SRECG: ECG Signal Super-Resolution Framework for Portable/Wearable Devices in Cardiac Arrhythmias Classification

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Abstract—A combination of cloud-based deep learning (DL) algorithms with portable/wearable (P/W) devices has been developed as a smart heath care system to support automatic cardiac arrhythmias (CAs) classification using electrocardiography (ECG). However, long-term and continuous ECG monitoring is challenging because of limitations of batteries and transmission bandwidth of P/W devices while incorporated with consumer electronics (CE). A feasible approach to address this challenge is to decrease sampling rates. However, low sampling rates lead to low-resolution signals that hinder the CAs classification performance. In this study, we propose a DL-based ECG signal super-resolution framework (called SRECG) to enhance low-resolution ECG signals by jointly considering the accuracies when applied to the DL-based high-resolution multiclass classifier (HMC) of CAs. In our experiments, we downsampled the ECG signals from the CPSC2018 dataset and evaluated their HMC accuracies with and without the SRECG. Experimental results show that SRECG can well improve the HMC accuracies as compared to traditional interpolation methods. Moreover, approximately half of the CAs classification accuracies of HMC were maintained within the enhanced ECG signals by SRECG. The promising results confirm that SRECG can be suitably used to enhance low-resolution ECG signals from P/W devices with CE to improve their cloud-based HMC performances.

Index Terms—Cardiac arrhythmias, consumer electronics, deep learning, electrocardiography, signal enhancement.

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I. INTRODUCTION

C ONSUMER technologies have changed the world, especially in healthcare, where traditional medicine has been pushed to transform into telemedicine, connected-health, ehealth, mobile-health, and smart health [1]. Through the growth of portable/wearable (P/W) devices, smart healthcare systems enable medical doctors to not only collect patient data without worrying about the physical distance in between, but also to implement instant treatment remotely [2]. Moreover, P/W devices make it possible to continuously monitor the health conditions of a patient and collect relevant data. This data is helpful in alerting medical professionals to critical conditions and implement medical actions and is valuable for providing new medical insights for further research.

Cardiac arrhythmias (CAs) are harbingers of cardiovascular diseases that are life-threatening to human beings [3]. A typical assessment tool relies mainly on electrocardiography (ECG) signals that record the electrical activity of the heart. ECG is a noninvasive and inexpensive method that is clinically applied to monitor heart functionality [4], [5], [6]. The diagnosis of CAs is based on wave-like features, such as the P wave, QRS complex, and T wave, of ECG signals. Medical doctors can manually examine the ECG features to evaluate the occurrence of CAs. The entire examination process for a long-term ECG record is time consuming and tedious. Therefore, it is essential to develop an automatic CAs classification system to support clinical diagnosis.

Several studies have developed cloud-based CAs monitoring systems that combine cloud services and advanced deep learning (DL) models for smart healthcare systems. These CAs monitoring systems collect ECG using P/W devices and convey heavy processing to cloud DL engines for further CAs classification [7], [8]. For a P/W device incorporated with consumer electronics (CE), long-term monitoring of CAs requires continuous sampling and interpretation of user data, which may lead to unacceptable power consumption. Factors such as the monitoring source, sampling frequency, electronic components, and transmission protocols affect the power efficiency of a P/W device. Reducing the monitoring sources and sampling frequency are two compromises for reducing power consumption. The latter has been extensively studied because at high sampling frequencies, up to 90% of power consumption of health monitoring systems is due to data sampling [9], [10], [11].

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Various power efficiency strategies have been applied to automatic CAs classification systems to enable long-term health monitoring services [10], [11], [12]. Maji et al. proposed an adaptive power management approach to classify critical and non-critical signals before further DL-based ECG classification [13]. Hierarchical classification models can efficiently reduce power consumption [13]. Furthermore, they employed typical data augmentation approaches, including SMOTE and BIRCH [14], [15], to improve performance in terms of power consumption [13]. Biagetti et al. proposed a low-cost wireless system to transmit the acquired surface electromyographic and ECG signals through a 2.4 GHz radio link based on the IEEE 802.15.4 physical layer [16]. This system can be combined with different mobile and wearable devices to support low-cost and low-power health-monitoring systems [16]. Raj proposed a tuned twin support vector machine (SVM) classifier using particle swarm optimization to classify 16 categories of ECG signals on benchmarked Physionet data, which was four times faster than the standard SVM and showed enhanced performance when implemented on the Internet of Things platform [17].

