

Introduction

With the rapid growth of computational power and complex algorithms, we propose a novel approach to detect arrhythmias in Phonocardiograms (PCGs). Typically, Electrocardiograms are used to diagnose arrhythmias, requiring medical-grade equipment to recognize cardiac illnesses accurately. However, PCGs provide ease of access to everyone who has a device capable of recording audio, allowing medical professionals to treat arrhythmias in the developmental stages. The new design comprises two subsystems: one is based on the relationship between Electrocardiograms (ECGs) and PCGs, and the other between PCGs and arrhythmias. The association between ECGs and PCGs is amended to translate from one space to another, where ECGs become dimensionally reduced, then reconstructed into a PCG signal. The second subsystem uses a Generative Adversarial Networks (GAN), in which both arbitrary PCG signals and generated signals are fed into a discriminator that detects if an arrhythmia is present or if the signal is false.

Background

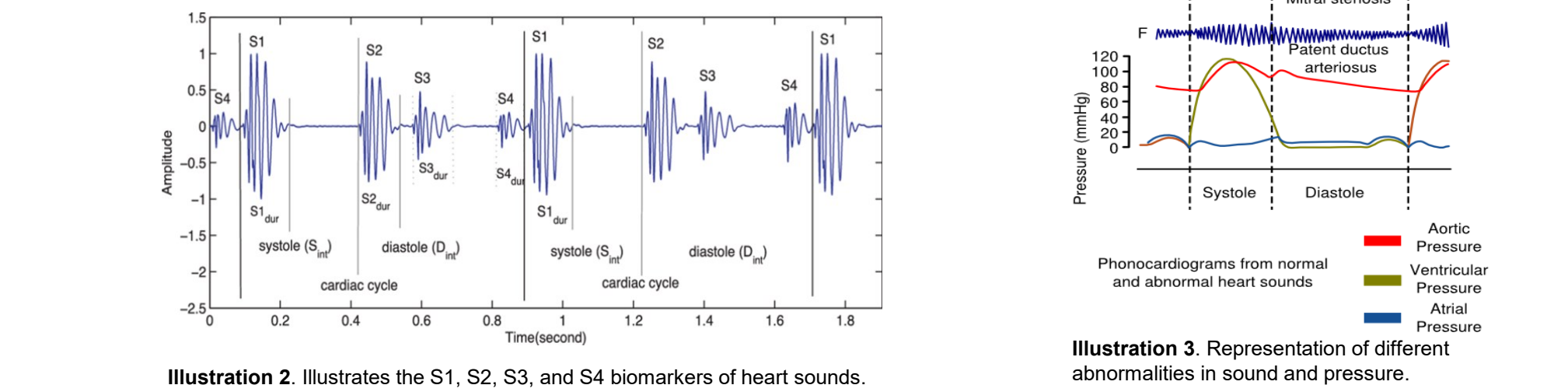
An estimated three million cases of arrhythmia occur in the United States yearly (Mayo Clinic), with 300,000 sudden deaths per year – an incidence rather high than stroke, lung cancer, or breast cancer (American Heart Association). Traditionally, non-invasive arrhythmia analysis is based on multiple electrodes that reflect the electrical activity on ECGs. This method, despite being accurate, limits the use case to hospitals and clinics with specialized equipment; thus, limiting the portability of diagnosing, let alone classification of the type of pathology.

Phonocardiograms (PCGs) are sounds that are created by the mechanical movement of the heart. This physical movement produces four distinct sounds: S1, S2, S3, S4, and murmurs. S1 and S2 are sounds created by a healthy heart; whereas S3, S4, and murmurs refer to diseases or anomalies. The first heart sound, S1, marks the start of Systole. Systole occurs when the heart muscle contracts and pumps blood from the chambers into the arteries. The second heart sound, S2, marks the end of Systole and the start of Diastole. Diastole is a phase of the heartbeat when the heart muscle relaxes and allows the chambers to fill with blood.

Related Works

Although heart sound databases do exist, these datasets are limited by the number of pathologies that are collected, often having to divide the dataset into two categories: normal and abnormal. Currently, only three major supervised PCG datasets exist: PhysioNet Classification of Heart Sound Challenge dataset, PASCAL Heart Sound Challenge dataset, and the Littman Heart Sound and Murmur Library. Additionally, little effort has been done to increase the development and labeling of comprehensive datasets of PCG signals that cover the complete range of pathologies.

In diagnosing heart sounds, two major challenges arise: localization and classification. Localization aims to find the position of biomarkers in heart sounds. By doing this, heart sounds can be segmented into signals containing a single heart sound. Furthermore, classification attempts to categorize heart sounds into normal and abnormal groups by exploiting the information extracted from localization. Conventional heart sound localization and classification methods involve time and/or frequency and are typically dependent on machine learning algorithms to enhance the results. These algorithms typically include artificial neural networks (ANNs), support vector machines (SVMs), self-organizing maps (SOMs), and are limited to the number of samples and pathologies covered in a given dataset. This leads to a surface-level analysis of the heart sounds.



Objectives

Problem: Current detection methods have limited performance in pathologies and lack real-time classification capabilities.

Question: Is it possible to use Generative Adversarial Networks (GANs) to accurately detect arrhythmias in PCGs to decrease the undiagnosed rate of arrhythmias?

