

BULLETIN OF THE POLISH ACADEMY OF SCIENCES TECHNICAL SCIENCES, Vol. 69(3), 2021, Article number: e136921 DOI: 10.24425/bpasts.2021.136921

# A new method of cardiac sympathetic index estimation using a 1D-convolutional neural network

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**Abstract.** Epilepsy is a neurological disorder that causes seizures of many different types. The article presents an analysis of heart rate variability (HRV) for epileptic seizure prediction. Considering that HRV is nonstationary, our research focused on the quantitative analysis of a Poincare plot feature, i.e. cardiac sympathetic index (CSI). It is reported that the CSI value increases before the epileptic seizure. An algorithm using a 1D-convolutional neural network (1D-CNN) was proposed for CSI estimation. The usability of this method was checked for 40 epilepsy patients. Our algorithm was compared with the method proposed by Toichi et al. The mean squared error (MSE) for testing data was 0.046 and the mean absolute percentage error (MAPE) amounted to 0.097. The 1D-CNN algorithm was also compared with regression methods. For this purpose, a classical type of neural network (MLP), as well as linear regression and SVM regression, were tested. In the study, typical artifacts occurring in ECG signals before and during an epileptic seizure were simulated. The proposed 1D-CNN algorithm estimates CSI well and is resistant to noise and artifacts in the ECG signal.

Key words: epilepsy; seizure detection; seizure prediction; convolutional neural network; deep learning; ECG; HRV; cardiac sympathetic index.

## 1. Introduction

Around 50 million people worldwide have epilepsy, making it one of the most common neurological diseases globally [1]. In some cases, it is not possible to prevent epileptic seizures by using drugs or resection of the part of the cerebral cortex that is the source of seizures [2]. Certain changes in the autonomic nervous system (ANS) in people suffering from epilepsy can be observed. Changes and abnormal heart rhythm are among the most important symptoms. The following changes occur before and during seizures: tachycardia, bradycardia and asystole, conduction disturbances, and heart ischemia. Their description is presented in many works [3–6]. As shown, heart rate changes are associated with sudden unexpected death (SUDEP) in patients with epilepsy. Most researchers believe that insular cortex changes lead to the dysregulation of the ANS in patients with temporal lobe epilepsy (TLE).

Therefore, methods of predicting epileptic seizures are being sought. For this purpose, attempts have been made to use electrophysiological signals, in particular EEG [7–9]. The effectiveness of the developed methods is not fully satisfactory. Therefore, any additional physiological signal that can support the prediction of a seizure is valuable. In this task, attempts are made to use the ECG signal [10]. Numerous methods of quantifying the heart rate variability have been proposed [11–13]. One of the features of the ECG signal is the so-called cardiac

the calculation of the CSI using both the standard Toichi algo-

rithm and the proposed method. The presented research is an

extension of the study described in the conference paper [25].

sympathetic index (CSI). CSI is based on the quantitative analyses of Poincare plot features. Several studies indicate that it is

possible to use CSI to detect and predict seizures [14–17]. For

CSI calculation, the method proposed by Toichi et al. in [18] is

most often used. In this article, the symbolic name "Toichi" is

used to refer to the algorithm described in this work. To deter-

mine CSI, first R-waves must be detected in the ECG signal.

For this purpose, the well-known Pan-Tompkins algorithm [19]

Video EEG (vEEG) with simultaneous ECG was recorded in patients with refractory temporal epilepsy for long-term monitoring of epileptic seizures during presurgical evaluation. The signals were acquired at the Medical University of Warsaw.

Manuscript submitted 2020-10-08, revised 2020-12-23, initially accepted for publication 2021-02-13, published in June 2021

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or another method can be used. Next, the RR intervals should be analysed using the Poincare plot [20].

The purpose of the article is to use a 1D-convolutional neural network [21–23] to calculate the CSI values based on the registered ECG signal. For a trained neural network, there is no need to perform R-wave detection [24] and Poincare plot analysis. It was assumed that deep learning, with many training examples from many patients, would allow the network to work effectively. A 1D-convolutional neural network was used to estimate the CSI. Then it was checked whether the CSI was reliable for the prediction of epileptic seizures. It was also examined how the loss of part of the recorded signal or artifacts affected

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In 40 patients who underwent epilepsy surgery in the years 2005–2015, ictal vEEG recordings provided localization information and identification of temporal seizure type as well as ECG recordings. After non-invasive presurgical evaluation, all patients had anterior tailored lobectomy with pre-resection electrocorticographic recordings (iECoG). The temporal ictal onset zone was localized based on seizure ictal semiology and ictal patterns. Thirty-two biosignals were registered (1 ECG channel and 31 EEG channels) using an EEG DigiTrack amplifier. The ECG signal was recorded with the sampling frequency fs = 250 Hz. For EEG registration, cup-shaped electrodes were used, and for ECG precordial disposable electrodes were located at V1 and V2 positions. During signal registration, patients were either sitting or lying, but they could freely move their heads and limbs. The registration time was from several minutes to several hours.

