A Study on Several Critical Problems on Arrhythmia Detection using Varying-Dimensional Electrocardiography

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Abstract.

As a major tool for the diagnosis of heart diseases, the electrocardiography (ECG) has long been an active research field among medical researchers, especially the standard 12-lead ECGs due to its clinical significance. Although many excellent algorithms have been developed and a number of successful systems have already been applied in clinical, there are still fundamental problems that have not yet been thoroughly studied and fully answered.

The PhysioNet/Computing in Cardiology Challenge 2021 (CinC2021) raised several such critical problems: can subsets of the standard 12 leads provide models with adequate information to give comparable performances for classifying ECG abnormalities? can models be designed to be effective enough to classify a broad range of ECG abnormalities? In this work, we present our effort on tackling these problems. Through extensive searching, we (team name "Revenger") discover several novel architectures within the framework of convolutional recurrent neural networks (CRNNs) that provide such effectiveness, and moreover they provide comparable performances on reduced-lead ECGs, even in the extreme case of 2-lead ECGs. In addition, we propose a "lead-wise" mechanism to facilitate parameter reuse of ECG neural network models. This mechanism largely reduces model sizes while keeping comparable performances. To further augment model performances on specific ECG abnormalities and to improve interpretability, we manually design auxiliary detectors based on clinical diagnostic rules.

To conclude, our work gives positive answers to the critical questions CinC2021 raises and lays solid foundation for further research in the future on these topics.

Keywords: multi-dimensional ECG, reduced leads, deep learning, neural architecture search, clinical rule based detector

1. Introduction

Electrocardiography (ECG), as a physiological signal that records the electrical activity of the myocardium, is widely adopted in clinic for screening heart diseases, the worldwide leading death cause [1]. Despite the rapid growth of the application of ambulatory ECG and other methods, the standard 12-lead ECG is still the most widely accepted in the clinical practice. However, the 12-lead ECG equipment for real-time monitoring of the cardiac electrical activities, especially the 24-hour Holter monitoring systems, usually collect an enormous amount of data. Therefore, how to properly process these ECG signals, so as to be rapid, accurate and complete for the diagnosis of a range of cardiac abnormalities as broad as possible, is still an urgent problem that needs to be solved.

For this reason, many algorithms have been proposed to accurately and automatically assist in the diagnosis of cardiac electrical abnormalities. Among them, deep neural networks (DNNs) have achieved great success [2–4] in recent years and have been dominating in this research field. These models are claimed to be able to achieve very high precision, even comparable to cardiologists [2]. However, these work have their drawbacks. For example, their models have only been validated on just a few typical ECG arrhythmias (for example atrial fibrillation, ventricular tachycardia, etc. [2]). Data redundancy is another aspect that these work did not consider [3, 4]. Hence the problem raised in the begging of the paper still needs to be studied in more depth.

In this paper, our effort of tackling this problem, which uses DNNs in combination with clinical rules providing better interpretability, will be described. Following the mission of The PhysioNet/Computing in Cardiology Challenge 2021 (CinC2021) [5–8], we apply our hybrid method on the problem of classifying a very broad range of cardiac abnormalities, and using reduced-lead ECGs which largely reduce the data amount while at the cost of only holding partial information of the standard 12-lead ECGs. The 5 lead-sets reduced from the standard 12-lead are listed as follows.

• Twelve leads: I, II, III, aVR, aVL, aVF, V1, V2, V3, V4, V5, V6

• Six leads: I, II, III, aVR, aVL, aVF

Four leads: I, II, III, V2Three leads: I, II, V2

• Two leads: I, II

Our findings will be introduced in more details in the rest of the paper, which is organized as follows: Section 2 explains the selection of data processing methods, including methods of data preprocessing and data augmentation. Section 3 introduces the deep learning models with novel network architectures. Section 4 that follows discusses the training setups for the neural network models, including the selection of loss functions, optimizers and learning rate schedulers, etc. In order to assist the deep learning models, we design several detectors based on clinical rules for easily distinguishable abnormal ECG signals in section 5. The performances of our solution to the CinC2021 challenge problems are summarized in section 6. Section 7 contains our

thorough ablation studies for the selection of model structures, loss functions and data augmentation methods, etc. Section 8 concludes this work.

2. Data Processing

Most ECG abnormalities are characterized by their subtle changes both morphological and spectral, while only a minority can be described easily by clinical diagnostic criteria, for example bradycardia, low QRS voltage, etc. DNNs which are particularly suitable for capturing such subtle structures in a neat and uniform way become the de facto default method for various ECG tasks, as stated in the previous section. It is commonly believed that DNNs are robust to "natural"; noises, hence require little preprocessing as opposed to traditional signal processing methods. Instead of traditional preprocessing techniques, the deep learning research community tend to use data augmentation techniques to broaden the distribution of the training data, which in theory help improve performances of DNNs on unseen data. However, to the authors' knowledge, very few numerical evidences can be found in ECG deep learning research work on this topic. Therefore, we include both traditional preprocessing techniques as well as data augmentation techniques which we think would be helpful and suitable for developing our deep neural networks. A simple numerical comparison will be given in Section 7.1. Since we also adopt clinical rule based detectors, hence the inclusion of traditional preprocessing is reasonable.