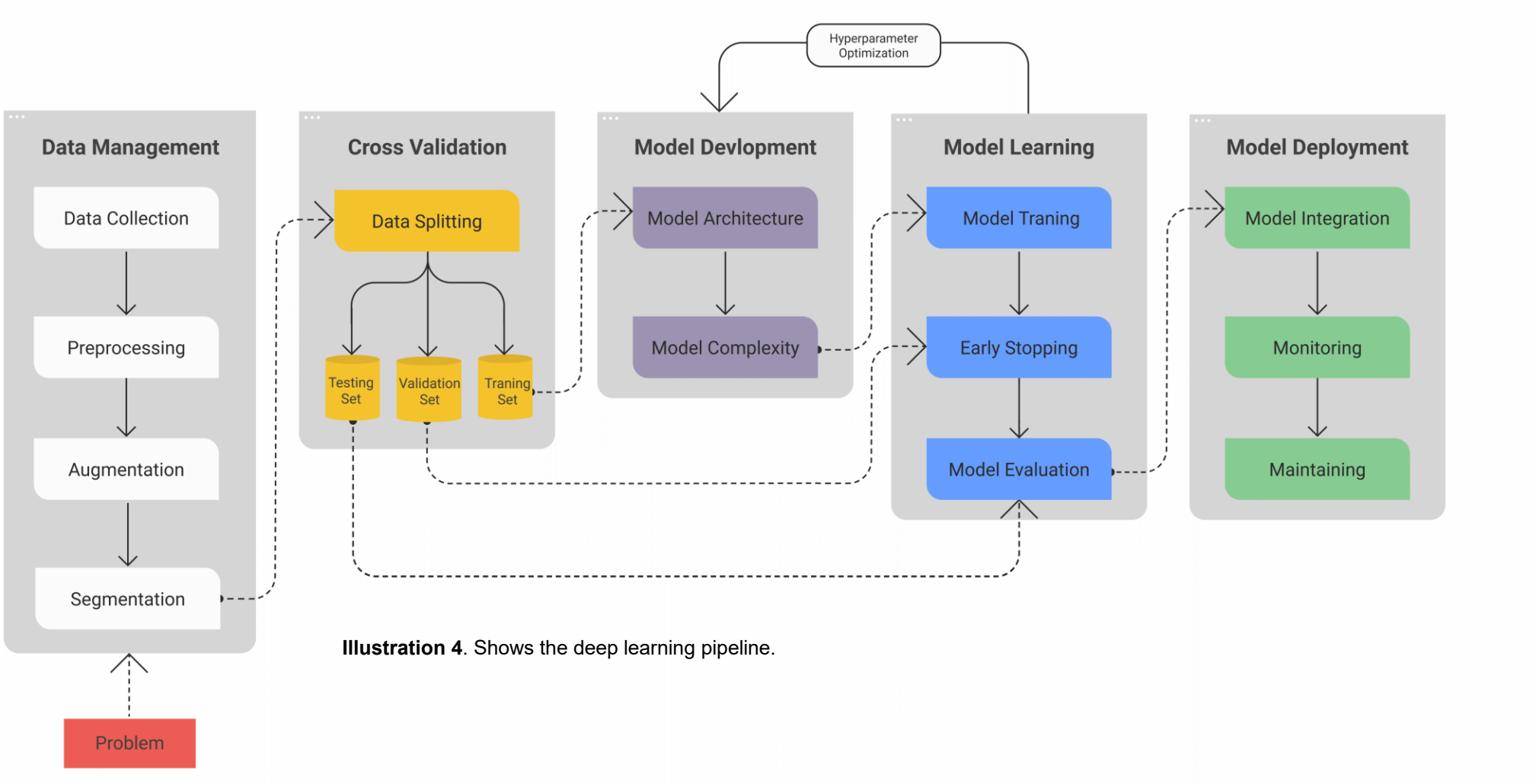
Hypothesis: If a novel heart sound analysis system is developed to detect a variety of arrhythmias, then arrhythmias will have a decreased undiagnosed rate.

Solution: The new semi-supervised approach is composed of two subsystems; the first subsystem uses a discriminator, in which PCG signals are classified into categories based on arrhythmias in the signal. The second subsystem – a generator, aims to generate data such that, when fed into the discriminator, the discriminator will classify the generated data as abnormal or normal. While the discriminator aims to adversarially classify the generated data as fake and classify the real PCG signals to their respective categories. This adversarial approach allows the discriminator to not only extract PCG signal specific features, but also introduces the discriminator potential noisy signals that should not be classified.

Engineering Goals

- 1. Increase the Number of Cardiovascular Pathologies analyzed in heart sounds-** Develop a model to construct heart sounds from pre-existing datasets that cover the complete range of pathologies that are likely to be encountered in clinical settings.
- 2. Develop an End-to-End System** - Create a system that is able to record, analyze, and predict (end-to-end system) heart sounds for Cardiovascular modalities without specialized equipment.
- 3. Real-World Testing** - Test the end-to-end system in a real-world environment to ensure the practicality and generality of the system.

