Imperial College London

Department of Electrical and Electronic Engineering

Final Year Project Final Report 2022



Project Title: Cuffless Estimation of Blood Pressure from Photoplethysmog-

raphy Signals using Transformers

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Final Report Plagiarism Statement

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Abstract (CHANGE AT END)

Abstract should consist of motivation, methods, results, conclusion. All concise & capturing the reader within 200 words. Your abstract needs working and finalising once you have your results and concluding statement.

The prevention, evaluation, and treatment of hypertension have attracted increasing attention in recent years. The advancement of wearable technology has resulted in increasing importance into the monitoring of non-invasive ambulatory blood pressure, compared to the traditional invasive blood-pressure monitoring methods. As photoplethysmography (PPG) technology has been widely applied to wearable sensors, the noninvasive estimation of blood pressure (BP) using the PPG method has received considerable interest. For this project, systolic and diastolic BPs are estimated using PPG signals. A Recurrent Neural Network (RNN) is used for estimation. Due to their being several alternative existing methods for estimating blood pressure, it was necessary to perform a comparison between the best performing Deep Learning based methods. Overall, the proposed method obtains better accuracy. The model achieves a mean absolute error of mmHg for systolic BP and mmHg for diastolic BP.

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1 Introduction

1.1 Motivation

Cardiovascular disease is one of the main causes of death around the world. High blood pressure (BP), which is also known as hypertension, is a common condition which can be a cause of cardiovascular disease [1]. According to the World Health Organization (WHO), the mortality rate due to hypertension is 9.4 million per year and it causes 55.3% of total deaths in cardiovascular patients [2]. If hypertension is detected early and prevented, this will greatly lower the number of deaths associated with cardiovascular diseases [2].

Recent developments in technology have made wearable sensors, such as Electrocardiogram (ECG) and Photoplethysmography (PPG) sensors significantly more popular in today's world. These sensors provide real-time 24 hour monitoring of the human bodily function. Hence there is great potential in using these sensors to diagnose medical conditions, such as hypertension, in real-time, thus helping to save lives [3]. Ambulatory BP monitoring is seen as a promising method for detecting early symptoms of hypertension [4]. There is a lot of existing research to predict ambulatory BP using methods which are cuff-less, continuous and non-invasive [5]. Hence wearables are seen as a viable option for this. ECG and PPG sensors have been discovered to be a potential estimator of blood pressure that cause minimal harm to patients compared to existing cuff-based methods [6] [7].

1.2 High-level problem statement

Needs rewriting. Aim: Your intention/what you hope to achieve.

Objectives: Statements of measurable outcomes/What will you be doing to achieve the aim/desire outcome. (this is the work you're going to do) The aim is not to implement and evaluate different techniques. The aim is to estimate cuffless BP using PPG for wearable technology purpose. I advise you to have a clear aim and clear objectives. Objectives can be listed as bullets or numerated. These objectives will reappear in your conclusion where you state if you completed them or not.

Based on the provided motivation, the main aim of this Final Year Project (FYP) is to estimate cuff-less blood pressure values using ECG and/or PPG signals, so that this estimation process can be integrated onto future wearable technology devices. Hence, these are the following objectives for this Final Year Project (FYP):

- Conduct a literature review to assess what are the most popular methods for estimating cuff-less BP values and to decide on the most feasible implementation for this FYP
- Develop a novel algorithm in the Python programming language to estimate cuffless blood pressure
- Assess the performance of this algorithm against existing methods
- Conclude whether this method is feasible for future wearable technology products

1.3 Overview of work

This section provides a chronological overview of the work that will be done in this FYP. The comprehensive overview of the work is provided in the Gantt Chart in the Appendix.

- 1.3.1 Autumn Term 2021
- 1.3.2 Spring Term 2022
- 1.3.3 Summer Term 2022

2 Background

In this chapter, the aim is to provide sufficient background information, in order to understand how the cuff-less estimation of blood pressure (BP) values can be achieved. An overview on all necessary fields will first be given (denoted by Subchapter) and then a literature review will be conducted to assess what is the most feasible implementation for BP cuff-less estimation for this FYP.

Add an intro to the background, rather than jumping into the lit review. Set the scene for the reader.

2.1 Medical background

In this chapter, all of the medical knowledge required to understand the basis of this project will be discussed.

2.1.1 Hypertension

The heart can suffer from a variety of diseases and pathologies. Low blood pressure, or hypotension, has the potential to cause a lack of oxygen flowing to the brain and other organs, causing shock [8]. Whilst hypotension is a serious issue, hypertension has been identified by the World Health Organization (WHO) as the most significant risk factor for cardiovascular diseases [9]. According to the 2017 American Heart Association guidelines for hypertension, the risk of developing stage two hypertension, ≥ 140 mmHg systolic or ≥ 90 mmHg diastolic is almost 90% [7] (see Table 1). Over 20% of adults have hypertension and its complications cause a major number of diseases, including heart attacks, strokes and heart failure. If hypertension is not diagnosed and properly treated it can even cause death [2].

Hypertension or high blood pressure (BP) is where blood continues to exert more and more pressure on the arterial walls. One particular disease linked to hypertension is hypertrophic cardiomyopathy, as indicated in Figure 1,

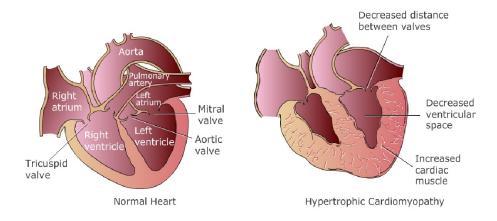


Figure 1: The effects of hypertension on the heart [10]

Hence it is clear that hypertension is one of the largest motivating factors for this project.

