Two Might Do: A Beat-by-Beat Classification of Cardiac Abnormalities Using Deep Learning with Domain-Specific Features

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Abstract

This paper proposes an efficient convolutional neural network to detect 26 different classes of cardiac activities from different numbers of leads in the Physionet/Computing data in the Cardiology Challenge 2021. The proposed CNN architecture is designed to utilize heart rate variation features from ECG recordings and waveform morphologies of heartbeats simultaneously. Also, the designed architecture is flexible for the implementation of a different number of leads with a varied length of ECG recordings. The proposed algorithm achieved a score of 0.38 using only 2 channels of ECG on all recordings for the hidden test set of the challenge, placing us 21, 20, 19, 20, 20th (Team name: METU-19) out of 39 teams for 12, 6, 4, 3, and 2-leads respectively. These results show the potential of an efficient, flexible novel neural network for beatby-beat classification of raw ECG recordings to a complex multi-class multi-label classification problem.

1. Introduction

Cardiovascular diseases are one of the leading causes of death worldwide [1]. Some works recently indicate that millions of deaths could have been prevented if the cardiovascular disease was detected earlier [2]. The electrocardiogram (ECG) signals are widely used for detecting different cardiovascular diseases, especially the clinical ECG setups are the main tools for detecting abnormal cardiac activity. However, the clinical ECG setup uses 12-lead recordings, making the cardiac monitoring limited outside the clinical hours. To solve this issue, people propose to use wearable devices with limited leads by utilizing the recent advancements in Internet-of-Things (IoT). However, this brings the question of how many leads are sufficient to diagnose a much larger number of cardiac problems? The Physionet/Computing in Cardiology Challenge 2021 [3,4] focus to identify clinical diagnoses from twelve-lead, sixlead, four-lead, three-lead, and two-lead ECG recordings. The differences in performances of the algorithms on the large-scale databases from multi-centers reveal the utility of reduced-lead ECGs compared to standard twelve-lead EGCs and answer the question.

This work proposed a novel, flexible neural network architecture to implement a different number of leads with a varied length of recordings to diagnose cardiac abnormalities using beat morphology and domain-specific features.

2. Methods

The objective is to create a model that can efficiently and accurately classify ECG recordings with different numbers of leads into multiple classes of 30 clinical diagnoses. Four pairs of classes were scored identically. Therefore, we considered these pairs the same and combined identical pairs as one class, so only 26 classes were considered. Although the datasets include varying length ECG recordings and our proposed model is designed to take beats as input and predict 26 different classes from a heartbeat, the overall decision is made using all heartbeats in a recording.

2.1. Preprocessing

The overall challenge dataset contained over 88,000 ECG records with varying lengths for training. Since our method is based on beat-by-beat classification, we have segmented each ECG recordings into beats using the Pan&Tompkins algorithm [5]. We take $F_s/4$ samples before and $F_s/3$ samples after the detected R-peak for segmentation. Additionally, the segmentation window is changed to $F_s/5$ samples before and after when the heart rate is higher than 180 beats per minute to prevent overlapping two heartbeats in the same window.

The sampling rate and the gain value of ECG recordings vary in different datasets. Due to this difference, we have resampled and normalized each extracted beat before feeding them to a classifier. All beats have been resampled to 200 Hz using Fourier resampling. The segmented beats were normalized by their maximum value to have the same signal amplitudes. Also, the Pearson cross-correlation of each extracted beat from the 10-second record is calculated by their linear dependence as Equation 1.

$$\rho_{xy} = \frac{1}{N} \sum_{i=1}^{N} \frac{x_i - \mu_x}{\sigma_x} \frac{y_i - \mu_y}{\sigma_y} \tag{1}$$

where each variable (segmented heartbeats as x and y) has N scalar observations (length of each heartbeat), and μ_x and σ_x are the mean and standard deviation of x, respectively, and μ_y and σ_y are the mean and standard deviation of y. Then using these correlations, the correlation matrix is formed as shown in Equation 2, where the K is the number of segmented heartbeats in a 10-second record.

$$P = \begin{bmatrix} \rho_{11} & \rho_{12} & \dots & \rho_{1K} \\ \rho_{21} & \rho_{22} & \dots & \rho_{2K} \\ \vdots & \ddots & \ddots & \vdots \\ \rho_{K1} & \rho_{K2} & \dots & \rho_{KK} \end{bmatrix}$$
(2)

Finally, each column, which refers to a heartbeat in P, is summed and normalized with the maximum value and compared with a threshold value found empirically and set to 0.8 value. If the normalized correlation coefficient of a segment is lower than 0.8, this segmented section is removed from the recording without further procession. This last step, based on cross-correlation of segmented beats, in preprocessing acts as a control mechanism to avoid a wrongly segmented heartbeat feeding as input to the classifier. Although this approach assumes that the performance of the peak detection algorithm is highly accurate and detects most of the correct peaks in the recording, it is observed that this preprocessing step reduces the number of wrong segments in a recording.