Methods

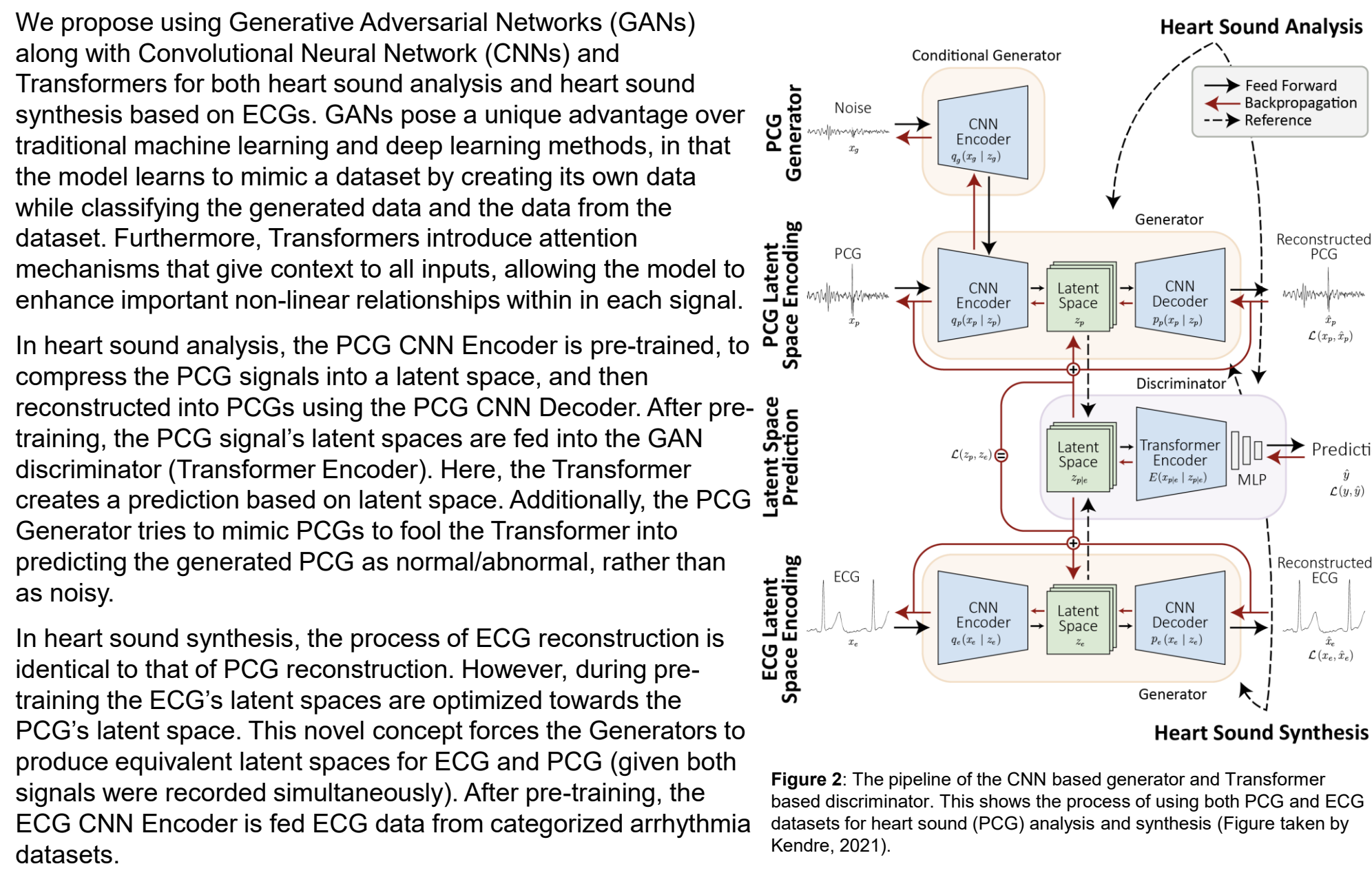


Employing Adversarial Machine Learning and Computer Audition for Smartphone-Based Real-Time Arrhythmia Classification in Heart Sounds

Data Management

Dataset	Dataset Type	Lengths	Environment & Recording Quality	Pathologies Ratios
Classification of Heart Sound Recordings - PhysioNet 2016	PCG & ECG	5-120 seconds	Extremely noisy and low signal quality	Normal: 3541 (77.1%) Abnormal: 551 (12.0%) Noisy: 501 (10.9%)
PASCAL 2011	PCG	1-30 seconds	Noisy and taken from digital stethoscope	Normal: 351 (40.0%) Murmur: 29 (14.7%) Extra: 86 (9.8%) Unlabeled: 247 (28.1%) Stenosis: 5 (1.3%) Septal: 1 (0.3%) Ejection: 5 (18.8%) Coronary: 1 (6.3%) Prosthetic: 1 (6.3%) Regurgitation: 2 (12.5%) Pericarditis: 1 (6.3%) Gallop: 2 (12.5%)
Littman Heart Sound & Murmur Library	PCG	2 seconds	Clean and taken from digital stethoscope	Normal: 9028 (24.2%) CD: 5486 (19.7%) MI: 5250 (18.9%) HTF: 4007 (17.8%) STTC: 2055 (9.5%)
PTB-XL	ECG	10 seconds	n/a	