Table 1: Categories of blood pressure in adults [9] [11]

| Blood pressure classification | Blood Pre | essure (mmHg) |
|--------------------------------|------------|---------------|
| Blood pressure classification | Systolic | Diastolic |
| Hypotension | ≤ 90 | $Or \le 60$ |
| Normal | 90-119 | And 60-79 |
| Prehypertension | 120-139 | Or 80-89 |
| Stage 1 hypertension | 140-159 | Or 90-99 |
| Stage 2 hypertension | ≥ 160 | $Or \ge 100$ |
| Isolated Systolic hypertension | ≥ 140 | And < 90 |
| Hypertensive crisis | ≥ 180 | $Or \ge 110$ |

2.1.2 Blood Pressure measurements

Blood pressure (BP) is the force of the blood pushing against the arterial walls as the heart pumps blood. It is measured in millimeters of mercury (mmHg) [3]. BP is measured in terms of systolic blood pressure (SBP) and diastolic blood pressure (DBP). These values are the maximum and minimum pressure values of an Arterial Blood Pressure waveform during a cardiac cycle respectively [11] [12]. An example of this structure is provided in Figure 2.

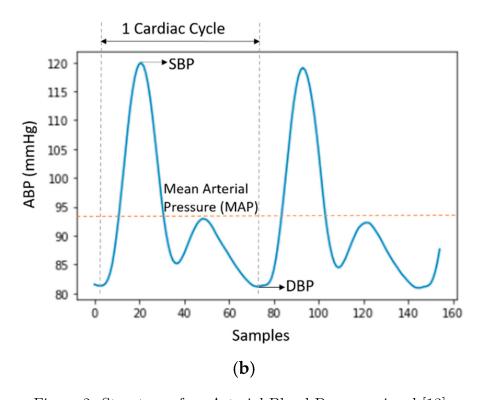


Figure 2: Structure of an Arterial Blood Pressure signal [13]

As shown in Figure 2, the Systolic and Diastolic Blood Pressure values of the waveform are defined by the maximum and minimum ABP values within the provided sampling window.

BP has oscillations or pulses that mirror the oscillatory nature of the heart. The blood is

propelled during systole, also known as heart contraction, and the blood is rested during diastole, known as heart relaxation, as illustrated in Figure 3.

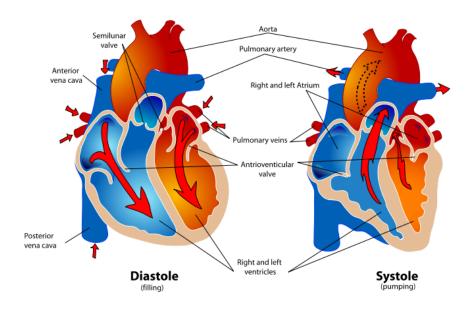


Figure 3: Visualisation for SBP and DBP [14]

There are two conventional methods for measuring blood pressure (BP). These are invasive and non-invasive methods.

If you're going to use firstly, there should be consistency. I.e where is secondly or finally? Check throughout report. Firstly, a description on invasive methods. The most popular form of invasive BP measurement is catheterization [5]. Invasive BP measurements are continuous and the most accurate from heartbeat to heartbeat. As a result these measurements are recognised as the gold standard internationally [1] [15]. However, this method is usually restricted to hospitals, as medical supervision is required [12]. In addition, this method poses several health risks, including bleeding and infection. As a result, invasive measurements are only utilised for critically ill patients in intensive care units and for use during surgery [5][15].

Check throughout the report: Type in 3rd person and not colloquial/chatty. Now a description on non-invasive methods. The gold standard for BP measurement is the use of a cuffed sphygmomanometer. Cuff-based methods provide BP measurements without any major side effects as opposed to BP measured invasively [15]. However, patients will feel uncomfortable with long term monitoring due to the painful cuff inflation which interrupts the regular blood flow [8]. In addition, these methods can only measure BP intermittently with intervals between measurements greater than at least two minutes. These devices are too cumbersome to wear during measurements. Also, it has been found that over three in ten home BP monitoring cuffs have produced inaccurate results [16].

As a result, the existing invasive and non-invasive BP measurement techniques are not feasible for an implementation involving continuous ambulatory BP monitoring [15]. Hence, after having assessed the viability of all aforementioned methods, it is clear that it is difficult for

these methods to be integrated with wearable technologies, which continue to gain popularity in commercial sectors and clinical practice [1].

2.1.3 Ambulatory Blood Pressure (ABP)

ABP monitoring (ABPM) is when BP is measured as the patient moves around, and it allows patients to still live their normal daily lives [17]. It has been classed as the gold standard for detecting and diagnosing hypertension and also assessing BP values over a 24 hour period [4]. ABPM provides data on several important and unique parameters [4]. This data can explain how changes in your BP may correlate with your daily activities and sleep patterns [17]. Conventionally, ABP is monitored by using a cuff attached to a portable device which is worn on the patient's waist [4]. In the data provided for this project, the blood pressure signals have been recorded from ICU patients. As a result, these waveforms are not ABP waveforms but are instead Arterial Blood Pressure waveforms.

2.1.4 Electrocardiogram (ECG) signals

ECG signals provide an overview of the electrical impulses occurring in the heart [11]. Electrical changes in the heart conduct through the body and are received at skin level. The record of these electrical fluctuations during the cardiac cycle is called the Electrocardiogram (ECG) [18]. The signals are recorded by measuring the electric potential difference by placing electrodes across the heart of an individual [8] [11]. These electrodes are connected to the ECG machine with recordings from 12 different places on the body, which is known as the 12-lead ECG. The standard ECG leads are I, II, III, aVF, aVR, aVL, V1, V2, V3, V4, V5, V6. Leads I, II, III, aVR, aVR, aVL, aVF are classed as the limb leads and the others are precordial leads [8].

The QRS complex of an ECG signal is detailed in Figure 4. This complex is first created through the generation of the electrical impulses from the heart. These signals then move along the electrical highway and as a result cause the ventricles to contract and pump oxygenated blood into the arteries. Physically, this whole describes the QRS complex [18].