2.2. Domain-Specific Feature Extraction

When we segment the ECG records into heartbeats and feed them to a classifier, it is observed that the heart rate variability information is lost, resulting in a significant performance decrease for some cardiovascular diseases, which are characterized by heart rhythm such as atrial fibrillation, bradycardia. To prevent this performance decrease for some classes, we have calculated the heart rate for each beat in all recordings using the following equation:

$$HR_{(i)} = \frac{60}{(R_{(i)} - R_{(i-1)})/F_s}$$
(3)

Where $HR_{(i)}$ is the heart rate of the i^{th} heartbeat segment in beat per minute unit. $R_{(i)}$ is the location of the R-peak in the i^{th} heartbeat segment, and F_s is the sampling rate of this record. Then, using the calculated $HR_{(i)}$ values, an HR array (\overrightarrow{HR}) is formed by adding them one after another, as shown in Equation 4.

$$\overrightarrow{HR} = [HR_0, HR_1, HR_2, ..., HR_i]$$
(4)

Finally, the heart rate variability (HRV) features are calculated from lead I and integrated into the designed deep learning model.

- Mean of $H\dot{R}$
- Median of $H_{\mathcal{K}}$
- Range of \overrightarrow{HR}
- Standard deviation of \overrightarrow{HR} values
- Root mean square of $H\dot{R}$ values
- pNN20

(percentage of HR intervals differing more than 20 ms)

In total, six different HRV features are extracted from a 10-second recording to be used for classification. Since these features are obtained using the complete recording instead of a single heartbeat and our proposed model is based on the beat-by-beat classification, we have used the extracted feature with all heartbeats simultaneously at the neural network. Although, at first glance, it seems a overfitting problem due to the same features are used with the different heartbeats in the same recording. It is observed that it works as a regularizer instead of resulting in overfitting.

2.3. Classifier

The proposed convolutional neural network classifies the segmented heartbeat into 26 different classes using the extracted HRV features from a 10-second recording. The proposed CNN architecture is designed to be flexible with the different number of leads. The CNN architecture for the heartbeat classification is designed both utilizing the residual connections introduced by He et al. in [6] and the dropout [7] that has been shown to achieve good performance for increasing the generalization ability of the neural networks. Especially, it is noted that the CNN architecture classification performance increases approximately 3-5% when the dropout with 0.1 is used at the lead ways as shown in Figure 1.

2.4. Network Training

L2-regularization with 0.0002 is applied for each convolution layer. We used the Adam optimizer [8] with the default parameters $\beta_1 = 0.9$ and $\beta_2 = 0.999$, and a mini-batch size of 90. The learning rate is initialized to 0.001 and reduced by a factor of 10 when the validation F1 score stopped improving for 75 consecutive epochs. The training continues until 100 successive epochs without performance improvements with a maximum of 500 epochs. 10% of training records were used for validation to implement early-stopping based on the F1 score.

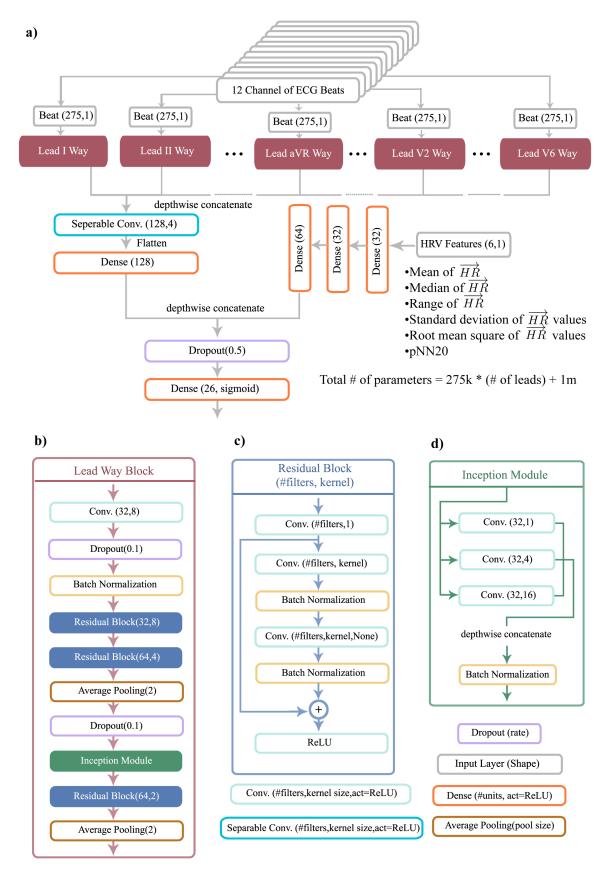


Figure 1: a) Complete model architecture, b) Channel block, c) Residual block, d) Inception block

3. **Results**

After training, the model predicts scores $p \in [0, 1]$ for each class. To binarize these scores, we have applied a threshold which is found empirically, and set it to 0.25 for all classes. As our model takes beats as input instead of recordings, the prediction outputs of beats in a record are combined into a single score for each class similar to our correlation matrix calculation. However, different from the correlation matrix, each row represents a beat in a recording, and each column refers to the scored classes in the challenge. After creating this scoring matrix for each heartbeat, the columns are summed and normalized with the maximum value. Finally, the normalized scores are compared with the threshold value and set to 1 if it is higher than the threshold.