#### 3. Methods

- **3.1. Pre-processing of the ECG signal.** The disturbance caused by the power line frequency (50 Hz) was removed from the ECG signal. For this purpose, a narrow-band (48–52 Hz) Butterworth digital filter was used [26, 27]. For the Toichi method, the R-waves were determined in the ECG signal using the Pan and Tompkins algorithm.
- **3.2. Cardiac sympathetic index.** Poincaré plots were used to analyse the nonlinear dynamics of ECG signals [28–32]. Each point in the plot is specified as n = 1, 2, 3, ...N ( $RR_n, RR_{n+1}$ ), where N represents the number of recorded R-waves. Such mapping reveals the correlation between consecutive RR intervals. The mapping gives useful information on short-term and long-term fluctuations of RR intervals. The mapping is shown in Fig. 1, along with the standard deviation parameters SD1 and SD2.

The SD1 shows beat-to-beat rapid changes, which are more related to respiratory sinus arrhythmia. However, SD2 describes

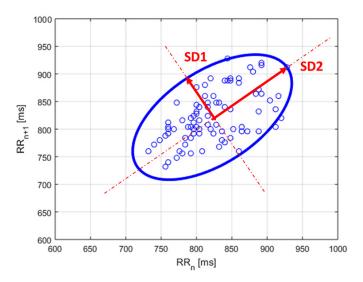


Fig. 1. Example of calculating SD1 and SD2 based on Poincaré plots

long-term beat-to-beat changes [33]. cardiac sympathetic index CSI = SD2/SD1 is calculated to describe the relationship between these components [18]. The SD1 and SD2 values in Poincaré mapping directly depend on the statistical values of the standard deviation of heart rate and two consecutive intervals of R peaks. Ellipses are described so that at least 95% of the RR data is in the ellipse area.

**3.3. 1D-convolutional neural network.** To determine the CSI value, we proposed the use of a 1D-convolutional neural network (1D-CNN). A one-minute ECG signal (15,000 samples) is fed to the network input, sampled at a frequency of fs = 250 Hz (Input Layer). The other layers in the network are Convolution, Batch Normalization, ReLU, MaxPooling, Dropout, Fully Connected, and Regression Output [21, 34–36]. The network structure is shown in Table 1.

Table 1 Structure of the neural network used

	Structure of the neural network used				
No.	Name of Layer	Parameters			
1	Input Layer	15,000 samples with zero centre normalization			
2	Conv_1	20 filters 150x1 convolutions with stride 1 and padding same			
3	Batchnorm_1	Batch normalization with 20 channels			
4	Relu_1	ReLU			
5	Maxpool	2x1 max pooling with stride 2 and padding 0			
6	Conv_2	40 filters 150x20 convolutions with stride 1 and padding same			
7	Batchnorm_2	Batch normalization with 40 channels			
8	Relu_2	ReLU			
9	Conv_3	60 filters 10x40 convolutions with stride 1 and padding same			
10	Batchnorm_3	Batch normalization with 60 channels			
11	Relu_3	ReLU			
12	Conv_4	60 filters 10x60 convolutions with stride 1 and padding same			
13	Batchnorm_4	Batch normalization with 60 channels			
14	Relu_4	ReLU			
15	Conv_5	60 filters 20x60 convolutions with stride 1 and padding same			
16	Batchnorm_5	Batch normalization with 60 channels			
17	Relu_5	ReLU			
18	Conv_6	60 filters 10x60 convolutions with stride 1 and padding same			
19	Batchnorm_6	Batch normalization with 60 channels			
20	Relu_6	ReLU			
21	dropout	50% dropout			
22	fc	1 Fully connected layer			
23	Regression output	Mean-squared-error with response			