Despite these strategies, the reduction in monitoring sources is still a typical and direct approach to saving energy, for example, reducing the recording of ECG sources from a full 12-lead ECG signal to a single lead [18], [19]. In our previous studies, we provided an empirical discussion over a 12-lead source to select the most suitable single-lead ECG source for DL-based CAs classification [20]. Another approach is to decrease the sampling rate of the P/W device, which is an efficient approach for reducing power consumption. However, in general, a lower sampling frequency is accompanied by a reduction in information, which is, low resolution. Low-resolution signals can cause degradation of CAs classification. For example, several important ECG characteristics are degraded in low-frequency signals, which may decrease the ability of the monitoring system to detect CAs automatically. Although there are studies regarding the sampling frequency of ECG, most are related to the association of heart rate variability, which is widely used as a noninvasive marker of the autonomic nervous system [21], [22].

A few existing studies have aimed to develop lowresolution automatic CAs classification. Sidek et al. proposed interpolation-based signal enhancement approaches, involving piecewise cubic Hermite interpolation and piecewise cubic spline interpolation, to increase the quality of signals and classification accuracy at a lower sampling frequency [23]. The CAs detection accuracy using the interpolation-based enhancement approach could reach ~99%, whereas without signal enhancement, it only reaches $\sim 97\%$ [23]. Mathews et al. proposed a DL-based CAs classification framework using a Restricted Boltzmann Machine and deep belief networks to improve the classification abilities at lower sampling rates, where a range of sampling frequencies from 72 Hz to 360 Hz was explored [24]. The experimental results validated the DL approach for low-resolution CAs classification [24]. However, these previous studies did not investigate automatic CAs classification at very low sampling rates, while several studies have developed P/W-based monitoring systems with an extremely low sampling frequency to support other long-term healthcare services, such as daily activity recognition and fall detection [10], [25].

To alleviate the loss of information at very low sampling frequencies, image super-resolution (SR) may serve as a viable solution to enhance low-resolution ECG signals for automatic CAs classification. Image SR has recently become a popular and well-studied area of interest [26], [27], [28]. The idea is to reconstruct the target information from the low-resolution data. The primary goal of image SR is to recover data from compressed or low-quality signals to facilitate better signal resolution for subsequent tasks. In particular, image SR has long been an important subject of image processing techniques in computer vision that aim to recover high-resolution images from low-resolution images, which in turn is a challenging task owing to its mathematically ill-posed nature [29], [30], [31], [32]. The rapid development of DL techniques in recent years has also driven researchers to tackle image SR tasks. In general, DL-based SR algorithms have three different components: (1) network architectures, such as convolutional layers, residual connections, recursive layers, and up-sampling layers [33], [34], [35]; (2) design of loss functions, such as pixel, perceptual, or adversarial losses [31], [36], [37]; and (3) learning principles and strategies [38], [39], [40], [41]. Some SR algorithms have reached the current state-of-the-art performance [42], [43], proving useful for improving other vision tasks [29], [30], [31], [32]. Task-driven SR can significantly enhance the accuracy of an object detector on low-resolution images, thus creating a positive impact on vision recognition [30]. Similarly, in signal processing, audio SR refers to the task of increasing the sampling rate for a given low-resolution (i.e., low-sampling rate) audio. Motivated by recent advances in learning-based algorithms for speech recognition [44], [45], [46], music generation [47], [48], and other areas [49], significant progress has been made in audio SR via the introduction of DL. The key advantage here is that one is allowed to directly model raw signals in the time domain [47], [50], [51], [52], and effectively capture long-term dependencies [53].

Although SR has had a leaping success in human vision tasks, to the best of our knowledge, no prior studies have attempted to apply the SR technique to ECG signals for automatic CAs classification. In this paper, we proposed and investigated a novel DL-based ECG signal SR framework (SRECG) as a processor. This framework is expected to help automatic CAs classification tackle the challenges of lowresolution ECG for P/W devices and achieve better accuracy. The proposed SRECG enhances low-resolution ECG signals with composite training loss in the cloud and, in turn, yields better utilization for the DL-based high-resolution multiclass classifier (HMC) of CAs. As shown in Fig. 1, compared to the typical workflow of a smart healthcare system, when employing SRECG as a computing processor, the ECG signals can be sampled at extremely low frequencies to reduce both energy consumption and data size for data transmission. The processor and HMC are both in the cloud; therefore, their computing power consumption is separated from that of a P/W device. Furthermore, we comprehensively analyzed and explored the



Fig. 1. Comparison of DL Workflows of Smart Healthcare System for a P/W Device. (a) Typical workflow. (b) The proposed workflow based on the proposed SRECG.

impact of sampling frequencies on the DL-based CAs classification of 12 different lead sources of the ECG signals. It would be beneficial to develop reliable DL-based CAs classification models for smart healthcare systems using P/W devices.