Model Development



Distribution Table

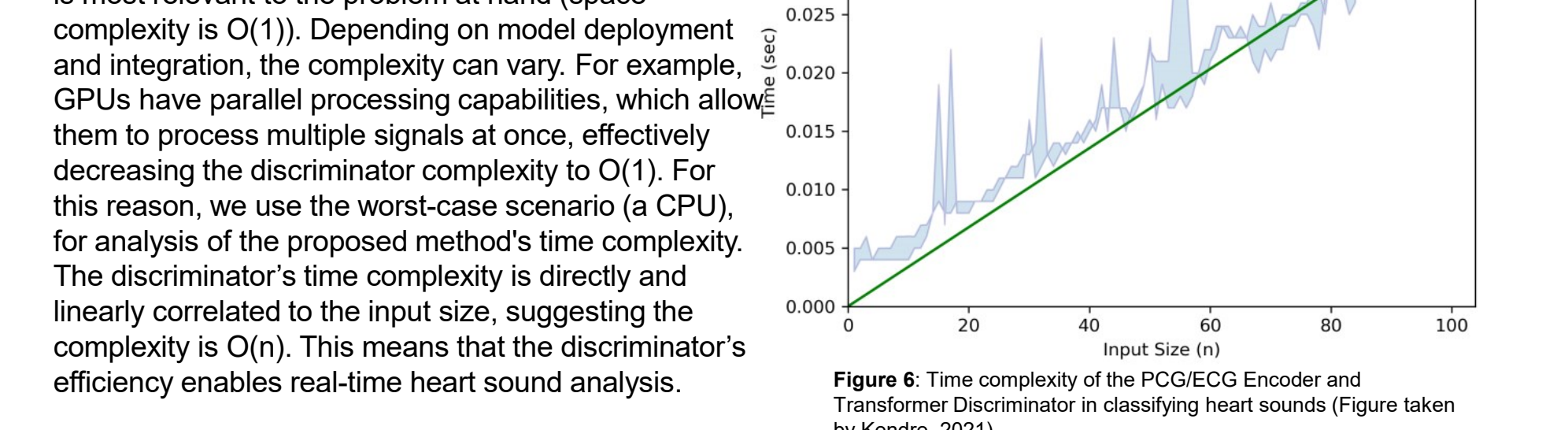
Dataset	Accuracy	F1 Score	Sensitivity	Specificity	PPV	NPV
PASCAL	73.7±0.02	75.0±0.04	75.0±0.05	72.2±0.04	75.0±0.02	72.2±0.04
PhysioNet 2016	89.5±0.03	64.4±0.05	48.0±0.06	85.7±0.02	96.5±0.04	85.7±0.03
Synthesized PTB-XL (AF)	83.0±0.03	94.3±0.03	99.5±0.03	90.3±0.03	99.3±0.03	91.8±0.03
Combined	95.0±0.03	94.6±0.04	94.3±0.04	99.5±0.03	90.3±0.03	99.3±0.02

Confusion Matrix

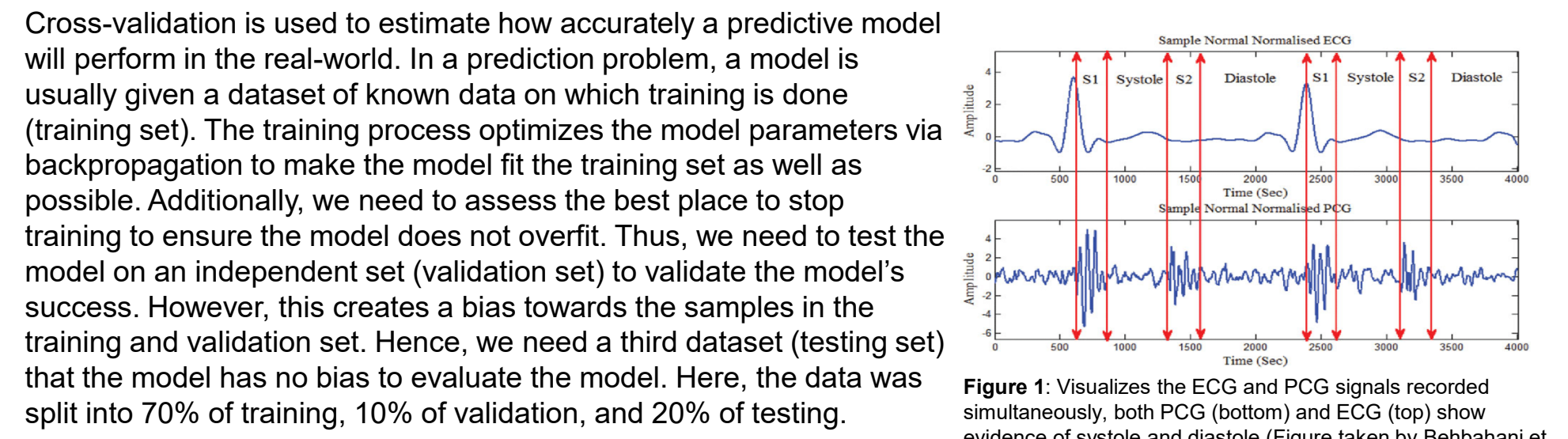
True label \ Predicted label	Normal	Murmur	Gallop	AF
Normal	0.959	0.0	0.041	0.0
Murmur	0.029	0.964	0.007	0.0
Gallop	0.094	0.062	0.944	0.0
AF	0.0	0.0	0.05	0.0
Noisy	0.0	0.046	0.0	0.954

Figure 4: Matrix of accuracy between categories (pathologies) classified by the model (predicted label) and the true categories (true label) (Figure taken by Kendre, 2021).