Table 1: Challenge scores for our final entry on the hidden validation set, and the hidden test set with the ranking.

Leads	Validation	Test	Ranking
12	0.57	0.38	21
4	0.57	0.38	19
2, 3, 6	0.57	0.38	20

Table 1 shows the Challenge scores for our final entry on the hidden validation and test set with rankings.

Table 2.	10 - fold	cross-validation results
Table 2:	10-1010	cross-vanuation results

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Database	AUROC	AUPRC	F-measure	Challenge Score
G12EC [3]	0.83	0.42	0.46	0.51
CPSC [9]	0.79	0.28	0.42	0.76
PTB-XL [10]	0.87	0.47	0.46	0.50

The results for the different sets are evaluated using the 10-fold cross-validation and showed in Table 2. Although the proposed method achieves high challenge score on the CPSC dataset, its performance decreases severely for the G12EC and PTB-XL. This fluctuation in the performance shows that the model's generalization capability heavily depends on the datasets' characteristics.

4. Discussion and Conclusion

This paper proposed a flexible CNN architecture for classifying raw ECG recordings with varying lengths and different numbers of leads. The designed CNN is not only utilized the heartbeat morphology but also employed the HRV features simultaneously to classify recordings. Also, novel preprocessing and postprocessing methods are introduced in this paper to integrate beat-by-beat classification to long, varying length ECG recordings. However, despite some promising results, questions remain about why the proposed approach's performance has severely fluctuated for different datasets. An explanation for these fluctuations can be the biases and quality of the recordings in the datasets. For example, if the extracted features from a class are calculated due to false peak detection, the performance decrease is inevitable since the features and segments are fed together with beats.

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References

- Heart Disease and Stroke Statistics—2020 Update: A Report From the American Heart Association. Circulation March 2020;141(9):e139–e596.
- [2] Rashid N, Al Faruque MA. Energy-efficient real-time myocardial infarction detection on wearable devices. In 2020 42nd Annual International Conference of the IEEE Engineering in Medicine Biology Society (EMBC). 2020; .
- [3] Alday EAP, Gu A, Shah AJ, Robichaux C, Wong AKI, Liu C, Liu F, Rad AB, Elola A, Seyedi S, Li Q, Sharma A, Clifford GD, Reyna MA. Classification of 12-lead ECGs: the PhysioNet/Computing in Cardiology Challenge 2020. Physiological Measurement January 2021;41(12).
- [4] Reyna MA, Sadr N, Perez Alday EA, Gu A, Shah A, Robichaux C, Rad BA, Elola A, Seyedi S, Ansari S, Ghanbari H, Li Q, Sharma A, Clifford GD. Will Two Do? Varying dimensions in electrocardiography: the PhysioNet/Computing in Cardiology Challenge 2021. Computing in Cardiology 2021;48:1–4.
- [5] Pan J, Tompkins WJ. A real-time qrs detection algorithm. IEEE Transactions on Biomedical Engineering 1985;BME-32(3):230–236.
- [6] He K, Zhang X, Ren S, Sun J. Deep residual learning for image recognition. 2016 IEEE Conference on Computer Vision and Pattern Recognition CVPR 2016;770–778.
- [7] Srivastava N, Hinton G, Krizhevsky A, Sutskever I, Salakhutdinov R. Dropout: A simple way to prevent neural networks from overfitting. Journal of Machine Learning Research 2014;15(56):1929–1958.
- [8] Kingma DP, Ba J. Adam: A method for stochastic optimization. CoRR 2015;abs/1412.6980.
- [9] Liu FF, Liu* CY, Zhao LN, Zhang XY, Wu XL, Xu XY, Liu YL, Ma CY, Wei SS, He ZQ, Li JQ, Kwee NY. An open access database for evaluating the algorithms of ecg rhythm and morphology abnormal detection. Journal of Medical Imaging and Health Informatics 2018;8(7):1368–1373.
- [10] Wagner P, Strodthoff N, Bousseljot RD, Kreiseler D, Lunze FI, Samek W, Schaeffter T. PTB-XL, a large publicly available electrocardiography dataset. Scientific Data 2020; 7(1):1–15.

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