The main contribution of this work is threefold: (1) it is the first work that investigates the ECG signals in the CAs classifier among different leads and sampling frequencies, which clearly shows the information loss with resolution reduction of ECG signals sampled from 500 to 1 Hz; (2) our SRECG framework enhances the CAs classification accuracies of HMC when using low-resolution ECG signals from a P/W device at low sampling frequency to resolve its energy consumption and transmission issue with CE; (3) training SRECG with composite loss can recover CAs classification accuracies of HMC better than using only single loss. We believe the proposed SRECG framework could be readily adapted for commercial applications in ECG diagnostics, where data storage is an important consideration.

II. MATERIALS AND METHODS

In this study, we attempted to enhance the ECG signals from 12 different lead sources at low sampling frequencies. Therefore, the open-source training dataset, benchmarked DL model, and categorical metric for performance evaluation are all from CPSC2018, which is a worldwide automatic ECG CAs classification competition [54].

A. CPSC2018 Open-Source Dataset

Detailed information on the CPSC2018 ECG database can be found in [54] by Liu et al. In comparison to the source of the publicly available ECG database with only 2-lead ECG signals of 48 recordings for most previous CAs prediction studies, such as the MIT-BIH Arrhythmia Database [55], [56], it contains 12-lead ECG signals of 6,877 recordings with different durations from seconds to minutes and a label of nine categories, including a normal type and eight abnormal CAs: AF, left bundle branch block (LBBB), right bundle branch block (RBBB), first-degree atrioventricular block (I-AVB), premature atrial contraction (PAC), premature ventricular contraction (PVC), ST-segment elevation (STE), and ST-segment depression (STD). Of the 6,877 recordings, 476 received more than one CA-type label. To simulate the measurements of ECG signals from different lead sources at a low sampling frequency, the CPSC2018 open-source ECG dataset with a single CAs label (6,401 out of 6,877 recordings) was downsampled from 500 Hz to 250, 125, 100, 50, 25, 10, 5, 2, and 1 Hz, and the signals of different lead sources were extracted from 12-lead signals separately.

The down-sampling approach was employed to obtain low-resolution signals of different lead sources from highresolution signals to explore the effects of different resolutions on the CAs classification performance. The lowresolution signals are gathered by taking an integer factor extitn, which refers to the skipping number of sequential units of time to collect the signal. In this study, n =1, 3, 4, 9, 19, 49, 99, 249, and 499 is applied to the CPSC2018 dataset to obtain sampling frequencies: 250, 125, 100, 50, 25, 10, 5, 2, and 1 Hz. To compensate for the unequal length of recordings, we simply applied zero padding before any recording that was shorter than the required length of maximum time of the whole CPSC2018 dataset. There was no preprocessing for the recordings.

B. CPSC2018 Open-Source Benchmarked DL Network

Our previous model architecture, winning CPSC2018, was selected as the reference model to create our HMCs and test the CAs classification performances at different sampling frequencies [20]. It was built on a combined architecture of five CNN blocks, followed by a bidirectional gated recurrent unit (GRU), an attention layer [57], [58], and a fully connected layer (see Fig. 2). In our architecture, each CNN block contained two convolution layers that were followed by a convolution-pooling layer, which is a convolution-based pooling layer, to reduce the number of parameters and computation in the network and to control overfitting [59]. All convolution layers share the same kernel number of 12 and kernel size of three, except for the five convolution-pooling layers with kernel sizes of 24, 24, 24, 24, and 48, sequentially. Furthermore, between these CNN blocks and other independent layers, including the one between the last CNN block and the bidirectional GRU layer, 20% of their connections were randomly dropped. Batch normalization was also used to adjust and rescale the results from the attention layer, which is a special mechanism proposed to generate importanceweighting vectors [60]. LeakyReLU activation functions were used in each layer other than the fully connected layer, where a sigmoid activation function was used [61]. This architecture was built using the Keras package supported by TensorFlow in GPUs [62], [63]. The model has 28,035 trainable parameters to ensure it is compact enough to explore the statistical relations using limited computing resources [20].



Fig. 2. Architecture of CPSC2018 benchmarked DL network (denoted by the function f_{judge}). Layers and blocks are specified in rectangle boxes; " $\times 5$ " indicates that five CNN blocks are tandem connected before being connected to the bidirectional GRU layer. The output layer at the bottom contains the probabilities predicted by the model for each of the nine types of the CAs classification. The type with the highest probability is the type predicted by the model for the input of an ECG signal.

