

# ME-GAN: Learning Panoptic Electrocardio Representations for Multi-view ECG Synthesis Conditioned on Heart Diseases

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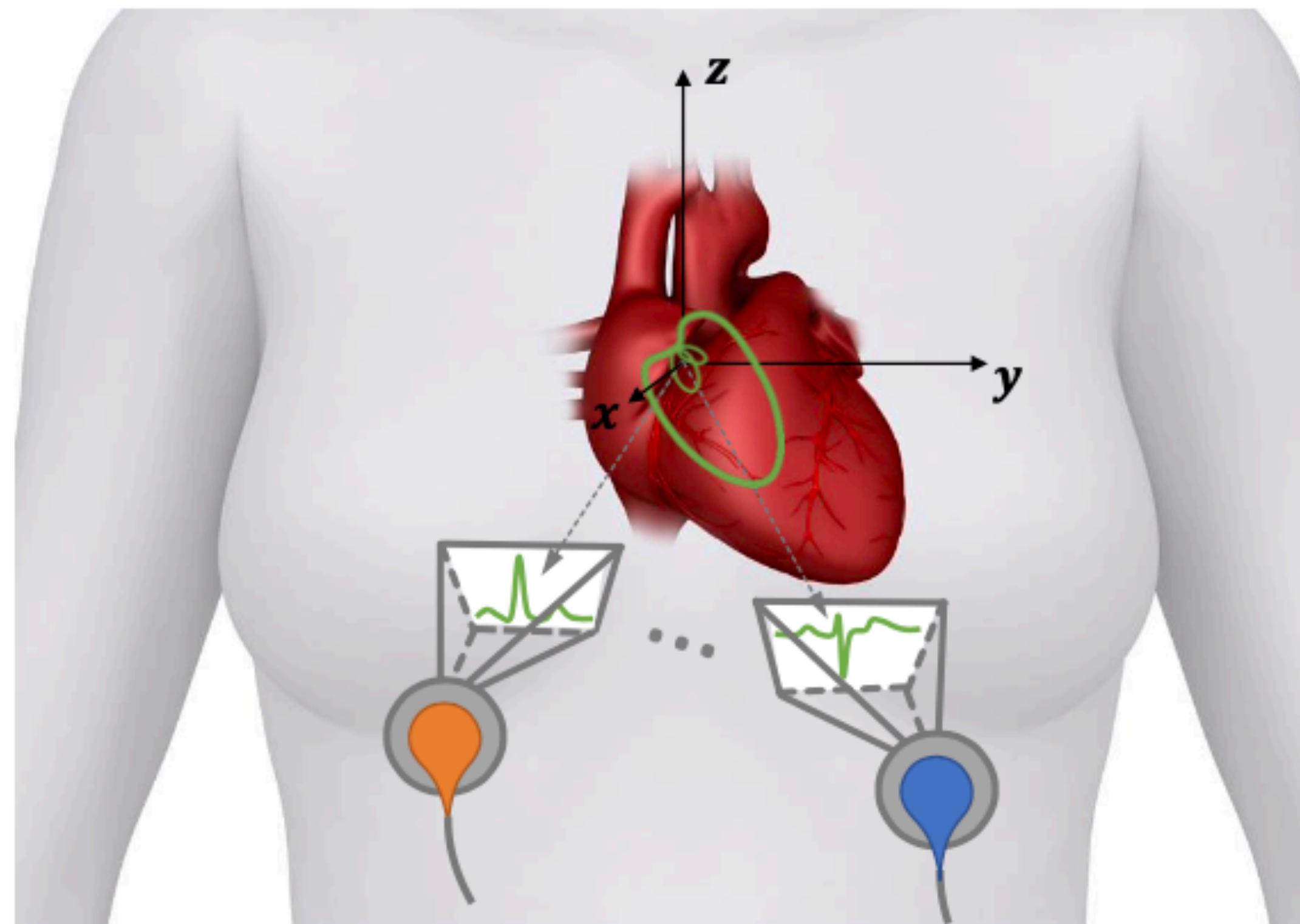
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# Background

- ECG is a widely used noninvasive diagnostic tool for heart diseases.
- Synthesizing ECG data is thus quite helpful to **increasing training samples, privacy protection, and annotation burden reduction**