The batch normalization method can be used to solve the problem of changing the distribution of hidden layer inputs [37]. This method is based on the proven property that the whitening of classifier inputs facilitates the optimization process during training. It also pays to normalize the inputs of all hidden layers in the network. rectified linear unit (ReLU) is the most commonly used activation function in the field of deep learning [38]. Neural networks equipped with ReLU functions obtained better classification accuracy than networks equipped with the hyperbolic tangent (tanh) function. Moreover, the authors of [39] found that the use of ReLU accelerates the training of the neural network four times, compared to the neural network using tanh. The most common operations that occur in the subsampling layers are MaxPooling [37]. These operations aggregate a group of adjacent samples, replacing them with samples with the maximum value. Dropout is a technique that improves the performance of neural networks across many different applications, including object classification, natural language processing, and scientific data analysis [40]. Dropout overcomes this problem by providing a way to approximately combine multiple neural networks into a single model. The method is to remove certain neurons temporarily during training. The group of neurons to be removed is selected randomly. The holding probability of a neuron is a tunable hyperparameter called the dropout rate. Neurons are drawn and removed each time a batch of samples is delivered to the network. In practice, a neuron is removed by multiplying its output by zero. The FC layer refers to a fully connected layer like in the case of a classical neural network [41].

In our research, electrocardiographic signals recorded for 40 patients were used to train CNN. Pre-seizure ECG fragments, starting ten minutes before the seizure, were selected. Windows lasting 1 minute and moving every 1 second were used for signal analysis. In that way, 540 examples were generated for each patient. A total of 21,600 examples were created for 40 patients. The data was randomly divided into three parts: 80% (17,280) for training, 10% (2160) for validation, and 20% (4,320) for testing.

The ADAM algorithm with the initial learning rate parameter of 0.001 was used to train the network. During training, GPU graphics processors with GeForce GTX 1070 graphics card were used. The process of training the network lasted several dozen minutes.

Figure 2 presents the procedure for the Toichi algorithm when calculating the CSI. The recorded ECG signal lasting one minute (ECG signal) is digitally filtered to remove the power line frequency (Notch filter). Then R-waves are detected in the ECG signal (R-wave detection). The well-known Pan and Tompkins algorithm is used for this purpose. In the next step, the time between the individual RR-waves is calculated (RR interval calculation). The calculated RR intervals allow the computation of the Poincare plot (Poincare plot). Based on the Poincare plot, the SD1 and SD2 values are calculated (SD1 and SD2 calculation). The determined values facilitate the calculation of the CSI coefficient.

Figure 3 presents the procedure for the algorithm we propose using 1D-CNN. The recorded ECG signal lasting one minute (ECG signal) is subjected to digital filtering, the purpose of

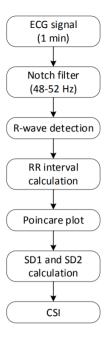


Fig. 2. CSI values calculated using the Toichi algorithm

which is to remove the power line frequency of 50 Hz (*Notch filter*). Then, each 1-minute segment of the ECG signal is fed to the input of the 1D-CNN. A trained 1D-convolutional neural network calculates the CSI.

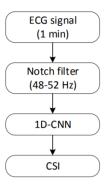


Fig. 3. CSI values calculated using the 1D-CNN

## 4. Results and discussion

To prevent overfitting of the neural network, a validation set was used. An expert verified all examples used for training, validation, and testing to assess the correct R-wave detection. The CSI calculation method proposed by Toichi et al. was considered as the reference, showing the correct values.

The difference between the estimations of the CSI value calculated using the Toichi algorithm and the 1D-convolutional neural network was calculated. We calculated the mean squared error (equal to 0.046) and the mean absolute percentage error (0.097) for the testing set as a measure of the difference. Additionally, to determine the correctness of the CSI estimation, the maximum, minimum, and average values of this index were calculated using both algorithms. The maximum CSI value

calculated using the Toichi algorithm was 7.873 while using a 1D-CNN 7.324. The minimum CSI value calculated using the Toichi algorithm was 1.001 while using a 1D-CNN 0.092. The average CSI value calculated using the Toichi algorithm was 1.956 while using a 1D-CNN 1.849. The obtained results proved the correct estimation of the CSI value by a 1D-CNN.

Figure 4 presents an example of the calculated CSI values with the Toichi and 1D-CNN algorithms for one of the patients. We can observe significant fluctuations in CSI values. Although the CSI values calculated with the Toichi method slightly differ from these calculated with a 1D-CNN, there is a large correlation between them,  $\rho = 0.97$ .