2.1. Data Selection

The main data resource of this study for conducting the experiments is the public training database of CinC2021 challenge [8], excluding the StPetersburg subset (the INCART dataset [9]) since the granularity of its labels is too coarse, being not suitable for training neural networks. Beside the INCART dataset, the public part of this database contains 4 other subsets, namely

- CPSC Database and CPSC-Extra database [10]
- PTB [11] and PTB-XL [12] database
- Georgia 12-lead ECG Challenge (G12EC) database [7]
- Chapman-Shaoxing [13] and Ningbo [14] database

There are totally 85056 records in 132 classes (abnormalities) in the CinC2021 database. We use only the 26 scored classes (equivalent classes counted as 1 class) in the challenge which cover 92.46% of the whole database.

2.2. Preprocessing Procedures

As our initial step of the data processing pipeline for training the neural networks, all ECGs are resampled to 500 Hz if necessary. Resampled ECGs are further cropped

‡ in contrast to man-made adversarial attack

or padded by zeros to a length of 10 seconds (5000 sample points) in order to utilize mini-batch training.

It is a common practice for traditional ECG signal processing to apply band-pass filters to the ECGs to remove noises as well as coarse baseline drifts. Sometimes an additional median filter is used to further remove the baseline. Another advantage of band-pass filtering is that such operations can largely bridge the potential gap between training and inference data distributions for DNNs. In almost all our experiments, we take band-pass filtering, using a finite impulse response (FIR) filter with pass-band 0.5 Hz - 60 Hz, as a part of the data processing pipeline. To validate its efficacy for DNNs, experiments are performed as a part of our ablation study in Section 7.1.

Another preprocessing procedure we take is the z-score normalization defined as

$$\frac{\mathbf{x} - \text{mean}(\mathbf{x})}{\text{std}(\mathbf{x}) + eps} \cdot s + m \tag{1}$$

where \mathbf{x} is the ECG signal, s, m, eps are fixed values. In our experiments, we take $s=1, m=0, eps=10^{-7}$. The small value eps is added to avoid division by zero for ECG records with constant values. After this transformation, the ECG will have mean m and standard deviation s (except for constant value ECGs). Numerical results verifying the validity of z-score normalization are also presented in Section 7.1. As z-score normalization has similar effect to a median filter, the latter is often omitted when preparing data for deep learning models.

2.3. Data Augmentation

To alleviate overfitting, data augmentation techniques including mixup [15], random masking [3], etc. are adopted stochastically (with certain probability) on the training data for the neural networks. Mixup performs convex linear combination of the training data as follows

$$\mathbf{x} = \alpha \mathbf{x}_1 + (1 - \alpha) \mathbf{x}_2, \mathbf{y} = \alpha \mathbf{y}_1 + (1 - \alpha) \mathbf{y}_2,$$
 (2)

where $0 < \alpha < 1$, $\mathbf{x}_1, \mathbf{x}_2$ are preprocessed ECGs, and $\mathbf{y}_1, \mathbf{y}_2$ are corresponding labels. No more augmentations, like random flip, are done, since they might completely change the interpretation of the standard-lead ECGs§, as will be seen in Section 5.

For dealing with the potential problem of overconfidence of the models, we adopt the label-smoothing regularization (LSR) technique [16] defined as

$$\mathbf{y}' = (1 - \varepsilon)\mathbf{y} + \frac{1}{K}\varepsilon\mathbf{1},\tag{3}$$

where \mathbf{y} be the one-hot label vector, K is the number of classes, $\mathbf{1}$ is the K-dimensional vector filled with ones, and $\varepsilon \in [0,1]$ is a weight factor. LSR generates the "soft label" \mathbf{y}' from the "hard label" \mathbf{y} . In our experiments, we fix $\varepsilon = 0.1$.

§ which might not be the case for wearable dynamic ECGs

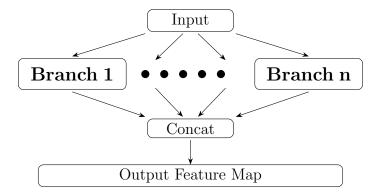


Figure 1: Sketch of the architecture of the multi-branch CNN family.

Similar to preprocessing procedures, we performed experiments validating the impact on model performances of the augmentation techniques in Section 7.1. The selection of an optimal data processing pipeline serves as a part of our post challenge work.

3. Neural Network Architectures

3.1. Convolutional Recurrent Neural Networks (CRNNs)

Inspired by previous work [3, 17], we base our pipeline for solving ECG classification problems on a CRNN framework. A CRNN consists of a CNN backbone extracting features from input ECGs. Following the CNN backbone is an optional RNN module (e.g. LSTM [18]) and an optional attention module (e.g. squeeze-and-excitation (SE) [19], global context (GC) [20]) to refine the feature maps, modeling long-range dependencies and making use of the sequence properties of ECGs. Thus obtained feature maps are fed into a multi-layer (typically 1 or 2 layers) perceptron (MLP) head to produce the vector of probabilities for each class.

The CNN backbone is the core of this CRNN framework and has been extensively studied by the ECG research community. Variants [2–4] from computer vision models, as well as networks distinctively designed for ECGs [17], have enjoyed great success in many ECG tasks. Design and selection of CNN backbones will be the central issue for this study.

3.2. Multi-branch CNNs

The first set of CNN backbones we use for solving the CinC2021 challenge problem are derived from the multi-branch CNNs proposed in [17], whose general architecture in depicted in Figure 1. This type of network has several branches, with each branch containing sequentially stacked convolutional layers, and with normalization layers, activation layers and dropout layers inserted after specific convolutional layers. The essence of this network family is that each branch uses dilated convolutions with different

	Kernel Sizes	Dilations
Branch 1		(1,1,1,1,1,1)
Branch 2	(11, 7, 7, 5, 5, 5)	(2, 2, 4, 8, 8, 8)
Branch 3		(4,4,8,16,32,64)

Table 1: Kernel sizes and dilations for the convolutions for a typical example of the CNN described in Figure 1. It has three branches, each with 6 convolutional layers. We used this CNN backbone in our official phase submission entry for the CinC2021 challenge.

dilation ratio, so that the receptive field of each branch matches specific waveforms of the ECG. The idea behind is similar to the ASPP (Atrous Spatial Pyramid Pooling) head originally proposed in DeepLabV2 [21], directly modelling the spectral characteristics. We used a 3-branch CNN backbone in our official phase submission entry for the CinC2021 challenge, whose key hyperparameters are listed in Table 1. This multibranch CNN serve as the CNN backbone for our final submission for the official phase of the CinC2021 challenge.