C. Performance Evaluation

1) F1-score

F1-score is an evaluation metric that considers both Precision and Recall by:

$$F1 - score = 2 \times \frac{Precision \times Recall}{Precision + Recall},$$
(1)

$$Precision = \frac{IP}{TP + FP},$$

$$TP$$

$$TP$$

$$(2)$$

$$Recall = \frac{II}{TP + FN},$$
(3)

where true positive (TP), false positive (FP), true negative (TN), and false negative (FN) are defined by

$$TP_j = \left| \left\{ (x_i, y_i) \middle| y_i = f_j(x_i) = 1, \ 1 \le i \le N \right\} \right|, \tag{4}$$

$$FP_{i} = \left| \left\{ (x_{i}, y_{i}) \middle| y_{i} = 0 \text{ and } f_{i}(x_{i}) = 1, \ 1 \le i \le N \right\} \right|, (5)$$

$$TN_j = \left| \left\{ (x_i, y_i) \middle| y_i = f_j(x_i) = 0, \ 1 \le i \le N \right\} \right|, \tag{6}$$

$$FN_j = \left| \left\{ (x_i, y_i) \middle| y_i = 1 \text{ and } f_j(x_i) = 0, \ 1 \le i \le N \right\} \right|, (7)$$

where (x_i, y_i) is the input and label of the *i*th sample out of *N* samples, along with the prediction from a *C*-categorical classifier $f = (f_1, \ldots, f_j, \ldots, f_C) \in \{0, 1\}^C$; each categorical prediction f_j is binary-valued, and $|\cdot|$ measures the cardinality of a set. The overall F1-score can be derived by averaging the F1-scores of all categories.

2) Predictive Power Recovery

To quantify the recovery of the CAs classification accuracies of the HMC by its high-resolution ECG input with the enhanced low-resolution ECG input, a new evaluation metric called predictive power recovery (PPR) is defined as follows:

$$PPR = \frac{E_R - C_N}{C_P - C_N},\tag{8}$$

where C_P and C_N are the F1-scores of the DL models as positive and negative controls, respectively, in the CAs classification trained at high-resolution and low-resolution ECG signals, respectively. E_R is the F1-score of the HMC with an enhanced low-resolution ECG input. Ideally, the relation $0 \le C_N \le E_R < C_P \le 1$ is expected to hold, such that $0 \le PPR < 1$. However, cases exist in which the relation fails (see Fig. 7), and the corresponding discussions can be found in Experimental Results.

3) 10-fold Cross-validation Procedure of Machine Learning The data were randomly divided into 10 equal parts to set up an 8-1-1 training, validation, and test scheme for machine learning. Under such data splitting, the model was trained for 100 epochs (which refers to one complete cycle feeding training dataset) to generate 100 models (one model at each epoch), and the model with the best performance on the validation set was selected as the best model for this training process. Subsequently, the best model was used to compute the F1-score on the test set. The procedure was repeated 10 times to complete a 10-fold training; thus, the 10 best models were selected for each fold. The median F1-score for the overall and each CAs label, including the normal type, for the 10 test sets was calculated using the Scikit-learn package [64]. The training process was implemented using the ADAM optimizer [65].

D. Traditional Spline Interpolation for Up-Sampling

Spline interpolation is a useful technique for enhancing low-resolution ECG signals from low sampling frequencies. Originally, spline was a term for curves bent to pass through a number of existing data points.

Spline interpolation is realized by a special type of interpolation that fits many piecewise low-degree polynomials between local data pairs, for example, fitting two polynomials between each of the pairs of three points, instead of fitting a single degree-three polynomial to all of them. Therefore, the oscillation at the edges of an interval can be reduced using low-degree polynomials for interpolation. In the numerical analysis, the spline interpolation curves are defined by a set of polynomials $\{q_i\}$ with $y = q_i(x)$ satisfying two data pairs (x_{i-1}, y_{i-1}) and (x_i, y_i) , and i = 1, 2, ..., n. There are *n* polynomials and n + 1 knots in this case: the first polynomial ends at (x_0, y_0) , and the last polynomial ends at (x_n, y_n) .

E. Proposed SRECG

Instead of using traditional spline interpolations to enhance the low-resolution ECG signals, we propose an SR-based method called SRECG, as shown in Fig. 3. The SRECG is modified from a well-known architecture, SRResNet [66] to have one convolution layer and 16 residual blocks [67],



Fig. 3. Architecture of SRECG (denoted by the function f). X^L is the low-resolution ECG signals input, and $f(X^L)$ is the model output as the enhanced ECG signals. Layers and blocks are specified by the rectangles; "×16" indicates that 16 residual blocks are tandem connected before joining the convolution layer.

each of which contains two convolution layers with residual connections, followed by four convolution layers for both up-sampling and signal reconstruction. All convolution layers share the same kernel size of 16 and kernel number of 64, except for the last two up-sampling convolution layers with kernel numbers of 320 (for five times up-sampling) and 128 (for two times up-sampling), respectively. In the residual blocks, batch normalizations were also used to adjust and rescale the input from the two convolution layers (downsampling layers were not applied). LeakyReLU activation functions were applied between convolution layers before they were connected to the residual blocks and within up-sampling. Finally, batch normalization is used in the convolution layer following the residual blocks. This architecture was built using the Keras package supported by TensorFlow in GPUs [62], [63].