Figure 5 presents the RR interval variability for subsequent heartbeats. We can observe significant variation in the RR interval. Figure 6 shows Poincare plots for one patient for an ECG signal registered 10 minutes before the seizure (blue) and one minute before the seizure (red). For this patient, the CSI value calculated using the Toichi algorithm was 1.535 for ten minutes

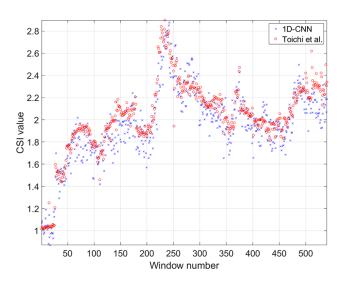


Fig. 4. CSI values calculated using the Toichi and 1D-CNN algorithms

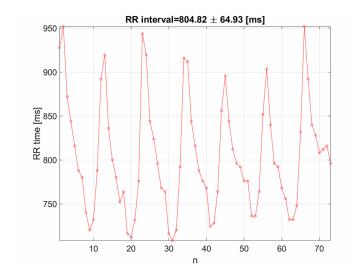


Fig. 5. Example of RR interval variation for the signal one minute before the seizure

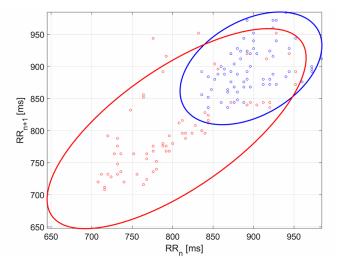


Fig. 6. Poincaré plots for one patient 10 minutes before the seizure (blue) and one minute before the seizure (red)

before the seizure and 2.417 for one minute before the seizure. In contrast, the CSI value estimated using a 1D-CNN was 1.549 for ten minutes before the seizure and 2.018 for one minute before the seizure.

The influence of disturbances that may occur during the registration of ECG signals on the CSI estimation error was also examined. For this purpose, a one-minute ECG signal recording was chosen, with correctly detected R-waves. The standard deviation for the selected ECG signal samples was  $\sigma=0.928$ . CSIs were calculated using the Toichi and 1D-CNN algorithms. Then, signal fragments (from 0.5 to 20 seconds) were removed from the centre of the ECG waveform, and CSI values were calculated again using the Toichi algorithm and a 1D-CNN. Such removal of the signal was intended to simulate the artifact of the temporary disconnection of the ECG electrode. Figure 7 shows the ECG signal fragment with the simulated removal of 8 seconds of the signal.

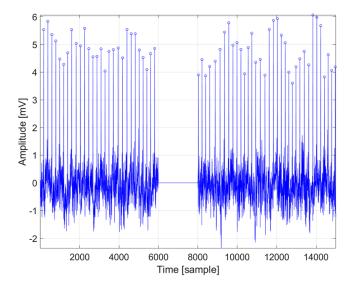


Fig. 7. A fragment of the ECG signal with the removal of 8 seconds of the signal

In the next stage, the ECG signal was disturbed by noise with a normal distribution and an amplitude of  $0.5\sigma$ ,  $1\sigma$ ,  $1.5\sigma$ , and  $2\sigma$  of the standard deviation of the ECG waveform. In this case, the signal noise was intended to simulate low-amplitude, long-term EMG artifacts. Then the ECG signal was disturbed by noise with a normal distribution and amplitude from  $2\sigma$  to  $6\sigma$  times greater than the ECG signal standard deviation. The duration of the signal disturbance ranged from 2 to 4 seconds. This signal noise was intended to simulate high-amplitude, short-term EMG artifacts. An example of such an artifact is presented in Fig. 8.

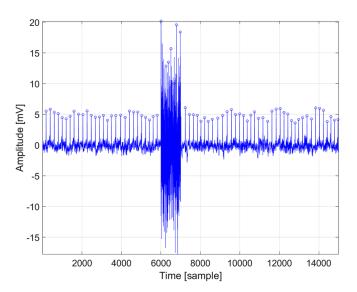


Fig. 8. A fragment of ECG signal with 4-second EMG artifact

Finally, the influence of sinusoidal disturbances was examined. Such kind of disturbance simulates floating electrode artifacts. For this purpose, a sinusoidal signal with amplitude 1 and frequencies from 0.05 Hz to 10 Hz was added to the ECG signal. An example of a disturbed signal with a sinusoidal artifact is presented in Fig. 9.