3.3. "Lead-wise" CNNs

Another novel structure of the CNNs we design during developing our solution to the CinC2021 challenge problem is the "lead-wise" convolution, which is implemented via grouped convolutions so that the number of groups are divisible by the number of leads of the input ECGs. Another key point is that normalization layers should use group normalizations [22] instead of batch normalization. Using this mechanism, training one model on multi-lead ECGs is equivalent to training multiple models on single-lead ECGs. This enables parameter reuse for training models on reduced-lead ECGs from models trained on the standard 12-lead ECGs, in which case one freezes the parameters of the CNN backbone and fine-tunes the rest part of the network. The "lead-wise" CNNs have significantly less parameters compared to their "normal" counterparts, while offering competitive performances. This would be shown via experiments in Section 7.2.

3.4. Variants of ResNet

For the purpose of searching for more effective model architectures other than the networks we used for the CinC2021 challenge, we experiment with various ResNet variants (e.g. [2, 4, 23, 24]) as CNN backbone for our neural networks. This constitutes a large part of post challenge work of us.

The original ResNet was proposed in [25] in 2016, and quickly became the most widely used neural network architecture. The TResNet [24] variants currently are even still almost the state-of-the-art (SOTA) models for multi-label classification on the MS-COCO dataset [26]. Until recently, variants of ResNet [2, 4] have long been dominating the ECG deep learning research community.

$$Input \longrightarrow \boxed{Stem} \longrightarrow \boxed{4 \text{ Stages}} \longrightarrow Feature \text{ Map}$$

Figure 2: A typical architecture of ResNet. The Stem typically consists of convolutions with large kernel sizes to capture coarsely-level features and down-sampling layers to reduce computations. The 4 stages consists of stacks of units sketched in Figure 3 producing the feature maps for final classification.

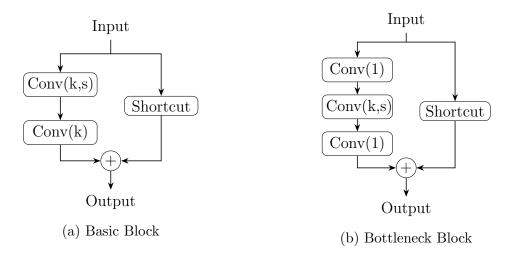


Figure 3: Sketch of 2 types of building blocks for ResNet. Conv(k,s) is convolution of kernel size k, and stride s; Conv(1) is convolution of kernel size 1 and stride 1. When s=1, the shortcut is just the identity map; when s>1, shortcut need to do down-sampling. When the main stream (the path passing from Input to Output through the convolutions) raises the number of channels, shortcut also needs to use convolutions of kernel size 1 to raise to the same number of channels. Other layers including activations, normalizations and dropouts are not plotted in this Figure. In (b), it is actually from the ResNet-B variant, rather than the original one.

A typical architecture of ResNet consists of an input stem, followed by 4 stages, as depicted in Figure 2. The stem consists of convolutions with large kernel sizes and with [23–25] or without [4] down-sampling layers. The 4 stages consist of different numbers of stacked building blocks of basic type (Figure 3a) or of bottleneck type (Figure 3b). Attention mechanisms, which are aforementioned in Section 3.1, are usually integrated into these building blocks to further improve the performance of the networks.

The SE-variant (with SE attention integrated in the building blocks) of the ResNet model proposed in [4] will serve as the baseline for our comparative studied. This baseline will be denoted as ResNet-NC-SE ("NC" is short for "Nature Communications"). Its bottleneck variant, denoted as ResNet-NC-BS, along with another variant ResNet-NC-BG, obtained by replacing the SE attention module with GC attention module in ResNet-NC-BS, are also experimented.

Another series tested is the TResNet family [24]. TResNets adopted many new techniques for acceleration and/or improving prediction accuracy and robustness. These

Network	Stage1		Stage2		Stage3		Stage4	
Network	rpt, type	chan						
ResNet-NC-SE	1 Ba-SE	128	1 Ba-SE	192	1 Ba-SE	256	1 Ba-SE	320
ResNet-NC-BS	1 Bot-SE	512	1 Bot-SE	768	1 Bot-SE	1024	1 Bot-SE	1280
ResNet-NC-BG	1 Bot-GC	512	1 Bot-GC	768	1 Bot-GC	1024	1 Bot-GC	1280
TResNet-N	2 Ba-SE	56	2 Ba-SE	112	2 Bot-SE*	896	2 Bot	1792
TResNet-P	1 Ba-SE	56	1 Ba-SE	112	1 Bot-SE*	896	1 Bot	1792
TResNet-F**	1 Ba-SE	32	1 Ba-SE	64	1 Bot-SE*	512	1 Bot	1024

Table 2: Architectures of Variants of ResNet. Types and number of repeats and output channels of the building blocks used are listed. "Ba" refers to the Basic type building block, "Bot" refers to the Bottleneck type building block.

^{**} Convolutions are separable convolutions.