Time Complexity



Cross-Validation



Model Learning

Mean Squared Error:
 $MSE(x, y) = \frac{1}{n} \sum_{i=1}^n (x_i - y_i)^2$

Cross Entropy Loss:
 $CE(y, \hat{y}) = -\log(\frac{\exp(\hat{y}_i)}{\sum_{j=1}^K \exp(\hat{y}_j)}) = -\log(\frac{\exp(\hat{y}_i)}{\sum_{j=1}^K \exp(\hat{y}_j)})$

Accuracy:
 $ACC = \frac{TP + TN}{TP + TN + FP + FN}$

Sensitivity:
 $TPR = \frac{TP}{P} = \frac{TP}{TP + FN} = 1 - FNR$

Specificity:
 $TNR = \frac{TN}{N} = \frac{TN}{TN + FP} = 1 - FPR$

Positive Predictive Value:
 $PPV = \frac{TP}{TP + FP} = 1 - FDR$

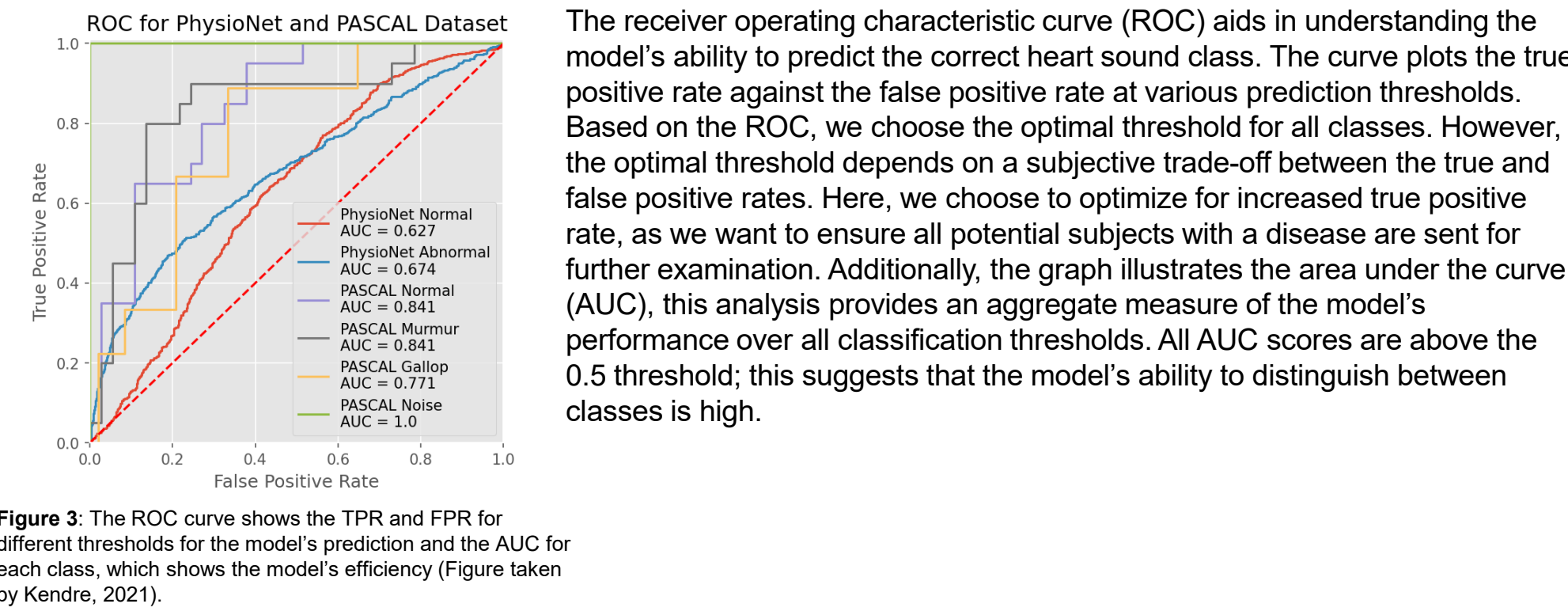
Negative Predictive Value:
 $NPV = \frac{TN}{TN + FN} = 1 - FOR$

F1 Score:
 $F_1 = 2 \times \frac{PPV \times TPR}{PPV + TPR} = \frac{2TP}{2TP + FP + FN}$