# Issues

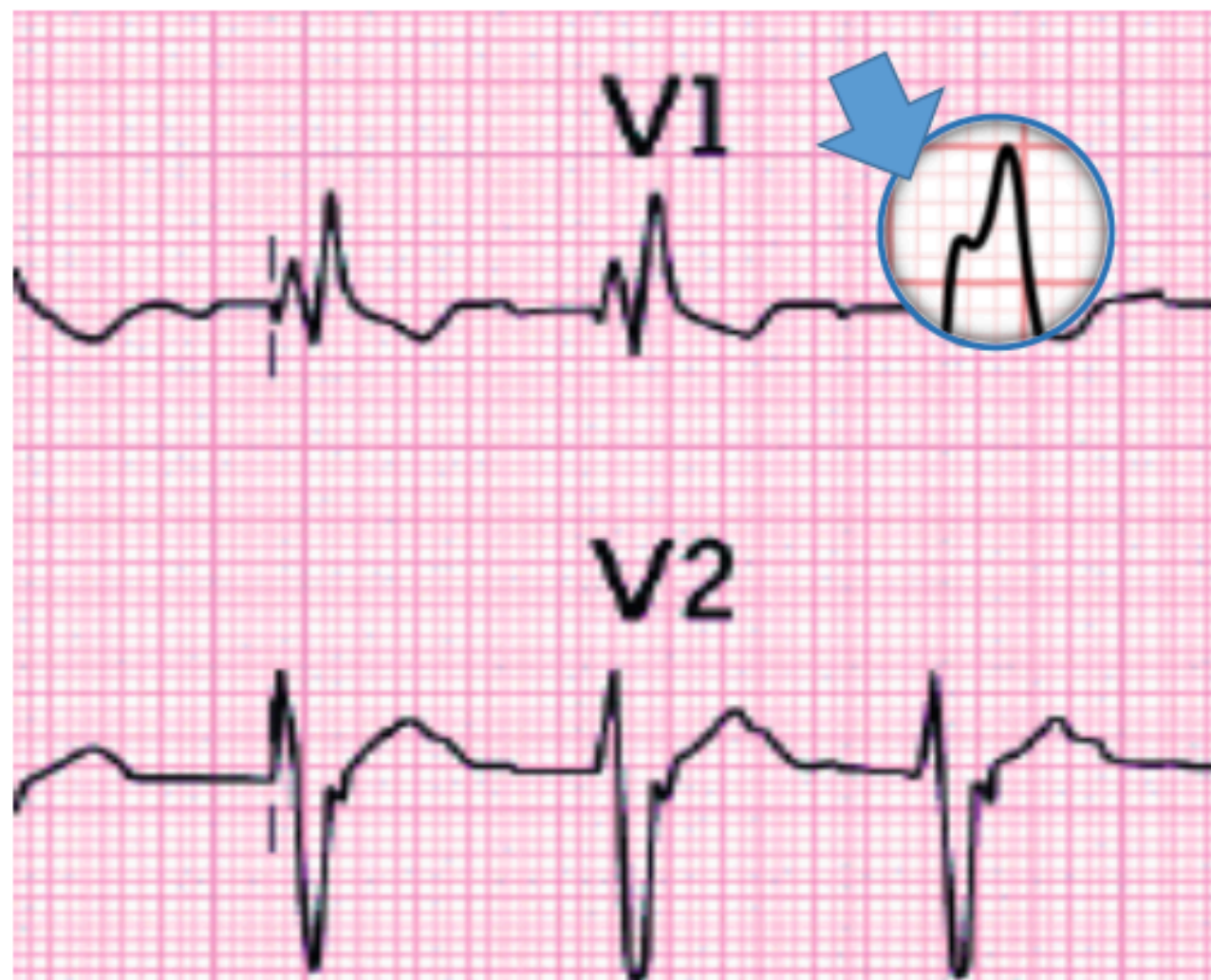
- How to ensure the multi-view ECG synthesis is trustworthy?
  - Different views were highly correlated because they represent the same heart beat signals from different viewpoints