Network	Ker	nel Size	Down-Sampling		
Network	Stem	4 Stages	Stem	4 Stages	
ResNet-NC-SE	17	17	1	(4, 4, 4, 4)	
ResNet-NC-BS	17	17	1	(4, 4, 4, 4)	
ResNet-NC-BG	17	17	1	(4, 4, 4, 4)	
TResNet-N	1	15	4	(1, 2, 2, 2)	
TResNet-P	1	17	4	(1, 2, 2, 2)	
TResNet-F	1	17	4	(1, 2, 2, 2)	

Table 3: Details of the kernel sizes and down-sampling ratios of the stem and the building blocks of the 4 stages in the ResNet variants.

techniques include space-to-depth (S2D) [27] stems, anti-aliasing (AA) down-sampling [28]. Since the smallest network (the TResNet-M) proposed therein has 21 building blocks, which is much too large for ECG processing, we design and experiment with 3 smaller variants, namely TResNet-N (nano), TResNet-P (pico), TResNet-F (femto). More details on architectures of the above networks are gathered in Table 2 and Table 3. Experiment results will be presented in Section 7.

4. Training Setups

The whole neural network development and validation work are done on a GPU server with 8×RTX3080. In most cases, we set batch size to be 64 and set the maximum number of epochs to be 50. 20% of the training data is left out for model evaluation and model selection. In most cases throughout this paper, when referring to "validation set", we mean this left-out set from the whole of the public training set, rather than the hidden validation set of the CinC2021 challenge. Early stopping is triggered if the

^{*} SE module follows the 2nd convolution of the block, the rest follows the last convolution of the block.

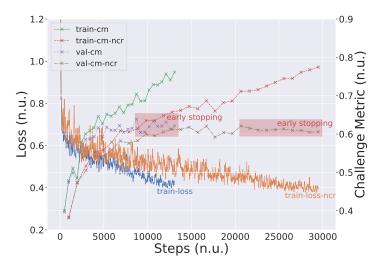


Figure 4: Two typical training processes using weighted BCE loss for neural networks with lead-wise branched CNN backbone. The suffix "ncr" refers to the challenge solution pipeline with **no clinical rule** based detector (ref. Section 5), in which case the whole 26 classes are included for training. "cm" is the abbreviation of the **challenge metric**.

challenge metric on the validation set does not grow for 8 epochs. Two typical training processes are depicted in Figure 4.

4.1. Loss Functions

Since the challenge data is highly unbalanced, having a long tail distribution, we tested 2 loss functions aiming at dealing with such data distribution. Initially, we used weighted binary cross entropy (BCE) as the loss function with weights inversely proportional to the number of records of the classes. After studying other challengers' solutions after the CinC2021 challenge official phase, we found that the asymmetric loss [29] is more widely adopted. The asymmetric loss is defined as follows

$$ASL = \begin{cases} L_{+} := (1 - \mathbf{p})^{\gamma_{+}} \log(\mathbf{p}), \\ L_{-} := (\mathbf{p}_{m})^{\gamma_{-}} \log(1 - \mathbf{p}_{m}), \end{cases}$$
(4)

where $\mathbf{p}_m = \max(\mathbf{p} - m, 0)$ is the so-called shifted probability, with probability margin m. The loss on one label \mathbf{y} of one sample is

$$L = -yL_{+} - (1 - y)L_{-} \tag{5}$$

All multiplications above are Hadamard products (element-wise products). By using this asymmetric loss, one is able to emphasize the contribution of positive samples or negative samples by adjusting the ratio of the focusing parameters γ_+ to γ_- . Typically in our experiments, γ_+ is fixed to be 0, and we set $\gamma_- = 0.2$. We observed augments of challenge metric by 0.03-0.05 on the validation set, which is very significant. More details can be found in Section 7.1. Therefore the asymmetric loss is our final choice of

the loss function for training the neural network models. Although the validation and selection of loss functions is only a small part of our post challenge work, it offers the largest augmentation of the model performances.

4.2. Optimizers and Learning Rate Schedulers

Parameters of neural network models are optimized using the AMSGrad variant of the AdamW optimizer (denoted as adamw_amsgrad) [30]. For learning rate, which is the most important hyperparameter of an optimizer, we use the OneCycle scheduler [31] with maximum learning rate 0.002 and cosine annealing strategy to adjust learning rate during training. It was reported in the deep learning research community that the combination of adamw_amsgrad optimizer with OneCycle scheduler gives the best performance most of the time. For the purpose of comparative study, we also conduction experiments without any learning rate schedulers starting from an initial learning rate of 0.001, letting the optimizer adamw_amsgrad do self-adaptation of the learning rate during training. Corresponding results will be gathered in Section 7.1.

4.3. Model Inference and Model Selection

Model inference is done in a self-adaptive way. More precisely, to make binary predictions from the probability vector, a hard threshold 0.5 is used. If none of the probabilities exceeds 0.5, then the class with the highest probability would be chosen as the binary output, along with other classes that are close enough in probability with the highest one (the difference < 0.03).

The monitor for model selection is the challenge score computed on the left-out validation set.

It should be emphasized that the platform (computing resources) of the CinC2021 challenge official submission system is different from the system that we mainly work on as described in the begging of this section, the batch size and learning rates are adjusted for the challenge submissions. The principles for adjustment are discussed in Section 7.1.