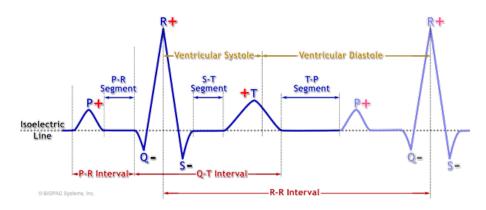


Figure 4: Structure of an ECG signal [19] [20]

2.1.5 Photoplethysmography (PPG) signals

Photoplethysmography (PPG) measures the blood volume changes per pulse. It is an optical and non-invasive technique that can determine a wide range of medical values, including an

estimate for BP [15]. Physically, the PPG signal is acquired by measuring the optical signal transmitted through or reflected from the subject's tissue [8]. The PPG sensor consists of two components. The first component is an Light Emitting Diode (LED) to light up the surface of the skin. The second component is a photodetector, which is utilised for measuring the changes in light absorption over a period of time [15] [18].

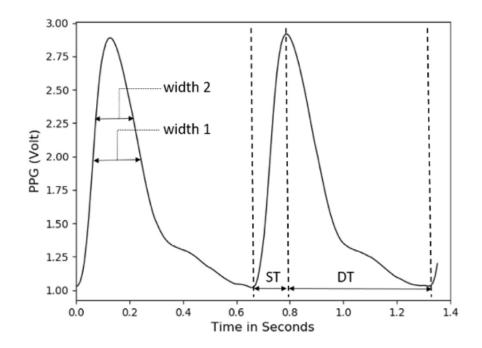


Figure 5: Structure of a PPG Signal [15]

In Figure 5, the four features are the Systolic upstroke Time (ST), Diastolic Time (DT), width at $\frac{1}{2}$ amplitude (width 1) and width at $\frac{2}{3}$ amplitude (width 2) [15]. PPG waveforms have a wide range of temporal features [15]. These features have been utilised in several experimentations, creating models to estimate blood pressure [12].

2.2 Cuff-less methods for deriving BP

Cuff-less methods have great potential in being used to estimate BP. This is because they provide continuous measurements, they cause minimal harm to the patients and they produce BP values over a long period of time [21]. There are three fundamental cuff-less methods which will now be discussed which can be used for deriving BP. These three methods rely on Pulse Transit Time (PTT), Pulse Arrival Time (PAT) and Pulse Wave Velocity (PWV) respectively [22]. These will each now be discussed in more detail.

2.2.1 Pulse Transit Time (PTT)

PTT is the time required for the arterial pressure wave to travel from the left ventricle to a distal arterial site. PTT holds an inverse relationship to blood pressure and as a result it is dependent on arterial compliance, arterial wall thickness, arterial radius, and blood density. PTT is conventionally found with the use of two PPG sensors [8] [9] [15], as indicated in Figure 6.

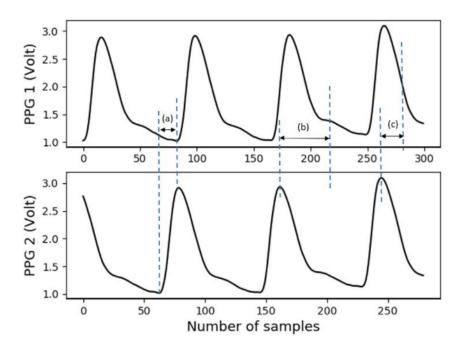


Figure 6: Pulse Transit Time (PTT) visualisation [15]

It is important to note for Figure 6 that the PTT can be measured at different points along the PPG waveforms. (a) represents a foot-to-foot time delay, (b) is a peak-to-dicrotic notch time delay and (c) is a peak to mid-point of the falling edge time delay [15]. As a proof of concept, increasing BP leads to an increase in the tension along the arterial wall tension, which therefore reduces the PTT. Hence, the opposite also applies [18].

2.2.2 Pulse Arrival Time (PAT)

The PAT is the difference in time between the R-peak of the ECG signal and the systolic peak of the PPG signal when measured during the same cardiac cycle, as indicated in Figure 7 [15] [6]. Physically, PAT is the interval in time between the activation of electrical impulses at the heart and the arrival of the pulse wave at a location on the body, such as the finger [23]. PAT is measured using two sensors, an ECG and a PPG sensor [15].

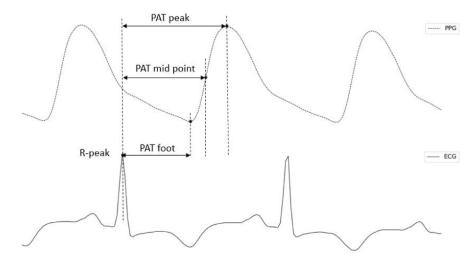


Figure 7: Pulse Arrival Time (PAT) visualisation [15]

The Pre-ejection Period (PEP) delay can also be briefly discussed. PEP is the time needed to convert the electrical signal into a mechanical pumping force and isovolumetric contraction to open the aortic valves,

$$PAT = PTT + PEP \tag{1}$$

2.2.3 Pulse Wave Velocity (PWV)

The PWV calculates the velocity of the pulse wave using two PPG sensors located on the same arterial branch at a known distance apart [12] [15]. The relation between PTT and PWV can be expressed as

$$PWV = \frac{d}{PTT} \tag{2}$$

where d is the arterial distance travelled by the pressure wave. PWV is related to the Young's modulus of the vessel wall by the Moens-Kortweg equation,

$$PWV = \sqrt{\frac{Eh}{\rho d}} \tag{3}$$

where

$$PWV = \text{Velocity of the pulse wave (m/s)}$$
 (4)

$$E = \text{Young's modulus of vessel wall (Pa)}$$
 (5)

$$h = \text{vessel thickness (m)}$$
 (6)

$$\rho = blood density (kg/m^3) \tag{7}$$

$$d = \text{arterial diameter (m)}$$
 (8)

The Young's modulus of the vessel wall is then related to the arterial pressure by the Bramwell-Hills equation,

$$E = E_0 e^{\lambda P} \tag{9}$$

where E_0 and λ depend on the thoracic and abdominal aortas and P is the vessel blood pressure (mmHg) [2] [8] [24].