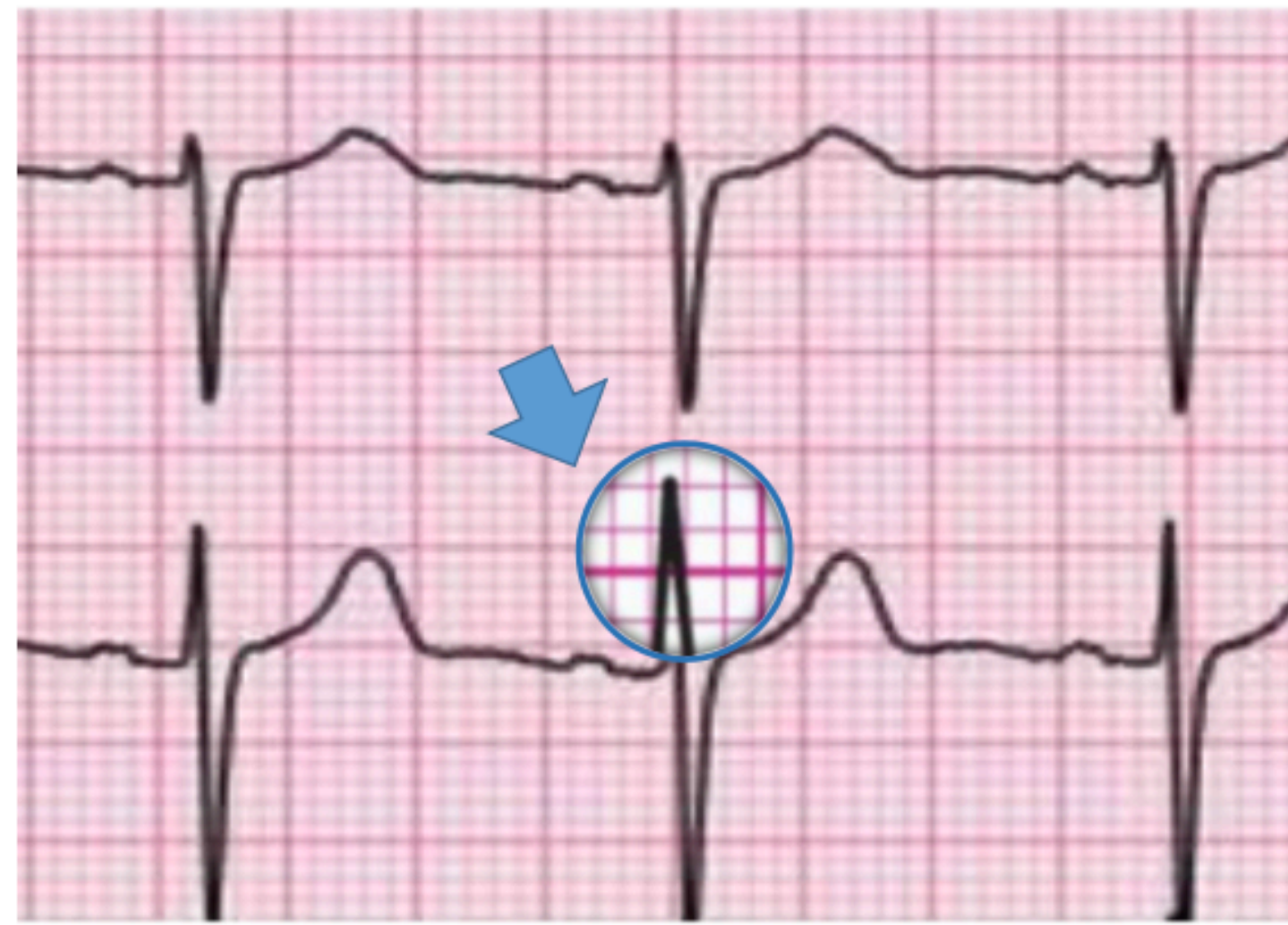


# Issues

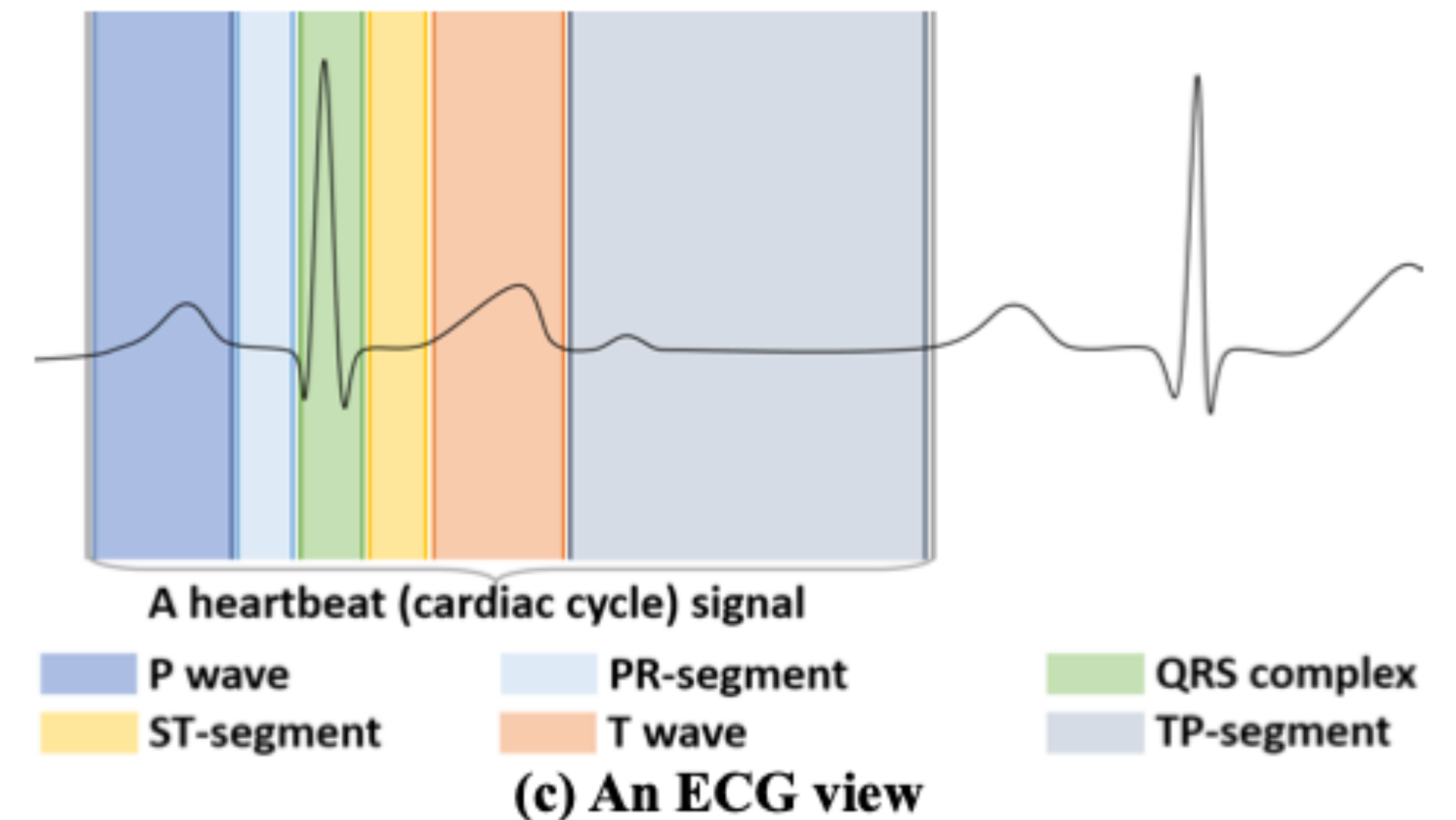
- How to synthesize ECG conditioned on heart diseases?
  - ECG manifestations of some heart diseases are often localized in specific waveforms and were shown as some fine-grained waveform deformations.