5. Clinical Rule Based Detectors

In order to assist the deep learning models, as well as to improve interpretability, we design clinical rules based detectors for several ECG abnormalities. From the authors' previous research experience as well as experiences in designing industrial ECG auxiliary diagnosis systems, rule-based post-processing is often a helpful supplement to machine learning models. We list a few of them as follows:

1. "LAD" (left axis deviation) and "RAD" (right axis deviation): these two abnormalities are detected by checking the positivity of the QRS complexes in leads I, II, aVF (the "3-lead" method) or in leads I, aVF (the "2-lead" method) as in [32]. More precisely

class	TP	FP	TN	FN
brady*	18905	5342	54403	306
LAD	6547	13547	57778	1084
RAD	927	3723	73953	353
LQRSV	787	2798	74671	700
PR	671	55	77421	809

Table 4: Confusion matrices of clinical rule based detectors on 5 typical classes in the CinC2021 challenge database. All 12 leads are used. Abbreviations are as follows: "brady" for bradycardia, "LAD" for left axis deviation, "RAD" for right axis deviation, "LQRSV" for low QRS voltage, "PR" for pacing rhythm; "TP" for true positive, "FP" for false positive, "TN" for true negative, "FN" for false negative.

* we merge the 2 classes "Brady" and "SB" (sinus bradycardia) in the CinC2021 challenge database into one for applying clinical rule based detectors.

- "2-lead" method:
 - "LAD": lead I is positive; lead aVF is negative;
 - "RAD": lead I is negative; lead aVF is positive.
- "3-lead" method:
 - "LAD": lead I is positive; lead II, aVF are negative;
 - "RAD": lead I is negative; lead II, aVF are positive.

the "2-lead" method is simpler, but might have false positives.

2. "LQRSV" (low QRS voltage): this abnormality is directly related to the absolute values of the ECGs, hence is distinctive from almost all of the rest ECG abnormalities. The detection is done by checking peak-to-peak amplitudes of the QRS complexes in the limb leads (I, II, III, aVR, aVL, aVF), or in the precordial leads (V1-V6). If the QRS detection were to fail, then peak-to-peak amplitudes are computed as the amplitudes of sliding windows of 0.12 second length, which is almost the duration of a normal QRS complex.

It should be emphasized that the a large proportion of clinical rule based detectors rely heavily on the quality of R peak detection, which is another research topic other than classification for ECG that we do not treat in this work.

Confusion matrices of clinical rule based detectors on 5 typical classes in the CinC2021 challenge database, using the standard 12-lead ECGs, is collected in Table 4. Many of the false positives are not truly false positives, as would be discussed in Section 8, and illustrated with examples in Appendix A.

It can be inferred from the clinical rules that a small proportion of ECG arrhythmias can not be reasonably detected under certain combination of reduced leads, resulting in potential drop in model performances, as would be illustrated in Section 6.

6. Results

In this section, our major findings for the problems raised in the begging of this work are presented.

6.1. Data Redundancy

For studying the problem of data redundancy of the standard 12-lead ECGs, we adopt the baseline model with the CRNN architecture with the following (sequential) components:

- CNN backbone: ResNet-NC-SE;
- RNN module: stack of 2 bidirectional LSTM layers with hidden size 192;
- Attention module: SE module with reduction ratio 8;
- MLP head: stack of 2 linear layers with 1024 intermediate features.

This baseline model is newly adopted in the post challenge work, instead of the model used for the CinC2021 challenge official phase. Note that in [4], the feature map from the CNN backbone are flattened (which can be called "depth to space") to feed into a linear layer for final prediction, while we just use a global max pooling layer before the linear layers.

The curves of challenge scores for this set of experiments are plotted in Figure 5. We find that the drop from using all the 12 leads to using only 2 of them for making ECG arrhythmia classifications is very slight (only 0.036 in terms of challenge score) and is reasonably acceptable. Another interesting phenomenon is that the model performance is higher on the 3-lead-set (I, II, V2) than on the 4-lead-set (I, II, III, V2), although the former is a subset of the latter. This phenomenon has already been observed in our CinC2021 challenge official phase solution, as well as other participants and the challenge organizers. The extra information from lead III seems to be harmful to the neural networks.

6.2. CNN Backbone Effectiveness

To verify the effectiveness of the CNN backbones, we remove extra components from the CRNN framework, leaving only the CNN backbone along with a linear layer for producing the probability vector. The CNN backbones we compare include 3 variants of [4] and 3 variants of TResNet as listed in Table 3, as well as the 3-branch CNN as described in Table 1 and its "lead-wise" variant which we used in the CinC2021 challenge official phase. Their performances in terms of the challenge score on the validation set against the model sizes in terms of the number of trainable parameters is shown in Figure 6. We can see in this figure that the variants of [4] are more effective compared against others while having moderate sizes, especially the ones with bottleneck building block. It should be noted that authors of TResNet [24] replaces all basic building blocks with bottleneck building block in their TResNet V2 in their GitHub repository. This

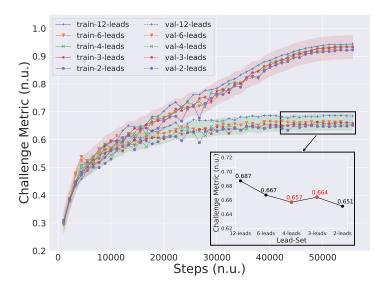


Figure 5: Curves of challenge scores on both the training and validation set from the baseline model tested on the 5 lead-sets.

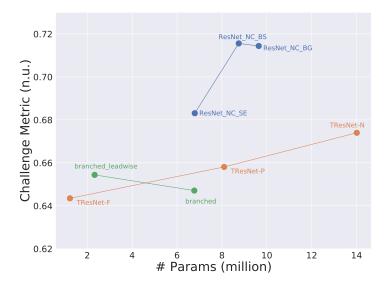


Figure 6: Plot of the performances of the CNN backbones against their sizes in number of trainable parameters.

might be another evidence of the superiority of the bottleneck building block over the basic building block.