In Fig. 4, the low-resolution ECG data are fed into this architecture for training under the same 10-fold crossvalidation procedure. Throughout the training process, the model attempts to reconstruct an enhanced ECG from a lowresolution signal. In contrast to the conventional SR task, which merely increases the resolution of an imaging system by minimizing the signal mean square error (MSE) difference, we intended to combine the CAs classification results of another DL-based judge model, which is, HMC, for assistance. Considering this goal, a special loss function was designed to simultaneously optimize the ECG signal reconstruction and predictability for CAs classification, wherein the final enhanced output is subsequently fed into the HMC for CAs judgment. Notably, the HMC was independently pre-trained with high-resolution ECG signals using the same 10-fold cross-validation scheme. The customized joint loss L_J for the SRECG is defined as follows:

$$L_J(\widehat{y}, y, \widehat{z}, z) = \gamma L_R(\widehat{y}, y) + (1 - \gamma) L_C(\widehat{z}, z), \gamma \in [0, 1], \quad (9)$$

where L_R denotes the MSE regression loss between the highresolution ECG signals extity and the enhanced ones \hat{y} :

$$L_{R}(\widehat{y}, y) = \frac{1}{M} \sum_{i=1}^{M} (\widehat{y}_{i} - y_{i})^{2}, \qquad (10)$$

where *M* denotes the total number of maximum time steps of *y* and \hat{y} . The categorical cross-entropy (CCE) loss L_C measures the difference between the true labels of HMC $z = (z_1, \ldots, z_C) \in [0, 1]^C$ and that of predictions $\hat{z} = (\hat{z}_1, \ldots, \hat{z}_C) \in (0, 1]^C$ by

$$L_C(\widehat{z}, z) = -\sum_{k=1}^C z_k \log(\widehat{z}_k), \qquad (11)$$

In our model, we let $z = f_{judge}(y)$ and $\hat{z} = f_{judge}(\hat{y})$ serve as the CAs probabilities using a (fixed) HMC f_{judge} such that the SR loss

$$L_J(\widehat{y}, y) = L_J(\widehat{y}, y, f_{judge}(\widehat{y}), f_{judge}(y)), \qquad (12)$$

is determined once the enhanced ECG signals \hat{y} are generated. It is also understood that all the above losses are averaged over the sample numbers for normalization during the actual implementation.

III. EXPERIMENTAL RESULTS

A. Impacts of Sampling Frequency on DL-Based CAs Classification

A range of sampling frequencies from 500 to 1 Hz of ECG signals from 12 different lead sources was tested to explore the impact of sampling frequency on the DL-based multiclass CAs classifier in Fig. 2 under the 10-fold cross-validation procedure (see Fig. 5). In our preliminary results of ECG among different leads and sampling frequencies, it was observed that the 25 Hz sampling frequency lost most of the information for the overall CAs classification, which was in contrast to the case of the 250 Hz sampling frequency that preserved the most information. Based on this observation, we applied the traditional spline interpolation method and the proposed SRECG model to enhance 25 Hz input to improve its F1-score of HMC, trained at 250 Hz.

B. Effects of Traditional Spline Interpolation Methods

Four representative traditional spline interpolation methods were demonstrated for 1-dimensional signals: spline interpolation using polynomials of degree zero (I_0), degree one (I_1), degree two (I_2), and degree three (I_3). Table I shows the comparison of median overall F1-scores of CAs classification from



Fig. 4. Flowchart of the proposed SRECG for enhancing low-resolution ECG data. The SRECG *f* receiving the low-resolution ECG data X^L as input generates enhanced ECG prediction $f(X^L)$ as output by considering the joint loss L_J combining the regression loss L_R and the CAs classification loss L_C in Equation (1). The detailed structures of DL-based CAs classification model and SRECG are depicted in Figs. 2 and 3, respectively.



Fig. 5. Predictive power changes of DL-based multiclass CAs classifiers trained at downsampled ECG data. 12 lines with different colors represent the median overall F1-scores on the 10-fold tests of the CAs classification model in Fig. 2, which are trained at ECG data of different lead sources, sampled at different frequencies (500, 250, 125, 100, 50, 25, 10, 5, 2, 1 Hz).

the C_N (trained at 25 Hz), C_P (trained at 250 Hz), and HMC (trained at 250 Hz ECG) with spline-interpolated input from 25 Hz. Under different ECG lead sources, we observed that none of the interpolations F1-scores surpassed C_N. Figure 6 shows the signal reconstruction performance. The ECG waveforms of a lead II heartbeat demonstrated that I₁ had better reconstruction ability, resulting in the lowest MSE of 1.11% and the highest Pearson correlation coefficient (R) of 72.12%, in contrast to I₀ (MSE:3.71%; R:24.55%), I₂ (MSE:1.26%; R:71.74%), and I₃ (MSE:1.32%; R:70.73%).