Abnormal waveform with LBBB.

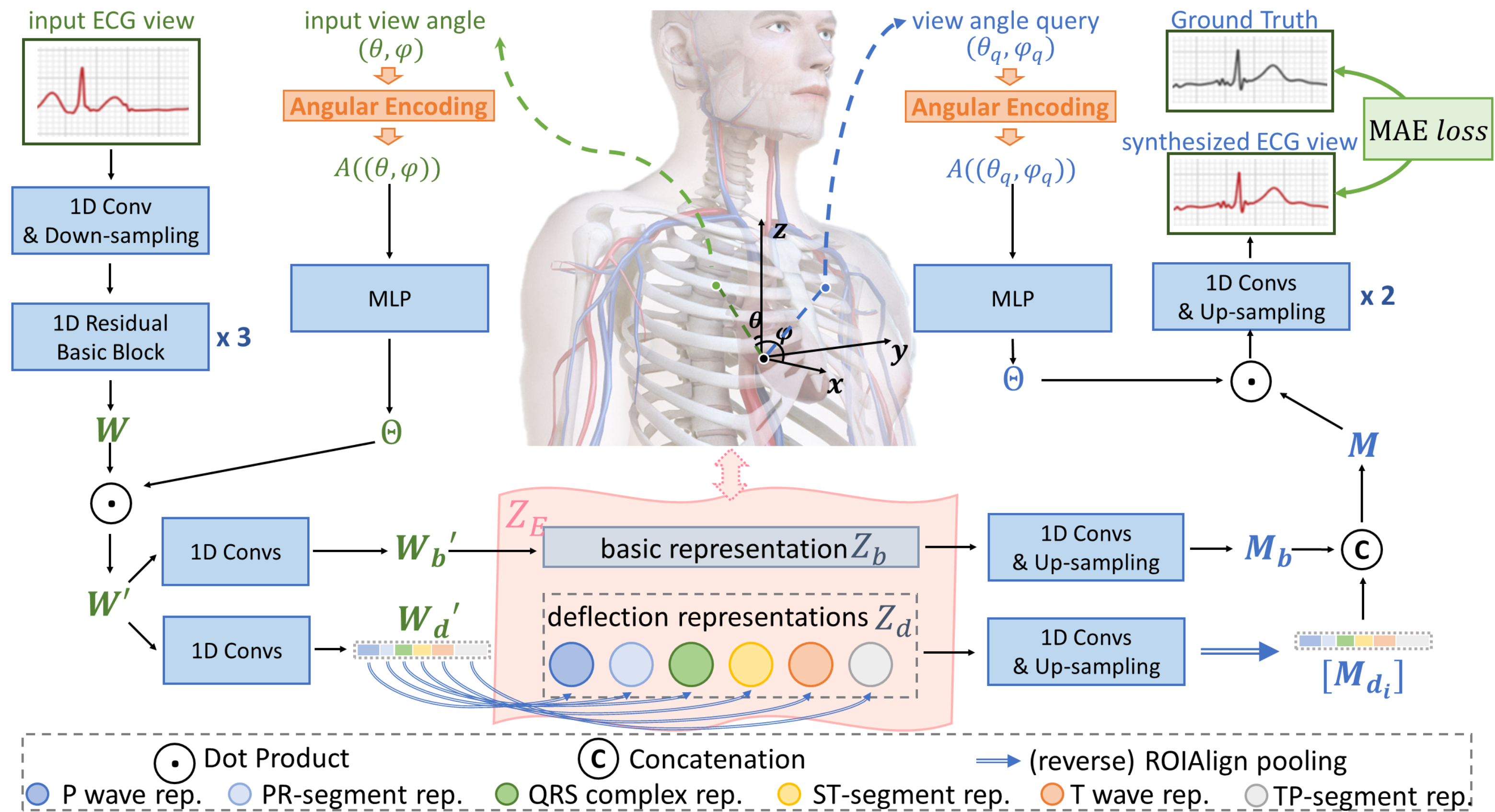


Normal waveform at R peak.