7. Ablation Study and Beyond

7.1. Effect of Loss Function, Normalization, Augmentations, and More

As is already mentioned in Section 4.1, we find that the asymmetric loss largely improves the performance of neural network models. We are going to show this point in this section. We also conduct various experiments on other techniques including band-pass filtering, normalization, augmentations, learning rate schedulers, etc., aiming at figuring out whether they truly help augmenting model performances or not. This servers as a part of our ablation studies in a broader sense, which we call Ablation Study 0.

We list the choices for the components for training neural networks that are altered in this part of ablation study:

- With or without band-pass filtering;
- Loss functions: asymmetric loss, weighted BCE loss;
- Normalization: z-score, no normalization;
- Augmentations: mixup, label smoothing;
- Learning rate scheduler: OneCycle scheduler, self-adaptive learning rate by adamw_amsgrad with initial value 0.001;

The baseline model for conducting this part of ablation study is the CRNN model used in Section 6.1.

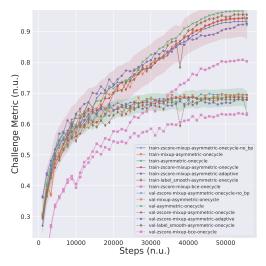
Figure 7a demonstrates the growth of the challenge score on both the training and the validation set. Mini-batch loss against the number of steps is plotted in Figure 7b. One can draw the conclusion that the asymmetric loss contributes most to the improvement of the baseline model. The OneCycle learning rate scheduler takes the second place but the augment is much less significant. Influences of other techniques are small. The loss curves on the training set with augmentation techniques have larger oscillations, while the loss curve without any augmentation techniques (the green "asymmetric-onecyle" curve with cross markers in Figure 7b) decreases more stably. It is surprising that without the band-pass filtering preprocessing procedure, the training loss curve has the largest oscillation while the trained model performs the best on the validation set.

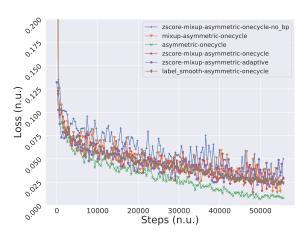
There are other phenomena we want to emphasize but not included in this ablation study. To name a few: performance of neural network models is severely deteriorated if low order band-pass filtered is added into the data preprocessing pipeline, in which case the loss in the pass-band is non-neglectable. Batch size also affects the model performance, in which case hyperparameters of the learning rate scheduler should be carefully tuned to fit the batch size. This problem has been discussed in literature [33] and in less formal papers [34].

7.2. Effect of Neural Network Complexity

Intuitively, increasing the network complexity helps improving model performances. However, it is not always true, sometimes even the opposite. We designed the following sets of experiments to check the contributions of various component of the whole network to the challenge task.

- 1. Ablation Study i: we fix the CNN backbone ResNet-NC-SE, and gradually shrink the other parts:
 - LSTM + SE attention layer + 2 linear layers (denoted lstm-se-2linear);





- (a) Curves of challenge scores on both the training and validation sets.
- (b) Curves of (mean value of) Mini-batch asymmetric loss on the training set.

Figure 7: Ablation Study 0. The techniques used are concatenated by "-" in the legend of the figures. The suffix "no_bp" indicates that the input ECGs are not band-pass filtered before fed into the model. The experiment using the weighted BCE loss is not included in (b), since it's meaningless to compare it with the rest 4. The curves in (b) are smoothed using exponential moving average (EMA) with weight 0.6.

- SE attention layer + 2 linear layers (denoted se-2linear);
- 2 linear layers (denoted 2linear);
- 1 linear layer (denoted 1linear).
- 2. Ablation Study ii: we change the CNN backbones, and fix the rest part of the network to be LSTM + SE attention layer + 2 linear layers. The CNN backbones includes the ResNet variants introduced in Section 3.4, as well as the multi-branched CNN introduce in Section 3.2 and its lead-wise variant introduce in Section 3.3.
- 3. Ablation Study iii: this ablation study has similar setting as in Ablation 2, the difference is that the rest part of the network other than the CNN backbone is changed to the simplest single linear layer.

One can observe in Figure 8 that the 4 combinations listed in Ablation Study i have very close performances on the validation set. However, the extra LSTM layer largely suppresses the overfitting on the training set. The other 2 combinations (se-2linear and 2linear) with extra components other than one bare linear layer have lowered performances. Increasing the network complexity does not always result in better performances.

With the help of extra modules (lstm-se-2linear), neural networks with different CNN backbones also have very close performances as illustrated in Figure 9. The gap between the challenge scores on the training set and on the validation set is significantly smaller when using the ResNet-NC-BS backbone. The bottleneck building block seems

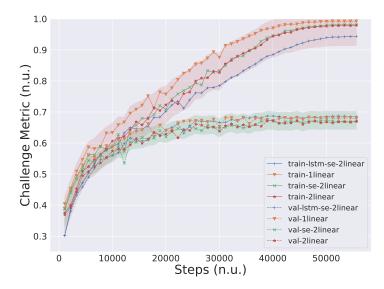


Figure 8: Ablation Study i: Curves of challenge scores on both the training and validation sets using 12 leads ECGs. Names of neural network components other than the ResNet-NC-SE backbone are concatenated by "-" in the legend.