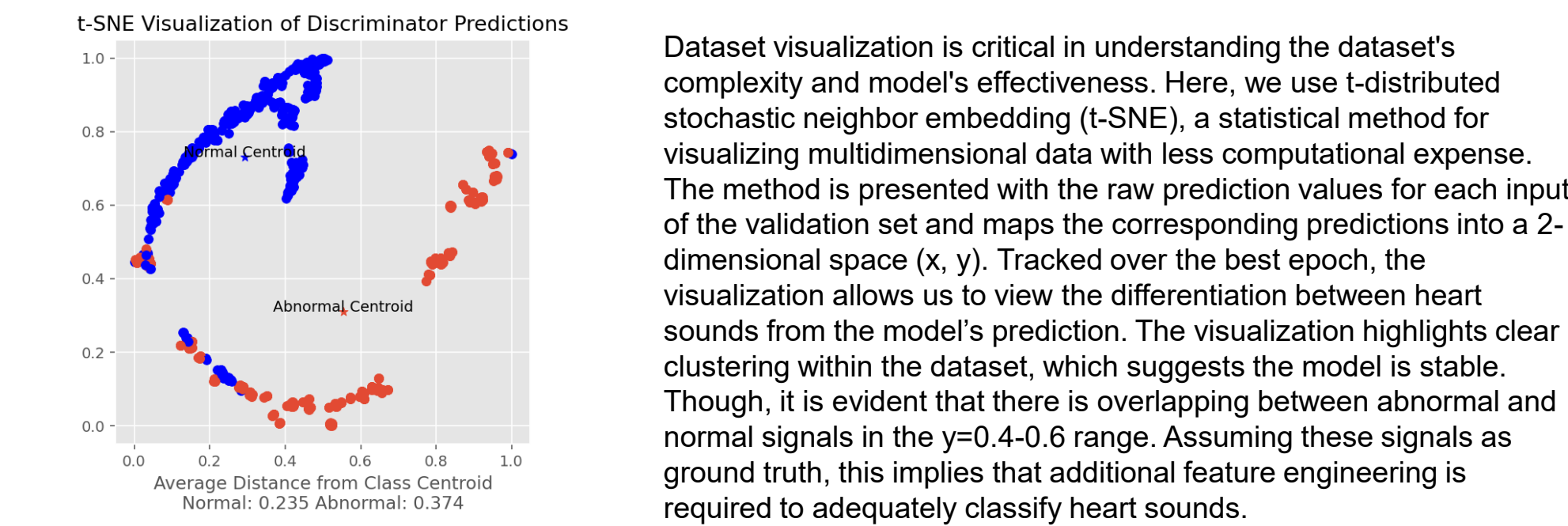
Time Complexity:
 $O(n) = \frac{1}{n} \sum_{i=1}^n m(\log(n) - n_i)$

Real-World Testing & Model Deployment
The model's viability is crucial for ensuring the model's success in the real world. Thus, we need to conduct trials with the end-to-end system to ensure we can deploy and integrate the model with ease.