From Table I and Fig. 6, we derived the observation that although traditional spline interpolations obtained good ECG signal reconstructions, CAs classifications (via F1-scores) were better preserved by the SRECG model (see C_N in Table II), which indicates that traditional spline interpolation does not recover sufficient CAs information during ECG signal enhancement.

C. Joint Effects of Two Different Training Losses for SRECG: Reconstruction and Classification

The composite loss function of the SRECG in Equation (9) was designed to simultaneously optimize two distinct objectives: CCE for CAs classification and MSE for signal reconstruction. Therefore, the ratio of the two losses can be weighted by a constant $\gamma \in [0, 1]$. We demonstrate three representative loss conditions in Equation (9): pure regression loss (L_R), pure classification loss (L_C), and half-mixture joint loss (L_J) by $\gamma = 1$, 0, and 1/2, respectively.

Table II shows the comparison of the median overall F1scores of C_N (trained at 25 Hz), C_P (trained at 250 Hz), and the HMC, trained at 250 Hz ECG, with SRECG enhanced input from 25 Hz by various losses.

Under different ECG lead sources, the highest F1-scores in the most enhanced CAs classification tasks were achieved under L_J, with the exception of V1, V2, and V3. In Fig. 6, the ECG waveforms of a lead II heartbeat show that: (1) L_R has better reconstruction power with the lowest MSE (0.14%) and highest R (97.10%); (2) ECG signals enhanced by L_C may not present a clear reconstruction capability (MSE:35.05%; R:-2.72%) and resemble ECG signals. Such enhancements derived higher F1-scores than those obtained by L_R (Table II), suggesting that the comprehension of machines can sometimes be beyond human intuition. (3) ECG signals enhanced by L_J share the characteristics of those enhanced by L_R and L_C (MSE:5.40%; R:50.49%). Thus, L_C may work as a constraint as both a penalty and incentive, restraining SRECG from overfitting and concentrating on crucial latent features.



Fig. 6. Enhanced ECG signals visualization of different spline interpolation degrees and SRECG losses. Representative ECG waveforms of a lead II heartbeat of 1-second lapse with (a) four spline interpolation degrees: I_0 , I_1 , I_2 , I_3 , and (b) three losses: L_R , L_C , L_J . The horizontal axis has the unit of time and the vertical axis indicates the ECG amplitude. Their MSEs and Rs of one second duration are also indicated in the subtitles to show the corresponding reconstruction quality.

 TABLE I

 MEDIAN OVERALL F1-SCORES (%) OF HMCs TRAINED AT 250 HZ WITH DIFFERENT LEADS AND SPLINE INTERPOLATION DEGREES

Leads / degrees	CN	Io	I ₁	I ₂	I3	Ср
Ι	61	56	47	43	39	73
II	61	57	47	29	24	71
III	52	50	45	36	31	65
aVR	63	56	41	38	36	75
aVL	49	49	42	38	36	64
aVF	57	56	52	35	29	70
V1	61	59	50	45	42	69
V2	55	34	32	23	18	71
V3	61	33	22	18	16	73
V4	63	39	20	13	12	74
V5	62	45	18	14	12	76
V6	62	53	31	26	22	75

The bold numbers indicate the highest scores close to their C_P.

 C_N : negative control (trained at 25 Hz), I_0 : degree zero spline interpolation, I_1 : degree one spline interpolation, I_2 : degree two spline interpolation, I_3 : degree three spline interpolation, C_P : positive control (trained at 250 Hz).

D. PPRs in CAs Classification With SRECG by Joint Loss

The previous F1-scores (Table II) demonstrated that, in most cases, SRECG by L_J can increase the CAs classification accuracy of HMC (trained at 250 Hz) from low-resolution ECG signals (25 Hz). Therefore, another experiment with L_J was conducted to further investigate the enhancement behavior of CAs classification accuracy, with PPR as a new metric for different ECG lead sources and CAs. The results in Fig. 7 show that the median PPR can reach up to overall 58%, yet variations across different leads and CAs exist. To visualize the median PPRs among the different leads and CAs, we labeled the median PPRs with color gradients based on their values. The colors from blue to green correspond to the median PPR values from high to low, and the black and red colors indicate anomalous median PPR values. The white color indicates the median PPR value that is not applicable (n/a) because its C_P , C_N , and E_R are all zero.