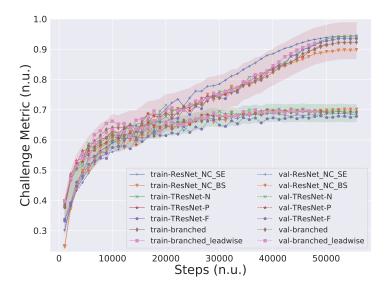


Figure 9: Ablation Study ii: Curves of challenge scores on both the training and validation sets using 12 leads ECGs from neural networks composed of different CNN backbones followed by an LSTM module, an SE module, and two linear layers.

to have the effect of regularization to some extent.

When CNN backbones are concatenated with only a bare linear layer for prediction, performances of the networks diverge, as can be observed in Figure 10. Networks completely without any extra structures (inner SE, bottleneck, etc.) degrade dramatically. Other networks that have higher complexities overfit very fast on the training set.

To fully validate the findings in Ablation Study i, we conduct an additional ablation

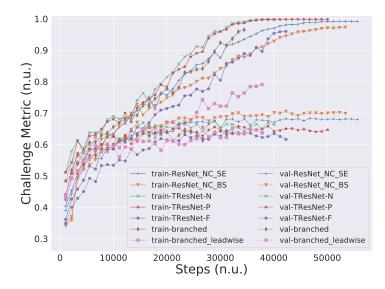


Figure 10: Ablation Study iii: Curves of challenge scores on both the training and validation sets using 12 leads ECGs from neural networks composed of different CNN backbones followed by only one linear layers. It should be noted that the training setups for this set of experiments have slight difference against experiments in Figure 6, hence the performance of ResNet-NC-BS slightly drops. But its superiority against others is unchanged.

study using the CNN backbone ResNet-NC-BS and with full set of all combinations (totally 8) of modules listed in Ablation Study i. The results are gathered in Figure 11. The simplest combination, i.e. a backbone ResNet-NC-BS with a linear output layer performs the best. Our claim that increasing the network complexity does not always result in better performances is verified again.

The overall statistics of all the ablation studies are gathered in Table 5.

8. Discussion and Conclusions

We can conclude from the statistics of the experiments and from the CinC2021 challenge results that the neural network architectures we designed in Section 3 provides effective solutions to the problem of detecting a wide range of ECG abnormalities. These neural network models are robust enough such that their performance deterioration on reduced-lead ECGs are almost neglectable. This gives positive answers to the critical problems raised by the CinC2021 challenge. Furthermore, the "lead-wise" mechanism offers the potential to reduce model size, as well as to reuse model parameters.

Another contribution of this work is that it exhibits the impact on the performance of neural networks from various aspects, for example from the data augmentation techniques through extensive ablation studies. It validates some principles of network design, choices of loss function, etc. This lays solid foundation for further research on related topics in the future.

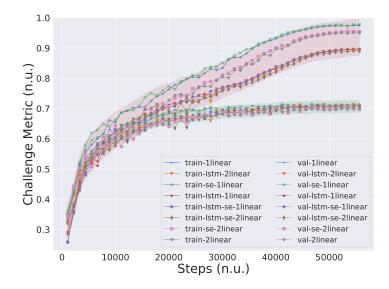


Figure 11: Curves of challenge scores on both the training and validation sets using 12 leads ECGs for a set of CRNN models. The CNN backbone is ResNet-NC-BS, the rest components of the models can be read from the legend of this figure.

However, our work still has its limitations and would be further improved in our future work. First of all, the neural architecture search is not so extensive since many other network architectures, especially those can be inherited from computer vision, have not been experimented. Other powerful modules from other application fields of deep learning are not studied as well, for example the transformer encoders [35]. Second, the mechanism of parameter reuse for "lead-wise" CNNs have not been fully established and implemented. This requires extracting channel-specific weights from the larger weight matrices in the neural networks trained on the standard 12-lead ECGs. An easier alternative method is to fill zeros to the input ECGs in the left-out leads.

The problem of label heterogeneity and insufficiency across the 5 datasets listed in Section 2.1 should also be noted. We observed labels that violates clinical criteria. For example, some "LAD" records violates the "3-lead" method which is more exact than the "2-lead" method mentioned in Section 5. An example is illustrated in Figure A1. A more common situation is that a dataset lacks labels of specific types of ECG abnormalities while some records inside this dataset should be diagnosed with such abnormalities. For example, the dataset CPSC [10] have labels related neither to abnormalities of electrical axis (namely "LAD" and "RAD") nor to abnormalities of amplitudes (e.g. "LQRSV", "PRWP" (poor R wave progression)). However some of its records should be diagnosed with one or more of such abnormalities, as illustrated in Figure A2 and in Figure A3. Directly training with such data would definitely harm the effectiveness of the models. The question of how to train models in a more proper way under such complicated circumstance is to be studied in our future work.

CNN backbone	LSTM	CE.	// 1:	# params	Best	Training	Inference
	LSIM	SE	# linear		Score	Speed(sig/s)	Speed(sig/s)
ResNet-NC-SE	×	×	1	6.79 M	0.683	399	2917
	×	×	2	7.14 M	0.673	391	2889
	×	√	2	7.17 M	0.674	386	2954
	√	√	2	8.92 M	0.687	352	2870
	×	×	1	8.75 M	0.716	344	2007
	×	×	2	10.1 M	0.700	338	2026
	×	√	1	9.16 M	0.712	334	2024
ResNet-NC-BS	×	√	2	10.5 M	0.702	326	2010
resner-nc-ps	√	×	1	11.9 M	0.710	313	1966
	√	×	2	12.3 M	0.713	310	1940
	√	√	1	11.9 M	0.708	302	1955
	√	√	2	12.3 M	0.708	302	1949
TResNet-N	×	×	1	14.0 M	0.674	244	1395
IResnet-N	√	√	2	18.4 M	0.703	182	1139
TD o gNo+ D	×	×	1	8.10 M	0.658	378	2046
TResNet-P	√	√	2	12.5 M	0.700	252	1553
TResNet-F	×	×	1	1.23 M	0.643	373	3325
	√	√	2	4.42 M	0.679	262	1785
Branched	×	×	1	6.77 M	0.647	408	515
	√	√	2	10.2 M	0.698	146	362
Branched-LW	×	×	1	2.33 M	0.654	357	536
	√	√	2	7.46 M	0.699	134	360