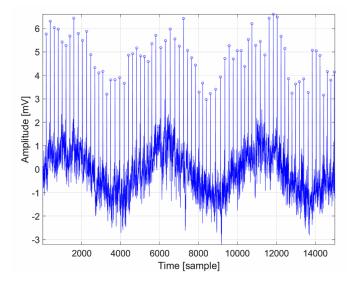


Fig. 9. A fragment of ECG signal with a floating electrode artifact

Then the CSI was calculated for each case using the Toichi and 1D-CNN algorithms. The calculated CSI values for the loss of the ECG signal fragments are presented in Table 2. For each calculated CSI value, an error was determined, that is, the difference between the calculated CSI value for the disturbed signal and the CSI value calculated for the undisturbed signal using the Toichi algorithm.

Table 2
CSI values for loss of fragments of the ECG signal

I am of simul	Toichi et al.		1D-CNN	
Loss of signal	CSI	Error	CSI	Error
Undisturbed	2.200	_	2.196	0.004
20.0 sec	1.101	1.099	2.037	0.163
10.0 sec	1.012	1.188	2.283	-0.083
4.0 sec	1.026	1.174	2.355	-0.155
2.0 sec	1.096	1.104	2.176	0.024
1.0 sec	1.265	0.935	2.152	0.048
0.5 sec	1.265	0.935	2.112	0.088

The proposed 1D-CNN algorithm estimates the CSI values much more accurately than the Toichi algorithm. Even for a loss of 20 seconds of the ECG signal, the CSI estimation correctness is acceptable. Table 3 presents the calculated CSI values for floating electrode artifacts.

Table 3
CSI values for floating electrode artifacts

Sinusoidal signal	Toichi et al.		1D-CNN	
	CSI	Error	CSI	Error
Undisturbed	2.200	_	2.196	0.004
0.05 Hz	1.920	0.280	2.233	-0.033
0.1 Hz	1.923	0.277	2.249	-0.049
0.2 Hz	1.607	0.593	2.223	-0.023
1 Hz	1.956	0.244	1.092	1.108
10 Hz	1.792	0.408	3.471	-1.271

For the ECG signal disturbed by a sinusoidal waveform with frequencies from 0.05 Hz to 0.2 Hz, we obtain a more accurate estimation of the CSI value using the proposed 1D-CNN algorithm than with the Toichi algorithm. Worse results of the CSI coefficient approximation were obtained for the sinusoidal signal in the range of 1 Hz to 10 Hz.

Table 4 presents the results of the calculated CSI values for ECG signal with added noise of normal distribution (simulated EMG artifacts). For the noise values from  $0.5\sigma$  to  $1.5\sigma$ , a more accurate estimation of the CSI value is obtained with the proposed 1D-CNN algorithm than with the Toichi algorithm. How-

ever, for a disturbance with an amplitude of  $2\sigma$ , it happened in one case that the value of the CSI coefficient determined by the Toichi algorithm was more accurate.

Table 4
CSI values for ECG signal with simulated low-amplitude long-term
EMG artifacts

Normal	Toichi et al.		1D-CNN	
distribution noise	CSI	Error	CSI	Error
Undisturbed	2.200	-	2.196	0.004
$0.5\sigma$	1.781	0.419	2.193	0.007
$0.5\sigma$	1.904	0.296	2.347	-0.147
$0.5\sigma$	1.616	0.584	2.103	0.097
1σ	1.220	0.980	2.403	-0.203
1σ	1.159	1.041	2.191	0.009
1σ	1.609	0.591	2.150	0.050
$1.5\sigma$	1.166	1.034	1.577	0.623
$1.5\sigma$	1.532	0.668	1.937	0.263
$1.5\sigma$	1.286	0.914	2.047	0.153
$2\sigma$	1.391	0.809	1.930	0.270
$2\sigma$	1.404	0.796	1.206	0.994
2σ	1.175	1.025	1.974	0.226

Table 5 presents the results of CSI value estimation for ECG signal with 2 seconds high-amplitude EMG artifacts. For the disturbance amplitude from  $2\sigma$  to  $6\sigma$ , again, the 1D-CNN algorithm gave better results than the Toichi algorithm. The presented results indicate that the 1D-CNN algorithm is more robust to artifacts specific to the ECG signal recordings.