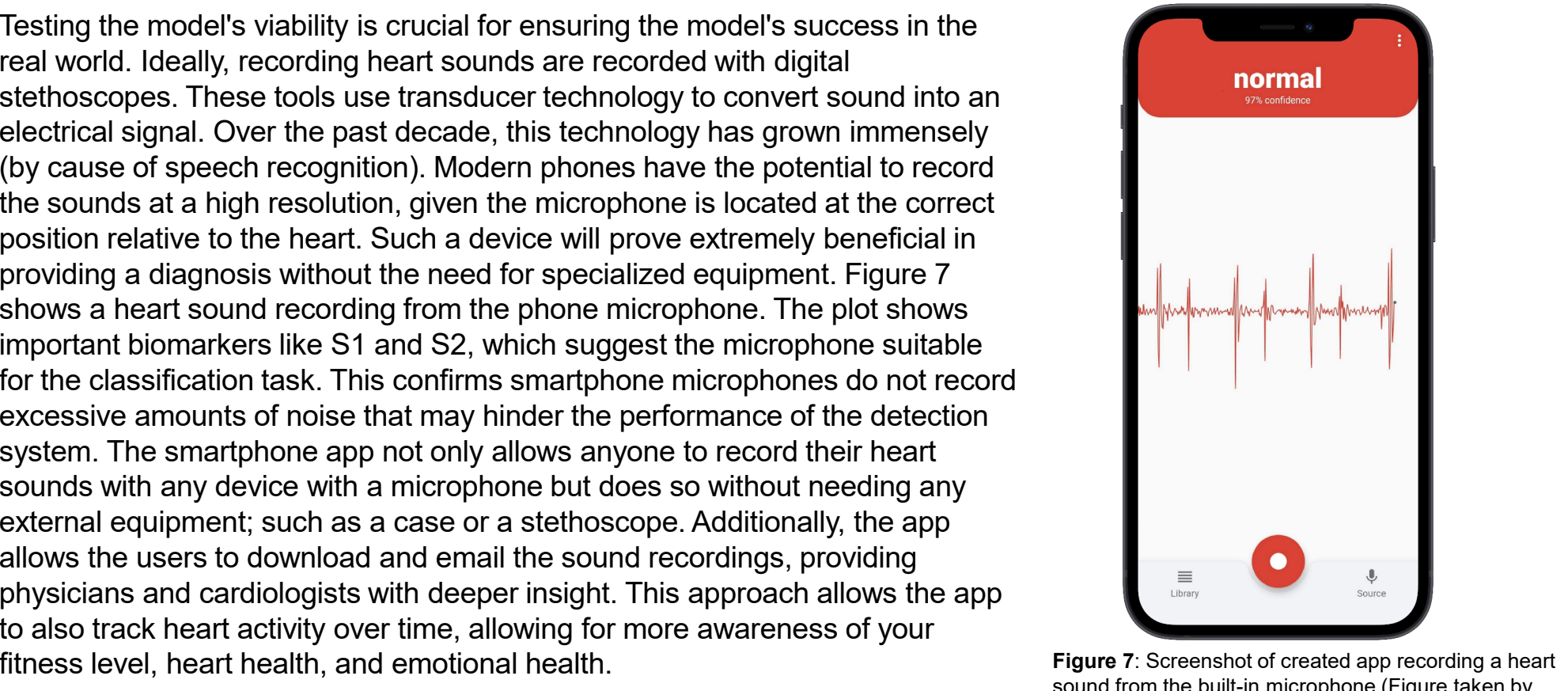
ROC/AUC



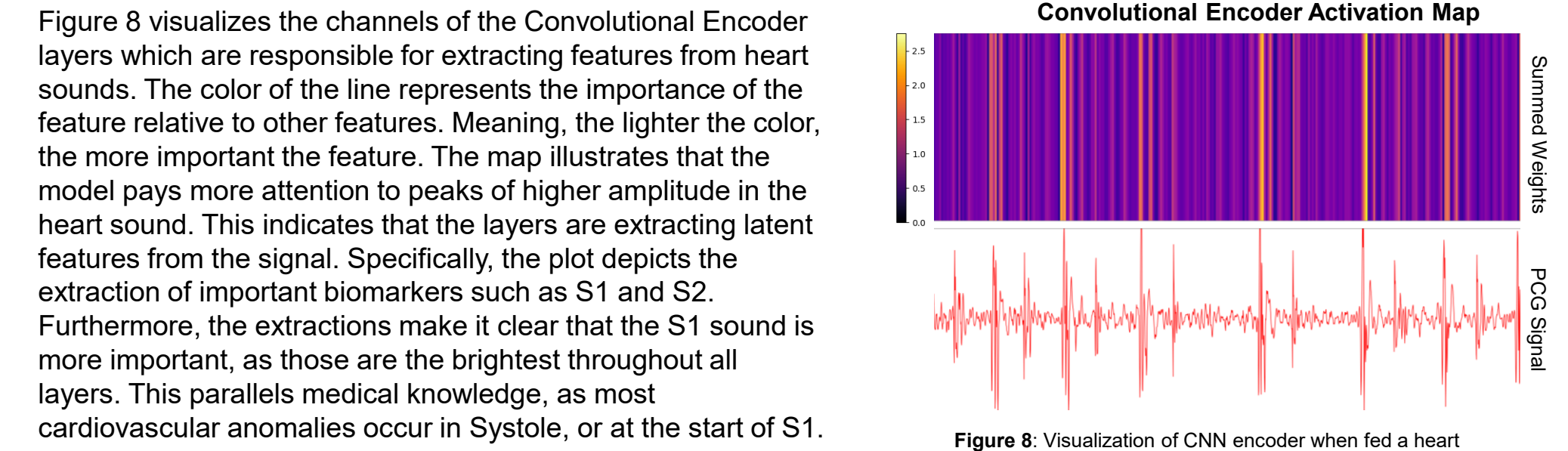
t-SNE Visualization



End-to-End System



Model Interpretation

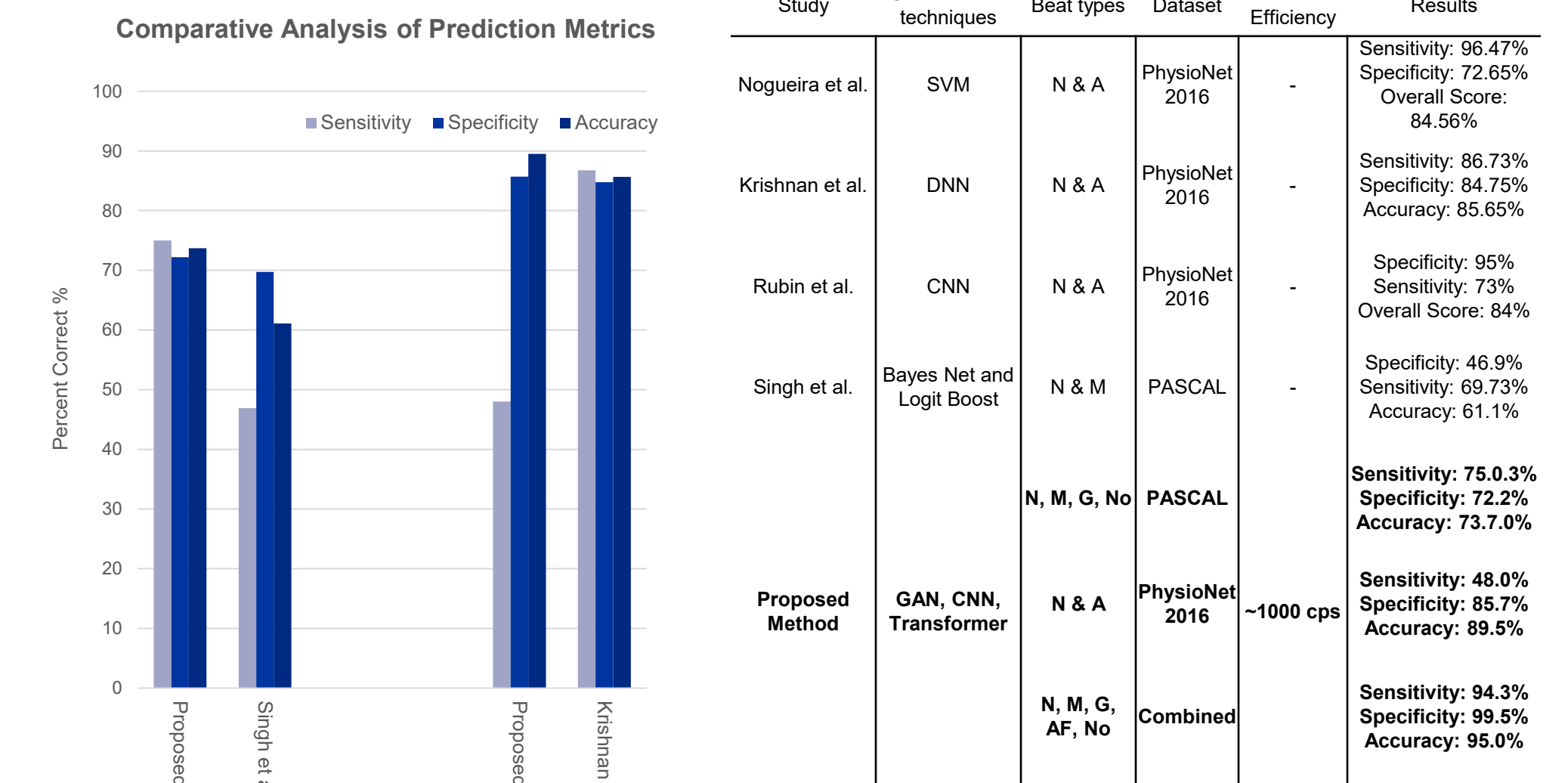


Comparative System Evaluation

Device	Frequency Range (Hz)	Sample Rate (Hz)	Amplification	Cardiac Landmark Guide	Cost
3M Littman 3200	20 – 2000	8000	Up to 24x	No	\$499
Eko Core	20 – 2000	4000	Up to 40x	No	\$349
Jabes	20 – 1000	8000	Up to 20x	No	\$229
Smartphone	20 – 2000	16000-48000	Up to 40x	Yes	n/a

Comparative Model Evaluation

Comparing different methods, we observe the 85.7% was the best accuracy reached on the PhysioNet dataset and 61.1% was the best accuracy reached on the PASCAL dataset. Conversely, our proposed method achieved 89.5% accuracy on the PhysioNet dataset and 73.7% accuracy on the PASCAL dataset. The increase in performance is attributed to our semi-supervised approach, where we used adversarial Conditional Generators to generate heart sounds. This exposed the Discriminator to a wide range of heart sounds which assisted the model in optimizing for generalized features. Additionally, we conducted a statistical significance test (t-test) to show the probability our proposed method's results are due to random chance. The test concluded the results were statistically significant as all p-values were less than 0.05. This implies that the null hypothesis can be rejected, and the results are statistically significant.



Conclusion
We proposed a Generative Adversarial Network (GAN), composed of a Convolution Transformer Generator and a Transformer Discriminator to detect abnormal heart sounds in a recording. The results from model testing and evaluation, along with results from the t-test revealed the proposed method reached better performance than the previous state-of-the-art methods. The introduction of heart sounds analysis with ECGs allowed for increased arrhythmia labels for classification and in a time-efficient manner. Furthermore, the proposed method showed real-world deployment capabilities for autonomous heart sound abnormality detection with recordings collected from a phone microphone.