From the color labeling in Fig. 7, it is noted that: (1) lead aVR, V4, V5, and V6 are shown to have great potential in the recovery of overall CAs classification accuracy (median PPR ranging from to 56-58%); (2) in contrast, lead V3 has the lowest median overall PPR:16%; (3) the recovery capability of different CAs is also observed to vary from lead to lead, for example, STE could be well recovered in lead I, II, III, aVR, V2, V3, V5, and V6 (median PPR \geq 50%), but not in aVF, V1, and V4 (median PPR < 50%). However, there are

Leads															
		Ι	Π	III	aVR	aVL	aVF	V1	V2	V3	V4	V5	V6		
CAs	Normal	0	70	34	65	63	32	44	18	40	44	50	58		≥100
	AF	38	11	36	43	47	27	48	17	0	38	55	67		90-99
	I-AVB	39	55	70	42	71	58	58	85	53	68	77	65		70-89
	LBBB	0	70	55	68	50	52	131	-13	4	67	42	98		60-69
	RBBB	19	41	35	54	33	61	31	-11	43	56	61	62		50-59
	PAC	57	32	26	67	49	44	62	34	25	25	65	62		30-49
	PVC	36	12	51	81	28	52	47	40	27	20	65	62		10-29
	STD	-39	10	69	40	64	50	23	60	19	70	44	65		0-9
	STE	75	97	53	106	n/a	0	0	105	75	33	67	54		n/a
	Overall	43	40	42	57	48	43	39	47	16	56	58	57		<0

Fig. 7. Median PPRs (%) of HMCs trained at 250 Hz with SRECG by LJ. Blue labels and green labels indicate relatively high and low median PPR values, respectively. Black labels indicate median PPR values greater than or equal to 100%. Red labels indicate negative median PPR values (<0%). The white label indicates the median PPR value that is not applicable (n/a) because its CP, CN, and ER are all zero.

TABLE II MEDIAN OVERALL F1-SCORES (%) OF HMCs TRAINED AT 250 HZ WITH DIFFERENT LEADS AND SRECG LOSSES

Leads / losses	CN	LR	Lc	L_J	Ср
Ι	61	60	65	66	73
II	61	64	64	66	71
III	52	51	56	58	65
aVR	63	68	71	71	75
aVL	49	48	55	56	64
aVF	57	59	61	62	70
V1	61	62	65	64	69
V2	55	62	64	61	71
V3	61	65	67	65	73
V4	63	65	65	68	74
V5	62	65	70	70	76
V6	62	57	69	69	75

The bold numbers indicate the highest scores close to their C_P.

C_N: negative control (trained at 25 Hz), L_R: SRECG by pure regression loss, L_C: SRECG by pure classification loss L_J: SRECG by half-mixture joint loss, C_P: positive control (trained at 250 Hz).

six anomalies (median PPR < 0% or median PPR \ge 100%) in Fig. 7, which in turn indicates that the F1-scores of HMCs with SRECG-enhanced ECG input are worse than those of C_N (median PPR < 0%) or better than C_P (median PPR $\geq 100\%$). This leads us to wonder if the enhanced signals may have unexpected predictive power changes for verifying the CAs.

E. Wilcoxon Signed-Rank Test for Four PPR Anomalies in CAs Classification With SRECG by Joint Loss

Conventionally, in statistics, a Wilcoxon signed-rank test is used to confirm that two sampled populations share significant differences if their p-value is less than 0.05. Therefore, the above six anomalies were subjected to the Wilcoxon signedrank test in comparison to their C_N or C_P, which are the 10-fold test F1-scores of low-resolution multiclass CAs classifiers trained at 25 Hz ECG signals (median PPR < 0%) or 250 Hz ECG signals (median PPR \geq 100%). However, we found that none of the anomalies passed the criteria (p-value < 0.05); therefore, we cannot assert that there are significant differences compared with C_N or C_P . Thus, we could not conclude that there were unexpected predictive power changes for verifying CAs in these six anomalies. Although the case of LBBB in lead V1 is still intriguing with a large increase (median PPR:131%), the main reason was no significant difference between C_N and C_P such that $C_P - C_N \approx 0$ which led to an enlarged value in $PPR = \frac{E_R - C_N}{C_P - C_N}$ according to its definition. Therefore, its increment of PPR is not valid.

IV. DISCUSSION

CE play an important role in smart healthcare to improve the quality of life. P/W devices incorporated with CE used for medical purposes have provided simple ways to monitor the health conditions of consumers in daily life, such as bracelets and wrist gadgets [68]. Several studies related to smart healthcare systems have utilized such devices to support visually impaired individuals and prevent falls in the elderly [69], [70], [71]. In addition, these devices can provide long-term physiological signal monitoring for clinical diagnosis and evaluation, such as electroencephalogram [72], ECG [73], [74], [75], and near-infrared spectroscopy [76].

Collectively, the combination of cloud-based DL algorithms with P/W devices is becoming a trend in smart healthcare systems. However, there are certain limits tied to the current capability of the hardware, such as battery life and computing power, for ECG long-term monitoring systems implemented in CE. With respect to the limitation on energy consumption, it is understood that more power is required when a higher frequency is used to record ECG signals [9], [10], [11]. Therefore, in this study, we attempted to alleviate this problem by recovering the predictive power of the HMC with a low sampling frequency input via the proposed DL-based SRECG framework, which is a novel SR-based method. The proposed SRECG is designed for P/W devices with the cloud-based workflow of a smart healthcare system (see Fig. 1).