# Related Work

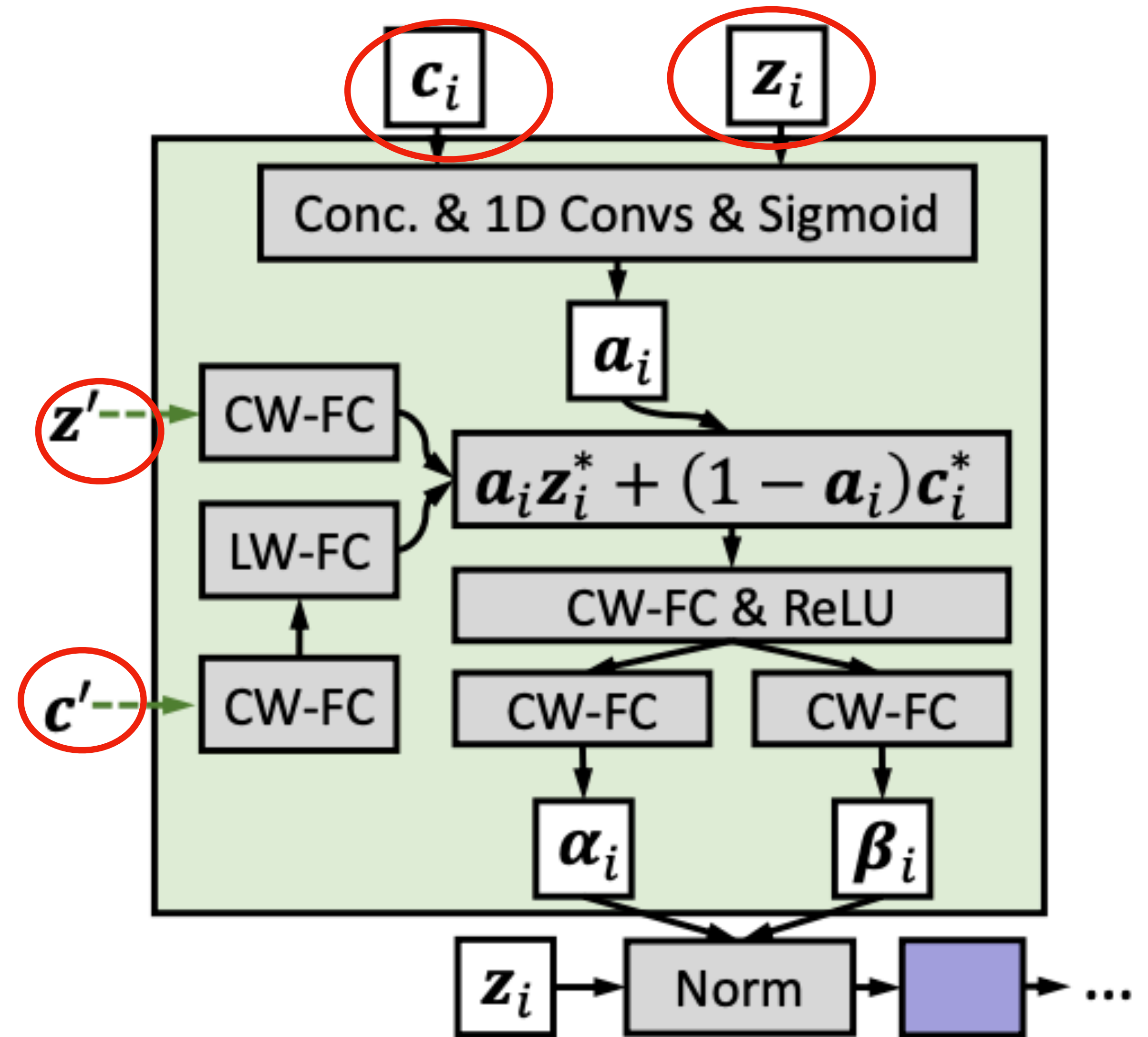


Chen, J., Zheng, X., et al. Electrocardio panorama: Synthesizing new ECG views with self-supervision. In *IJCAI*, 2021.