By equating and solving Equations 2 and 3, the final equation for estimated blood pressure is expressed as,

$$P = \frac{1}{\lambda} \ln \left(2r \rho \frac{\Delta X^2}{E_0 h} \right) - \frac{2}{\lambda} \ln \left(PTT \right) \tag{10}$$

where

$$r = \frac{d}{2}$$
 = arterial radius (11)

$$\Delta X = \text{distance from heart to vessel}$$
 (12)

2.2.4 Limitations

The blood pressure (BP) can be derived through mathematical models as soon as estimates have been calculated for PTT, PAT and PWV. Although these models are common approaches for BP monitoring in an environment that is non-invasive and cuff-less, there are many challenges to these implementations. As a result, none of these techniques by themselves have been established as a reliable indicator for the estimation of BP.

Firstly, all three of the aforementioned methods require two separate measurements from two synchronised sensor devices. This can be a very inconvenient process for patients who are uncomfortable with this method [23].

In addition, there is a very likely possibility that these sensor devices will have different real-time sampling rates. Their operability depends on rather complicated arterial wave propagation models [15].

In order to be able to continuously measure BP, constant calibration of the methods is required. This is due to individual patients having different physiological parameters [23].

Finally, even with per-person calibration, these models can only provide BP estimation for a short period of time. As a result, this makes the models unreliable for the estimation of BP with every heartbeat [15].

To conclude this chapter, there is a lot of potential in the use of the three above parameters in the estimation of Ambulatory BP. However, there are still overarching limitations which currently hinder the progress of these parameters.

2.3 Neural Networks

You need an intro for neural networks. Again, you can't assume the reader knows what it is. Include a summary to explain what neural network is, before diving into particular types. Also summarise the type of networks you'll be discussing. You need to improve on organising and structuring here..

2.3.1 Artificial neural networks

Due to advancements in technology related to machine learning, there has been a lot of research into neural networks algorithms that can offer continuous BP measurements that are non-invasive and also cuff-less [12]. However, in this case BP estimation is motivated by how much data is available to the algorithm [15].

Artificial neural networks (ANNs) are a machine learning method that can be used to estimate blood pressure [12]. For this report, the aim is to experiment with Recurrent Neural Networks (TNNs). However, in order to properly understand RNNs, it is necessary to first introduce ANNs. ANNs are based on the neural networks found in the human body and aim to replicate their behaviour [24]. The structure of the ANN consists of multiple individual units called neurons, as shown in Figure 8.

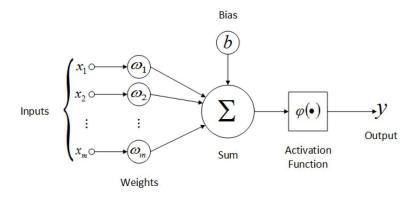


Figure 8: Structure of a neuron [25]

Each neuron has five critical components. These are inputs (\boldsymbol{x}) , weights $(\boldsymbol{\omega})$, transfer function (Σ) , activation function $(\phi(\cdot))$ and bias (b) [26]. In order to mathematically express the neuron unit, it is important to first understand its goal. The neuron applies a linear transformation to an m-dimensional input feature vector \mathbf{x} by applying a dot product with the weights $\boldsymbol{\omega}$ and adding a scalar bias b to this dot product. After this, a non-linear activation function $\phi(\cdot)$ is applied to the linear mapping. This enables the neuron to model non-linear relationships. This is expressed mathematically as follows,

$$y = \phi(\sum_{i} \omega_{i} x_{i} + b) = \phi(\boldsymbol{\omega}^{T} \mathbf{x} + b)$$
(13)

where $\omega \in \mathbb{R}^m$ and y, b are scalars. When each of these neurons are connected together with several other neurons across several layers, this forms a neural network, as shown in Figure 9. Each of the neurons in Figure 8 are represented by a grey unit in Figure 9. By having a network of multiple neurons, it is possible to model more complex relationships than just a

single neuron, provided that the activation functions $\phi(\cdot)$ are not linear for all neurons (since the combination of linear operations results in a linear operation). In addition, the output of the neural network can have as many units as needed depending on the task at hand (e.g. 5 neurons are needed for classification problems with 5 classes using one-hot encoding).

The network of a single fully-connected layer is mathematically expressed using Equation 14,

$$\mathbf{y} = \phi(\mathbf{\Omega}\mathbf{x} + \mathbf{b}) \tag{14}$$

where $\Omega \in \mathbb{R}^{N \times M}$ is the weights matrix, $\mathbf{b} \in \mathbb{R}^N$ is the bias vector, $\mathbf{y} \in \mathbb{R}^N$ is the output vector and $\phi(\cdot)$ performs element-wise non-linear transformations.

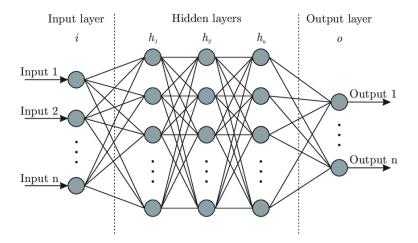


Figure 9: Structure of a multi-layer ANN

2.3.2 Recurrent Neural Networks (RNNs)

Traditional neural networks have found success in many fields, However it has been demonstrated that they cannot capture temporal dependencies in the data, making it unsuitable for signal processing applications. A Recurrent Neural Network (RNN) is a specific type of architecture that is widely used to deal with time-varying data [27]. RNNs contain additional memory states that retain and process information from previous time steps.