In terms of future development, we propose conducting prospective clinical trials with patients that have different types of arrhythmias. This will allow us to truly test the generalization capabilities of the model and smartphone app in the real world. Depending on these results, we may opt to develop a low-cost DIY and clinical solution for increased sensitivity in heart recordings. Also, applicable fields include medical emergencies that are time constraint (ER) and developing rural communities that don't have access to arrhythmia expertise.

Applications with the development of the multiview approach include language and time series processing. Specifically, we can train models to convert language to speech and speech to language without the need for a supervised dataset of language A and language B. Rather, the model can be trained to convert language A to an intermediary language (language C), this language can then be converted into language B. Moreover, we can train the model to reconstruct speech recording directly from electrical signals (EEGs) from the auditory cortex or reconstruct vision from the visual cortex.

- Deploying the model with an app that is available to 3rd world countries that can't afford to conduct in-depth testing regularly
- Integrate and serve the model using FastAPI
- Use abnormal heart sound unsupervised datasets as a basis of categorical arrhythmia classification
- Using low dimensional visualization techniques like t-SNE or UMAP
- Cluster data using methods like K means and hierarchical clustering
- Create a classifiable latent representation of PCG signal biomarkers that can be represented with accuracy and precision
- Created by VAE that are fed the PCG signals directly instead of a spectrogram
- Investigate training a heart sound discriminator from generated PCG data from ECG datasets
- Reconstructing speech (wav) recordings from the human auditory cortex (EEG) using techniques used for PCG construction

Future Applications