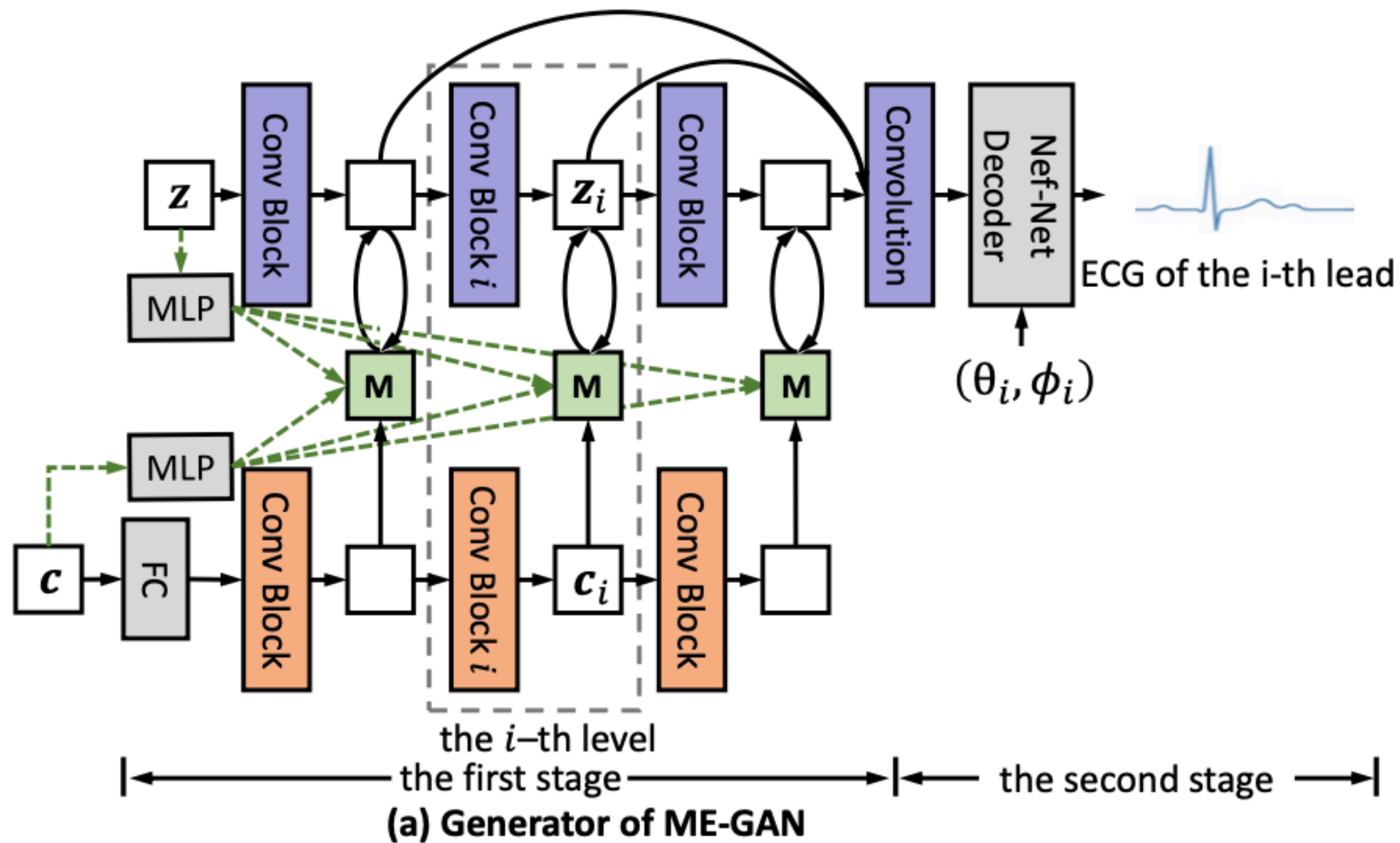
# Method

- **Mixup Normalization Layer**
  - Synthesize fine-grained waveformed conditioned on heart diseases