RNNs are called recurrent since they apply the same operation to each of the input sequences, with the output of an individual element being dependent on the previous one. Theoretically, RNNs establish a connection between the actual input and all the previous ones [27]. Although this is assumed, in the practice, RNNs have proven to only remember a limited number of inputs. In other words, RNNs have a memory that allows them to remember previous elements and use their information to deal with the current input [26]. Figure 10 shows the simplest version of an RNN, which can be easily derived from a simple feedforward architecture by adding a single loop:

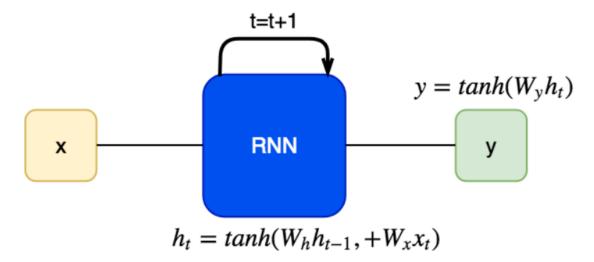


Figure 10: Simplification of a RNN

During training, the hidden state h is iteratively updated based on the input value x and the learned weights W_h and W_x . The final output y is estimated from the current state h_t and the matrix W_y . Although RNN can assure short-term dependencies within the network, simple RNNs become unable to learn to connect information as the gap between past and present information grows [28]. To overcome this limitation, in practical applications LSTM unit is adopted, that is a special RNNs architecture composed of multiple interacting layers.

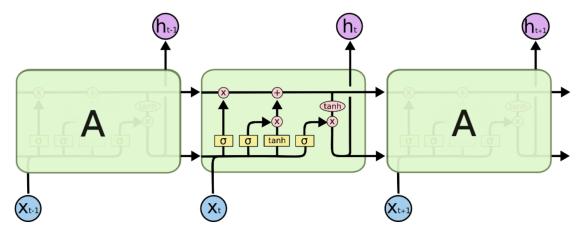


Figure 11: LSTM network

2.3.3 Activation Functions

Activation functions transform the output of a neural network unit element-wise, allowing it to model non-linear functions. For this project, the estimation of BP is treated as a regression problem. Hence,

2.3.4 Loss Functions

Loss functions are objective functions that the neural network aims to minimize when being trained. For this project, the loss function of concern is the Mean Squared Error (MSE) loss

function, which is defined in Equation 15.

$$l_{\text{MSE}}(\mathbf{y}, \hat{\mathbf{y}}) = \frac{1}{N} \sum_{i=1}^{N} (\mathbf{y}_i - \hat{\mathbf{y}}_i)^2$$
(15)

where N is the number of training examples, \mathbf{y}_i is the target output vector, and $\mathbf{y}_{\hat{y}_i}$ is the predicted output vector. The MSE is effective for ensuring that the trained model has no outlier predictions with huge errors, since the MSE puts larger weight on these errors due to the squaring operation of the function.

2.3.5 Neural Network Training

In the same manner for any other machine learning algorithm, neural networks aim to learn an underlying pattern present in the data by minimizing an error measure defined by a loss function given some sample data (a process known as training). Formally, this problem is done by finding the optimal weights of the model per layer as defined in Equation 16, where $\hat{\mathbf{y}}$ is the estimate of the true label \mathbf{y} and l is the loss function.

$$\Omega_{opt} = \arg\min_{\mathbf{\Omega}} \{ \mathbb{E}(l(\mathbf{y}, \hat{\mathbf{y}})) \}$$
(16)

At the end of this chapter, I advise you to summarise what the reader should take from this chapter. What is the takeaway message and how are you moving forward from this knowledge. Here you restate your aim and objective and how you are going to move forward. (The summary should flow into the next chapter)

2.4 Overview on Cuff-Less BP device options

Wearable technology provides an opportunity for real-time monitoring of human vital signs, thus enabling the possibility for preventive, timely notification and real-time diagnosis. Unlike commonly-used BP sensors, which demand a specific measurement procedure, modern wearable bio-sensors monitor vital signals online and all day long, presenting no additional burden other than wearing the device. A wide range of wearable devices achieve very positive results, even those wearables which are cheaper in price [3]. Some of the systems developed for the purpose of non-invasive BP monitoring will be discussed in more detail now. In addition, they will be critically analysed against two other viable alternatives.

The motivation behind this project is to replace the current cuff-based BP devices. Cuff-based devices often require the supervision of an expert to work correctly and do not provide continuous measurements for BP. In addition they can cause irritation and inconvenience for patients due to cuff inflation and deflation. As a result, current clinical cuff-based BP devices are not suitable for providing continuous BP monitoring which could play a significant role in the early detection of diseases which affect the heart [15].

2.4.1 Smart Watches

Smart watches are becoming an increasingly popular form of wearable technology [7]. Most of the existing smart watch produces currently measure BP through Pulse Transit Time (PTT) from a pulse wave measurement at the wrist. The Heartisans BP smartwatch uses ECG and PPG signals to measure PTT and estimate BP. The watch requires a motionless 20 second scan with the device held at heart level for measurements and provides systolic and diastolic BP readings. Calibration with a validated cuff-type BP device is required prior to standalone use. Despite its availability on the market, the Heartisans Watch has not undergone a formal validation study [7].

Another wearable method is through arterial tonometry. The BPro device, developed by the London company HealthSTATS Technologies, is one such device.

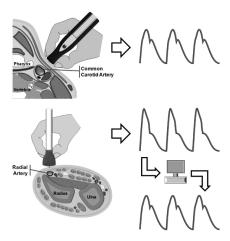


Figure 12: Arterial Tonometry

Figure 12 shows two different methods for using arterial tonometry to measure blood pressure. The first image shows that the recording is taking at the carotid artery level in order to estimate the central blood pressure waveform. In the second image, the recording is taken at the radial artery level to estimate the arterial pulse wave. The central waveform is then reconstructed from this pulse wave using software.

The BPro allows for continuous 24 hour waveform analysis of BP. Calibration to the brachial artery BP is required using a validated upper arm cuff device prior to use. The BPro has been validated in a non-ambulatory study following the Association for the Advancement of Medical Instrumentation (AAMI) standards and the European Society of Hypertension (ESH) protocol.