Table 5: Overall statistics of Ablation Study i, ii, iii and the additional ablation study. Best scores are computed on the validation set. Training speed is computed under the settings stated in Section 4. Inference speed is computed using one RTX3080 based on a single run on the training set.

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Code Availability

Code, configurations and auxiliary data are all available at https://github.com/DeepPSP/cinc2021

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Figure A1: Plot of the first 6 seconds of leads I, II, aVF of the record "HR05631" from the PTB-XL subset. It has "LAD" in its scored label list. Leads I and II are positive, lead aVF is negative. It will be classified as "LAD" by the "2-lead" method detector proposed in Section 5, while classified as non-"LAD" by the "3-lead" method detector.



Figure A2: Plot of the first 6 seconds of leads I, II, aVF of the record "A0226" from the CPSC subset. It has label "CRBBB". Lead I is positive, leads II and aVF are negative, hence this record should be classified as "LAD" by the "3-lead" method detector proposed in Section 5. If using the "2-lead" method, this record would definitely be classified as 'LAD". There are many more such examples in the CPSC subset, which contribute many false positives in Table 4.

Appendix A. Examples of Label Heterogeneity and Insufficiency

In this section we present several examples of the issue of label heterogeneity and insufficiency across the subsets of the CinC2021 database mentioned in Section 8.

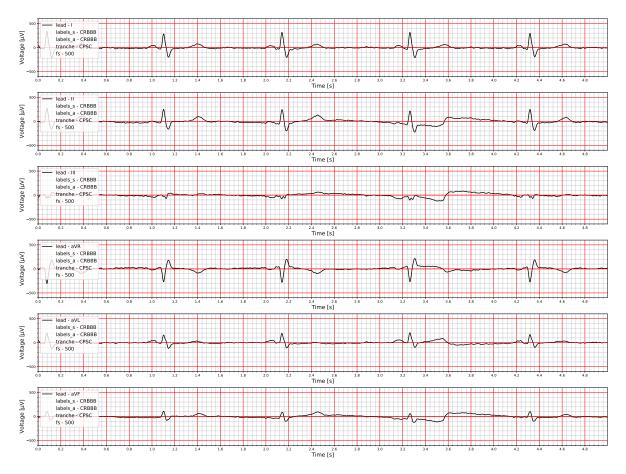


Figure A3: Plot of the first 6 seconds of the limb leads (I, II, III, aVR, aVL, aVF) of the record "A0095" from the CPSC subset. It has label "CRBBB". All its values are < 5mm (0.5mV) in the limb leads, hence should be classified as "LQRSV". Moreover, it should be classified as "LAD" by the "2-lead" method detector proposed in Section 5, although which is a mis-classification.

Appendix B. Probability Matrix of a Typical Neural Network Model

We plot the matrix of predicted probabilities on the validation set for the best neural network model ResNet-NC-BS in Figure B1. It should be noted that the data processing pipeline for this model includes the Z-score normalization. We can see from this figure that the model performs particularly bad on 2 classes, namely "LQRSV" (low QRS voltage) and "PRWP" (poor R wave progression). The former is described in Section 5, clinical diagnostic criteria for the latter is "absence of the normal increase in size of the R wave in the precordial leads when advancing from lead V1 to V6". These 2 classes are exactly the only 2 among the 26 scored classes in the CinC2021 challenge database that are directly related to the amplitudes of the ECGs. The normalization operation destroys such characteristics. With the help of clinical rule based detectors, such performance deterioration of the neural network models could be alleviated. (ref. the "TP" column and "FN" column of the Table 4.) Clinical rule based detectors are

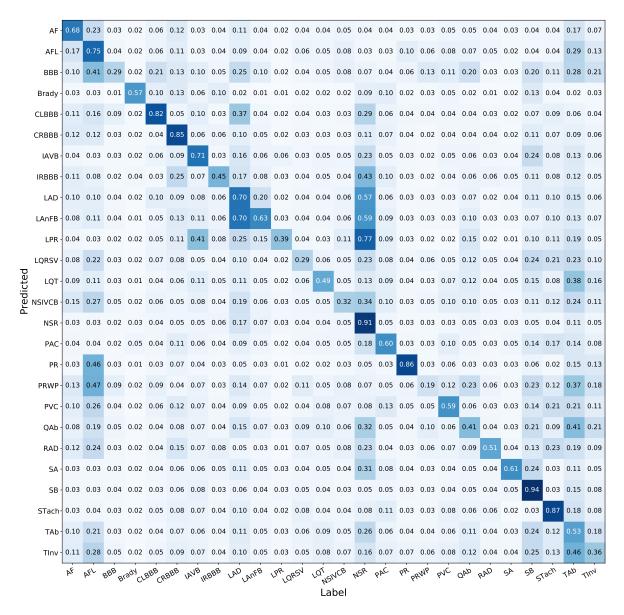


Figure B1: The matrix of predicted probabilities on the validation set for the best neural network model ResNet-NC-BS

truly helpful assists to neural network models.