$$\tilde{z} = \alpha \odot \frac{z_i - \mu}{\sigma} + \beta_i$$

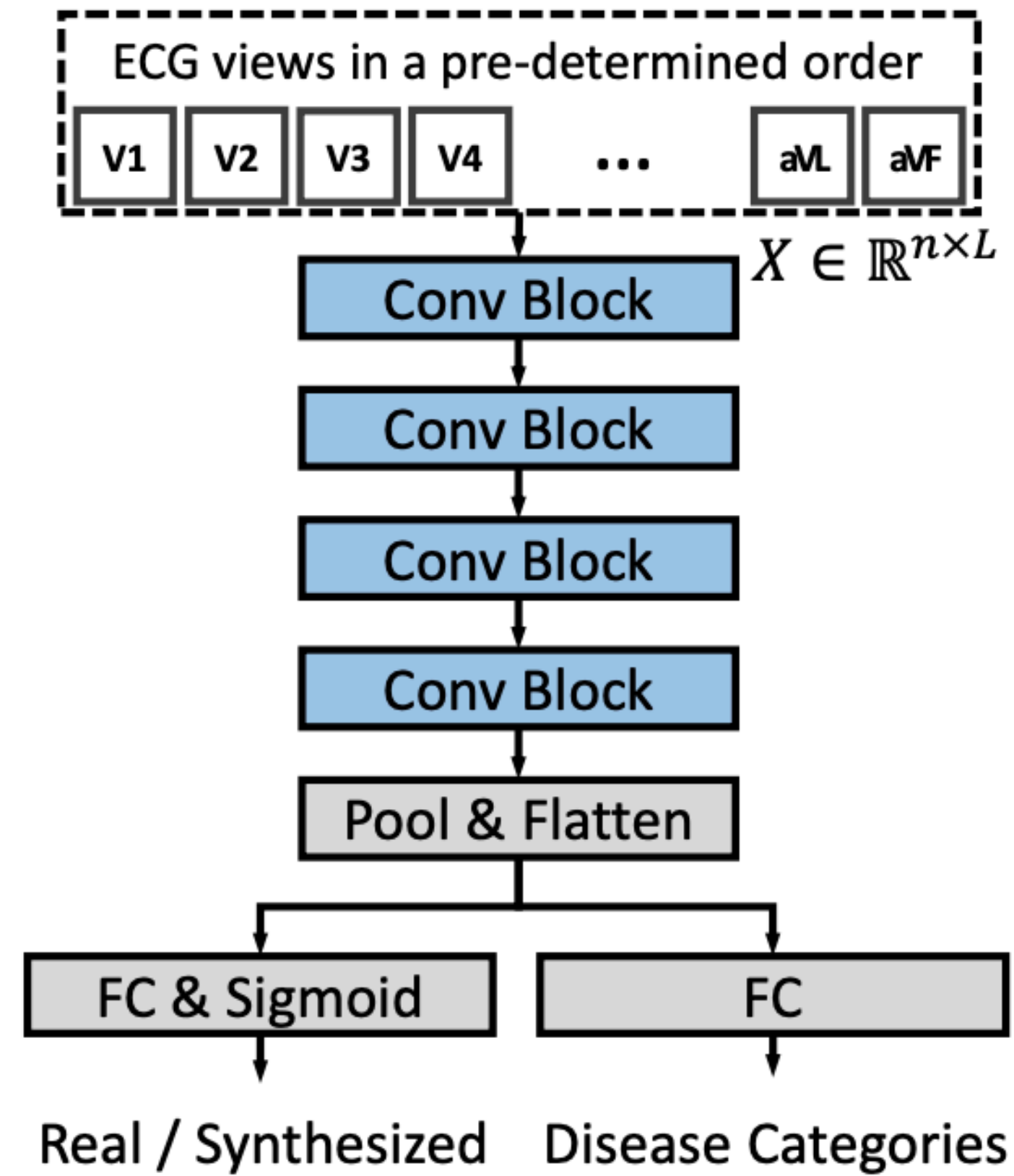


# Generator

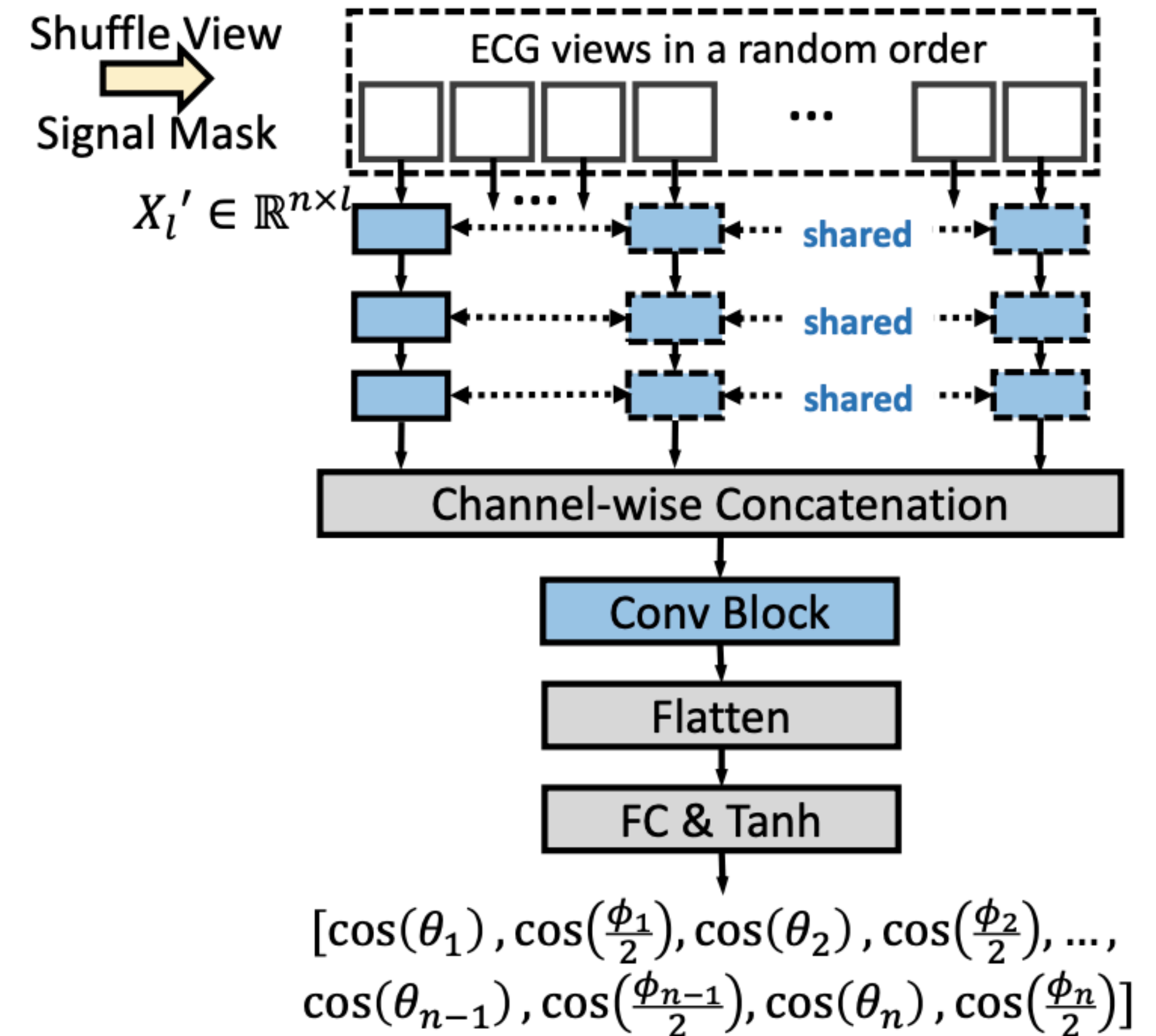




# Discriminator



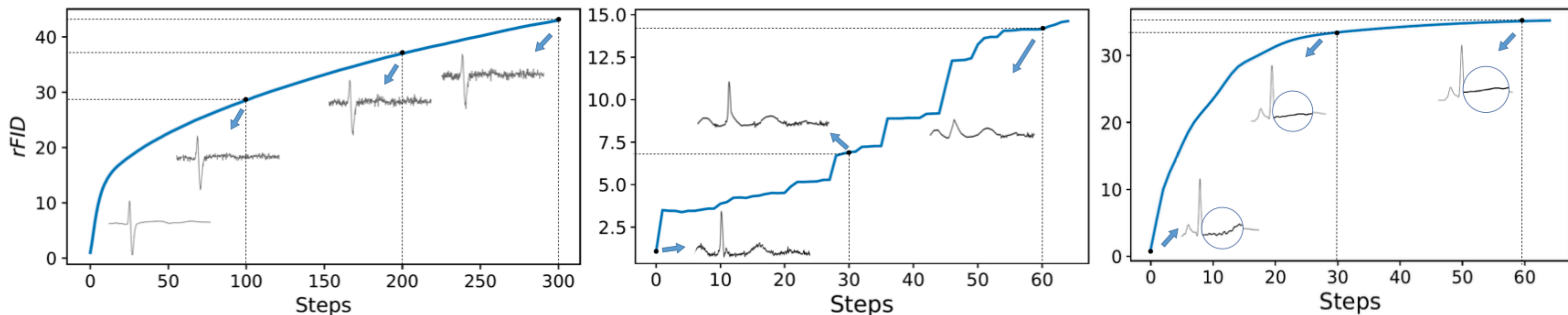
(a) Major Discriminator



(b) View Discriminator



# rFID



(a)  $rFID$  changes with continuously added noises. (b)  $rFID$  changes with ECG signals replaced by line segments. (c)  $rFID$  changes with ECG signal blurring.

**Figure 5. Illustrating  $rFID$  changes during a process in which noise is continually added onto ECG signals.** The “V2” view of an ECG signal is presented for visually showing the signal quality.

# Result

**Table 1. Synthesis performances of various GAN models.** The lower *rFID* score is the better, and the accuracy score of 1-NN classifier (1NNC) is better if it is closer to 0.5. The best performances are marked in **bold**.

Methods	Tianchi w/ diseases								Tianchi w/o diseases		PTB w/o diseases	
	overall		<i>LAD</i>		<i>RBBB</i>		<i>RAD</i>		overall		overall	
	1NNC	<i>rFID</i>	1NNC	<i>rFID</i>	1NNC	<i>rFID</i>	1NNC	<i>rFID</i>	1NNC	<i>rFID</i>	1NNC	<i>rFID</i>
WGAN-GP	0.722	7.534	0.697	5.359	0.683	3.509	0.676	6.892	0.640	5.494	0.656	42.046