2.4.2 Smartphone Applications

Smartphones offer a great potential to expand the continuous recording of BP if their sensors can be correctly utilised [7]. A formal validation study on one particular iOS application, the Instant Blood Pressure app, revealed poor accuracy of BP measurements. Mean absolute differences of 12.4 mmHg for systolic BP and 10.1 mmHg for diastolic BP were found between the Instant Blood Pressure application and a reference device. This resulted in approximately four out of five hypertensive individuals being falsely classified as normotensive [7]. The My BP Lab application measures BP through measurements of Pulse Transit Time (PTT). However, there is currently a lack of experimental data to justify it's dominance in the market.

2.4.3 Medical Tricorders

A medical tricorder is a handheld portable device used by consumers to self-diagnose medical conditions and take basic vital signs measurements. The BodiMetrics Performance Monitor uses ECG and PPG signal data to estimate BP through PTT. The Bodimetrics tricorder obtains measurements with a 20 second scan of the user's fingertip at heart level but only provides systolic BP data. This tricorder has a large spread in absolute bias against an automated sphygmomanometer, hence it is unlikely to meet formal accuracy and precision standards [7].

The FreeScan Personal Cardiovascular Monitor, which is developed by the Taiwanese company Maisense Inc., also estimates BP from PTT. However the device uses a force sensor to capture the systolic arterial waveform rather than PPG. This requires the user to physically apply the force sensor directly to the radial artery for around 10 seconds for measurements. The Freescan device has been verified according to the AAMI protocol [7].

The SOMNOtouch NIBP is a non-traditional medical tricorder that utilizes PTT data collected in a similar fashion to the Bodimetrics Performance Monitor. The device has met the ESH standards, however it begins to lose accuracy for higher SBP and DBP values [7].

2.4.4 Conclusion

After having discussed three viable devices for cuff-less measurements of BP, it has been finalised that smart watches are the most viable option. Whilst existing smart watch devices do suffer from inaccuracies in BP estimation due to motion, it is clear that they produce the most acceptable results in line with the AAMI and ESH standards. This chapter can be seen as a forward looking overview of how the estimation methods discussed in this report can be used to benefit future products. With regards to this project, this chapter can be treated as a supplementary overview.

2.5 SECTION CLARIFYING WHAT COMPLEXITY MEANS FOR THIS PROJECT?

2.6 Literature Review

Initial remarks: Where is the use of the PRISMA criteria? Where is the inclusion, exclusion diagram? After reading remarks: This is not a literature review. By only placing papers into a table as a summary is not sufficient. - what comments do you have on the results you've tabulated? - You mention factors were considered.. why? and when you considered them, what about them? Why is it important? - What should the reader be left with?

What are the advantages and disadvantages?

Currently, the literature review is below average. You need to work on this.

This chapter provides a detailed account of the literature review conducted for this FYP. The literature search equation used will first be discussed, followed by an explanation of the PRISMA flow diagram and how it was used to benefit this literature review. To help the reader, a table of the scientific papers used in this project is provided. Finally, a critical analysis will be given on the literature review and what can be concluded as a result. WHAT DO YOU AIM TO GET OUT OF THIS?

2.6.1 Survey Equation

At the beginning of the FYP, the only information provided was the FYP mission statement (see Appendix Item 10.1) and two published papers, A review of machine learning techniques in photoplethysmography for the non-invasive cuff-less measurement of blood pressure [15] and Continuous Blood Pressure Estimation From Electrocardiogram and Photoplethysmogram During Arrhythmias [21]. These resources pserved as an introduction to both the medical background and machine learning knowledge for myself. After reviewing this information, the next step was to perform an informal search of literature databases using keywords extracted from the FYP brief. These keywords can be divided into two fields:

Background knowledge

- "Ambulatory"
- "Blood Pressure"
- "Electrocardiogram"
- "Photoplethysmography"
- "Wearable technology"

Implementation strategy

- "Accuracy"
- "Algorithm"
- "Computational complexity"

These keywords were entered into 3 official literature databases, as shown in Table 2.

Table 2: Official online databases used to conduct the literature review [29]

| Literature database | Description |
|---------------------------------------|--|
| ACM Digital Library | The digital library of the Association for Computing Machinery |
| Engineering Village | Database platform for Physics, Electrical Engineering, Electronics and Computing |
| IEEE Xplore | Digital library containing full text of IEEE journals, conference/meeting papers and standards |
| National Library of Medicine (Pubmed) | Biomedical and life sciences literature |

This informal search enabled a clearer understanding of how the relevant published literature phrased their titles. Based on the findings of the informal search, the following literature survey search equation was used to identify the literature that best fits the needs of the FYP requirements. The equation chosen was:

• (Extraction OR Estimation OR Review) AND (Blood OR Arterial OR Ambulatory OR Cuffless) AND (Pressure) AND (ECG OR PPG) AND (Machine Learning OR Signal Processing).

Hence, this equation was entered into the four databases displayed in Table 2.

2.6.2 PRISMA checklist

After applying the chosen equation to the four databases in Table 2, the following PRISMA checklist was created:

The papers returned were then filtered to only those published between 2012 and 2022, i.e. the last 10 years. As a result the following tables summarise the most relevant papers for this project.

2.6.3 Literature survey table

As a reference, the original literature survey matrix can be found on the Github repository [30]. Firstly, in Table 3, a simplified literature survey has been detailed out for the best performing methods which do not employ machine learning methods.

Table 3: Overview of performance of the best non-invasive non-ML cuff-less methods for measuring BP

| Study | Source | No. Subjects | Age | Implementation | MAE SBP |
|-------|----------------------|--------------|-------|---------------------|------------------|
| [31] | ECG, PTT-CP | 10 | 24-63 | Numerical solution | ±5.93 |
| [32] | ECG | 5 | N/A | Analytical solution | 9 ± 5.6 |
| [33] | PPG | 16 | 18-48 | Frequency analysis | 0.8 ± 7 |
| [34] | ECG, PPG, PTT | N/A | N/A | Analytical solution | 7.49 ± 8.8 |
| [35] | PTT | 127 | N/A | Wavelet transforms | ± 7.63 |
| [36] | PTT, PPG | 27 | 21-29 | Analytical solution | -0.37 ± 5.21 |

Table doesn't fit on the page. Consider either turning it landscape or redesigning the table to fit.

