 80 Page 81 Page 82 Page 83 Page 84 Page 85 Page 86 Page 87 Page 88 Page 89 Page 90 Page 91 Page 92