For a conventional SR task, the loss function is typically only regression loss L_R, whereas our experiments indicated that enhancements by L_R alone did not suffice to improve the CAs classification accuracies of the HMCs. As a result, it is viable to introduce the concept of teacher-student learning by including the CCE loss using the judge model's predictions, which are from a well-pretrained HMC for CAs classification [77]. Correspondingly, the experimental results showed that our joint loss L_J obtained better performance on CAs classification (via F1-scores) in most of the ECG lead sources, with a few exceptions for leads V1, V2, and V3 (Table II). Moreover, the predictive power in the models of lead V5 trained at 250 Hz sampled ECG signals achieved up to 58% of their median overall PPR in the classifications of 25 Hz sampled ECG signals (see Fig. 7). It is interesting that the best reconstructed ECG signals (with the lowest MSEs and highest Rs) did not have the highest F1-scores when applying to HMCs for CAs classification. A possible reason for this is that the proposed model combining reconstruction and classification tasks pays more attention to reconstructing the features or signals that could help in ECG classification. In other words, the proposed approach is limited to processing fine-grained ECG signals that are relatively ineffective for classification tasks. Therefore, it could reach the highest model F1-scores without holding the lowest MSEs and the highest Rs. A similar phenomenon occurs in the field of automatic speech recognition (ASR) in a noisy environment [78], [79], [80]. They pointed out that removing background noise using speech enhancement models can improve signal quality, but it often degrades ASR performance. This is because the distortion caused by speech enhancement often deteriorates the ASR performance.

Clinically, DL-based ECG diagnosis of CAs has shown its feasibility and significance in improving diagnosis accuracy compared to that of general physicians and cardiologists [81], [82]. Different leads of ECG signals provide different ECG features for physicians and cardiologists to recognize specific types of arrhythmia [83], [84], [85], [86], [87]. Our experimental results in Fig. 7 show that lead V6 obtained the highest and standout median PPR of 98% in LBBB among all leads, which concurs with the clinically known LBBB diagnosis criteria by distinguishing QRS complex morphology at leads I, aVL, V1, V2, V5, and V6 [83]. In contrast, lead aVR also demonstrated the highest and distinguished median PPR of 81% in PVC, similar to lead V4 in STD (median PPR:70%) and lead V6 in AF (median PPR:67%). These observations suggest that the subtle ECG features hidden in leads aVR, V4, and V6 may serve as important judging criteria for DL models to distinguish PVC, STD, and AF, instead of human eyes.

Although the results demonstrated the capability to recover the predictive power of HMC trained at high-resolution ECG signals (250 Hz) with low-resolution input (25 Hz) by our SRECG, it becomes more challenging when trying to recover from even lower-resolution signals (< 25 Hz) to higherresolution signals (> 250 Hz). Our method requires more validation on various types of datasets, for example, noisy datasets collected by embedded devices, to verify the prediction stability and effectiveness in practical situations. In the absence of a systematic evaluation approach and the lack of standardized datasets, it is difficult to address these limitations at the present stage. We consider these obstacles as part of our future study.

On the other hand, with the increasing concern about medical data privacy issues [88], it is necessary to propose encryption methods to make the data undistinguishable to satisfy the requirements of new privacy-preservation regulations, for example, GDPR [89]. In Fig. 6, we accidentally found that the ECG signals enhanced by the classification loss L_C and joint loss L_J can generate undistinguishable ECG waveforms while improving the predictive power for CAs classification (Table II), which may provide an alternative way to encrypt ECG data.

As machine learning and DL methods continue to advance and become user-friendly via P/W, the role of ECG will gradually open as a physiological information container. Subsequently, a DL model can be used at the right time to extract information from the container for ignored or unattended predictions such as gender and age [90]. Therefore, it is promising to expect ECG to provide more diagnostic power in many upcoming applications on P/W devices.

V. CONCLUSION

To support automatic clinical diagnosis and long-term smart healthcare services on P/W devices while incorporated with CE, this work develops a novel DL-based SRECG as a cloud processor to enhance low-resolution ECG signals sampled from low frequency for CAs classification of HMC to achieve better accuracy. In future work, we will not only further investigate the effect of training SRECG with compound loss using different lead sources and their combinations but also explore different preprocessing methods for SRECG. Since DL-based diagnosis of ECG signals has great potential in deciphering other physiological conditions, it is worth applying SRECG for P/W devices in other diagnostic criteria of healthcare, such as mortality and heart failure, in addition to CAs.

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