Redesigning could be making a key for different sources and/or the method. E.g ECG = filled square, PPG = filled circle.

If you are abbreviating or using symbols, state clearly in your table captions.

Table 4: Overview of performance of the best non-invasive ML cuff-less methods for measuring BP

| \mathbf{Study} | Source | No. Subjects | Age | Method | MAE SBP |
|------------------|---------------|--------------|------------------|--------------------------|-------------------|
| [24] | ECG, PPG | 14 males | 17-43 | ANN | 7.99 ± 10.34 |
| [37] | PPG | 65 | 22-65 | Wavelet, SVM | 5.1 ± 4.34 |
| [38] | PPG | MIMIC II | Adults | Linear Reg., ANN, SVM | 13.84 ± 17.56 |
| [3] | ECG | 51 | 16-83 | Complexity analysis + ML | 7.72 ± 10.22 |
| [6] | PPG | 72 | N/A | ANN (MLP) | 4.02 ± 2.79 |
| [12] | ECG, PPG | MIMIC II | Adults, neonatal | ANN (150 neurons) | 5.76 ± 6.39 |
| ∞ | ECG, PPG | 39 | 20-100 | ANN-LSTM | 1.10 |
| [39] | PTT, ECG, PPG | MIMIC I | N/A | SVM, Lin Reg. | 3.27 ± 5.52 |
| [40] | PTT | 250 | MIMIC I | ANN-RBM | 3.70 |

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2.6.4 Critical analysis of literature survey table

2.6.5 Conclusions of literature survey

The Mean Absolute Error (MAE) of the Systolic BP (SBP) was used as the uniting accuracy measure in this paper, as it was the most readily available parameter in all of the aforementioned papers.

It is important to note that other factors were considered in this literature survey. These factors were,

- Range of SBP and DBP values
- Sampling frequency
- Denoising and detection techniques used
- Computational complexity
- Feasibility in a wearable context

However, due to a lack of regular occurrences of these factors over all the papers, they were not included in the above two tables.

The results show that the established non-ML methods do produce MAE values noticeably lower than the majority of the ML methods. However, a main factor to consider about these results is that machine learning based methods and neural networks are data driven. As shown in Table 4, there is a very limited number of subjects available for each study. If these studies had been extended to include more test patients, it is possible that these MAE SBP values were lower. An additional point is that another study [41] did a study into using a 4-layer LSTM architecture to estimate BP from ECG and PPG signals with 84 patients. This study resulted in an RMSE SBP of 3.9 mmHg but no available MAE. Hence there is a lot of potential in ML methods when there is sufficient data available.

3 Analysis and Design

3.1 Choice of programming language

Python is used as the sole programming language for this project. Python has a wide variety of easy to use and powerful libraries [42]. The scientific libraries from Python that are used for this project are numpy [43] and pandas [44]. In addition, the machine learning libraries used are tensorflow [45], keras [46] and scikit-learn [47]. In addition the heartpy [48] and wfdb-python [49] packages were installed, which are libraries of tools for reading, writing, and processing Waveform-Database (WFDB) signals and annotations.

MATLAB was also considered as a potential programming language to use, due to it having a wide range of signal processing and machine learning add-on toolboxes. However Python has been shown to offer a wider set of choices in graphics packages and toolsets, such as through matplotlib [50], and it also produces more compact and readable code.

3.2 Choice of dataset

Give a better explanation of the dataset you're using. Age range, gender split, any medical conditions.. This can all be found in the dataset website or the relevant papers linked to the dataset. As previously discussed in the literature review in Chapter 2, the chosen dataset is the Medical Information Mart for Intensive Care (MIMIC) dataset. Did you? I must have overlooked it. The MIMIC Database includes data recorded from over 90 ICU patients. The data in each case include signals and periodic measurements obtained from a bedside monitor as well as clinical data obtained from the patient's medical record. The recordings vary in length; almost all of them are at least 20 hours, and many are 40 hours or more If you know how many, state it. Don't be chatty.. In all, the database contains nearly 200 patient-days of real-time signals and accompanying data [51]. I don't understand.. You say there are over 90 ICU patients (again state the exact number because it is known). So where do the 200 come from? (again can't be nearly 200.. Be exact.) - Did you use all 90 or 200 patients data. Everything you write in your methods needs to be clear for someone to repeat exactly what you have done. This will also help us review your code and run it to produce the exact same results you will be showing in the report.

- Choice of signal channel 3.3
- Feature extraction 3.4
- Decision between signal-processing based or ML based method 3.5
- 3.6 Choice of ML model to use

3.7 Performance metrics

The two considered error calculations used in this experimentation are the Mean Absolute Error (MAE) and Root Mean Square Error (RMSE). They are defined by the following equations,

$$MAE = \frac{1}{N} \sum_{i=1}^{N} |a_{i_M} - b_{i_M}| \tag{17}$$

$$MAE = \frac{1}{N} \sum_{i=1}^{N} |a_{i_M} - b_{i_M}|$$

$$RMSE = \frac{1}{N} \sqrt{\sum_{i=1}^{N} |a_{i_M} - b_{i_M}|^2}$$
(18)

In the context of BP estimation, b_{i_M} and a_{i_M} represent the true value and BP estimate respectively for the Mth element of the time sequence.

4 Implementation

4.1 Description of dataset

As discussed in Section 3, the MIMIC Database includes data recorded from over 90 ICU patients. For the purposes of this project, a subset of this data is used for experimentation. Further details of the data used are shown in Table 6.