ACGAN	0.699	15.944	0.668	7.464	0.672	10.942	0.684	13.144	–	–	–	–
LSGAN	0.870	13.427	0.795	6.151	0.676	8.984	0.746	11.367	0.775	17.341	0.673	43.481
CGAN	0.757	12.479	0.585	8.495	<b>0.607</b>	5.489	<b>0.594</b>	10.849	–	–	–	–
SMDCGAN	0.998	38.619	0.934	23.985	0.823	18.370	0.924	31.408	0.617	2.618	0.623	20.182
BC-GAN	0.832	12.701	0.723	8.299	0.617	5.975	0.634	10.685	0.983	42.545	0.997	158.698
CBL-GAN	0.826	6.990	0.673	5.322	0.739	3.471	0.694	5.573	0.611	6.173	0.713	65.750
ME-GAN (Ours)	<b>0.643</b>	<b>3.722</b>	<b>0.567</b>	<b>2.662</b>	0.663	<b>1.491</b>	0.712	<b>3.433</b>	<b>0.582</b>	<b>2.343</b>	<b>0.618</b>	<b>15.282</b>

# Result

*Table 2. Classification performances of 1D ResNet-34 trained on augmented training sets.*

Method	Diseases (PR-AUC)			
	<i>LAD</i>	<i>RBBB</i>	<i>RAD</i>	Mean
1D ResNet-34 (baseline)	0.925	0.801	0.911	0.879
WGAN-GP	0.916	0.826	<b>0.922</b>	0.889
ACGAN	0.912	0.817	0.913	0.881
LSGAN	0.918	0.858	0.910	0.895
CGAN	0.906	0.867	0.904	0.892
SMDCGAN	0.913	0.832	0.921	0.889
BC-GAN	0.896	0.820	0.898	0.871
CBL-GAN	0.910	0.857	0.896	0.888
ME-GAN (Ours)	<b>0.927</b>	<b>0.870</b>	0.908	<b>0.902</b>



**Thank you for listening**