Why a subset? Why not all? If you're doing a split, explain this. Are you only using 8 patients out of the 90?

| Patient Number | Gender | Age | Health issue faced by the patient |
|----------------|--------|-----|---|
| 224 | Male | 21 | Sepsis |
| 225 | Male | 73 | Pulmonary edema |
| 230 | Female | 75 | Cardiac Heart Failure/Pulmonary edema |
| 232 | Male | 68 | Myocardial infarction/Cardiogenic shock |
| 235 | Female | 67 | Myocardial infarction/Cardiogenic shock |
| 240 | Male | 68 | Angina |
| 252 | Male | 52 | Respiratory Failure |
| 255 | Male | 67 | Cardiac Heart Failure/Pulmonary edema |

Table 6: Characteristics of the patients from the MIMIC-I database

4.2 Extracting the ground truth blood pressure values

The ground truth Systolic and Diastolic blood pressure values are calculated by taking the respective maxima and minima of the arterial blood pressure signal within each window of the signal.

4.3 Signal preprocessing steps

4.4 Windowing of PPG and ABP data

4.5 Convolutional Neural Network (CNN) model

What type of information should the reader gather from this figure? It needs explaining and annotations/labelling.

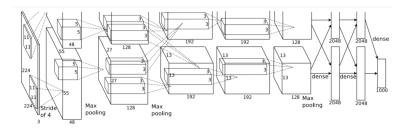


Figure 13: Overview of the AlexNet architecture

4.6 ResNet model

5 Overview of Results

For all results:

- 1. Ensure all results have units (if it doesn't have units, it's A.U for arbitrary units)
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5.1 Overview of testing parameters

5.2 Performance of AlexNet architecture

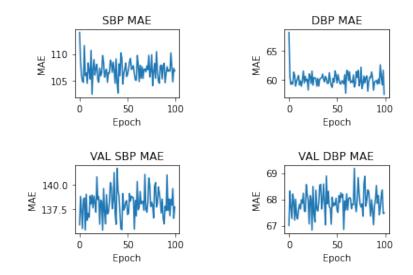


Figure 14: MAE of SBP and DBP for AlexNet architecture

5.3 Performance of ResNet architecture

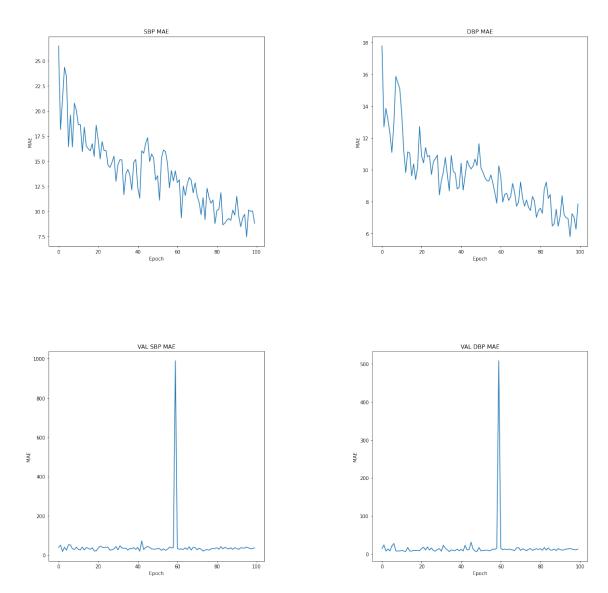


Figure 15: MAE of SBP and DBP for ResNet architecture

6 Evaluation of results

This Chapter (or possibly section of the conclusions) is distinct from your results. It must contain your critical evaluation of your work as compared to previous analysis, algorithms, products, and when related to your original objectives. To what extent have your original objectives been fulfilled? If they have changed, what is your rationale for this? What are the advantages, disadvantages of your approach compared with related work? How does the scope of your work differ from related work? Examiners expect your project report to show evidence of your ability to think as an engineer, and that includes the ability to critically reflect on your own work and evaluate its significance.

Material here will compare project outcomes with initial objectives and requirements captured. Usually your Interim Report will contain these. Where these have changed significantly over the course of the project this should be explained and reasons given. This section should not require examiners to read your Interim Report, and will not reference it. Changes between final and initial objectives should be explained in a self-contained manner.

Note that here you will reference and summarise, rather than repeat, your description of Requirements Capture earlier in the Final Report.

7 Conclusions and Further Work

7.1 Summary of project achievements

- Design choices: Aim was to minimize complexity of the neural network model in order to effectively monitor BP
- What was most difficult: 2 parts to this: Firstly the choice of how to preprocess the signal for effective analysis. This was solved by surveying the relevant literature and assessing the most feasible set of steps. Secondly, designing the ideal neural network architectures
- What I learned: Despite what was expected from the theory, the CNN Resnet performs better on the PPG signal data than the LSTM networks

•

7.2 Future work

• Discuss the transformer papers and the attention mechanism. The main reason the transformer model was not explored for this paper is due to its high model complexity. However, this could be tested on in the future and see if the results are any/much better than the Resnet/LSTM models

•

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8 Appendix

8.1 FYP Mission Statement (correct to June 2022)

'Ambulatory blood pressure monitoring has become increasingly relevant due to the advancement of wearable technology. Modalities such as Electrocardiogram (ECG) and Photoplethysmography (PPG) have provided an indirect method of blood pressure estimations compared to a traditional blood-pressure monitor. Although algorithms exist for calculating blood pressure with ECG and PPG, it is vital that their computational complexity is minimal, whilst maintaining accuracy, due to the power limitations of wearable technology. The goal of this project is to establish the tradeoffs between using PPG and ECG to quantify blood pressure, in the context of wearable technology, where power must be kept to a minimum. This project is ideal for students interested in signal processing, who have excellent programming skills in Matlab.'

8.2 Health standards requirements for blood pressure estimation

- The Advancement of Medical Instrumentation (AAMI) standard requires a mean BP difference of ≤ 5 mmHg with a standard deviation of ≤ 8 mmHg against auscultatory reference measurement
- Significant variation in BP measurements(> 12 mmHg systolic or > 8 mmHg diastolic) from the validated reference device is an exclusion criterion in the AAMI protocol [7]
- The European Society of Hypertension (ESH) protocol requires that the majority of subjects have investigational BP readings within ≤ 5 mmHg of the reference measurement.

8.3 Complete FYP Gantt chart