Table 5
CSI values for ECG with simulated high-amplitude, short-term
EMG artifacts

Normal	Toichi et al.		1D-CNN	
distribution noise	CSI	Error	CSI	Error
Undisturbed	2.200	-	2.196	0.004
$2\sigma$	1.440	0.760	2.248	-0.048
$2\sigma$	1.332	0.868	2.177	0.023
$2\sigma$	1.065	1.135	2.238	-0.038
$4\sigma$	1.611	0.589	2.072	0.128
$4\sigma$	1.645	0.555	2.142	0.058
$4\sigma$	1.588	0.612	2.065	0.135
6σ	1.259	0.941	2.022	0.178
6σ	1.071	1.129	2.077	0.123
6σ	1.765	0.435	2.071	0.129

The CSI calculation time for one minute of the ECG signal was tested using the Toichi and 1D-CNN algorithms. The average calculation time for 1D-CNN was 0.0048 seconds, while for Toichi 0.0029. For both algorithms, the CSI calculation time is comparable and can be used in practice.

It was also checked whether an advanced neural network (1D-CNN) is really needed to approximate the CSI coefficient. For this purpose, the performance of MLP with one hidden layer was compared with the results obtained for 1D-CNN. During the research, the number of neurons in the hidden layer was changed. The Levenberg-Marquardt algorithm was used to train the MLP. The same data was used for the MLP and 1D-CNN: 17,280 examples were used for training; 2,160 for validation; and 4,320 for testing. Table 6 presents the mean squared errors for a particular number of neurons in the hidden layer of MLP. The obtained results indicate that the proposed 1D-CNN facilitates a much more accurate estimation of the CSI values. The smallest MSE error (0.637) for the MLP was obtained for 300 neurons in the hidden layer. This is a bigger order of magnitude than for the 1D-CNN.

Table 6
MSE error comparison for MLP and 1D-CNN

Neural network	Number of neurons in the hidden layer	MSE
MLP	1,000	0.704
MLP	500	1.600
MLP	300	0.637
MLP	100	0.726
MLP	50	0.667
MLP	10	0.804
1D-CNN	_	0.046

The performance of the 1D-CNN was also compared with other regression methods. Two algorithms, linear regression and SVM regression (quadratic kernel) were selected for comparison. The PCA method was used to reduce the input data set. For this purpose, components explaining 95% of the variance of the set were selected. The same data was used for regression and 1D-CNN training and testing. Table 7 presents the MSE errors for the testing set for individual regression methods. Again, the MSE error for 1D-CNN is a smaller order of magnitude.

Table 7
MSE error for the regression and 1D-CNN methods

Regression method	MSE
PCA + Linear Regression	0.878
PCA + SVM Regression	0.860
1D-CNN	0.046

Figure 10a presents an exemplary 5-second segment of the ECG signal fed to the input of the 1D-CNN and three exem-

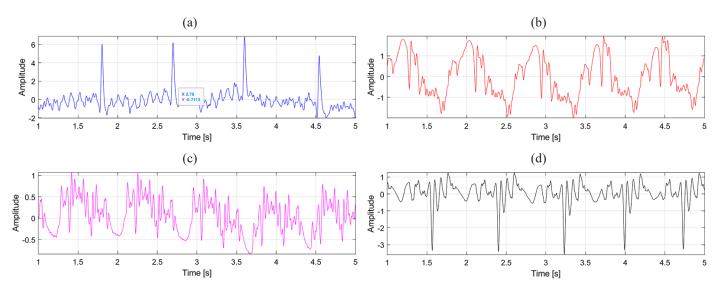


Fig. 10. a) An example of a 5-second segment of the ECG signal fed to the neural network input and the exemplary signals after the different filters created by the first layer of 1D-CNN; b) filter number one; c) filter number two; d) filter number three

plary ECG signals after passing through the network digital filters of the first convolutional layer (Figs. 10b–10d). The interpretation of the received signals is difficult. However, for the third signal (Fig. 10d), we can observe the effective "extraction" of R-waves. Further layers of the neural network use the created filters to calculate new features and, as a result, to calculate the CSI.

In another experiment, the influence of the size of the first convolution layer of 1D-CNN on the accuracy of the obtained results was checked. Table 8 shows the MSE error for sample filter size (50, 150, 300, and 500). In each case, very similar MSE error values were obtained. However, the best results were obtained with a filter size of 150.

Table 8
Comparison of the MSE error for 1D-CNN for different filter sizes of the first convolution layer

Filter size	50	150	300	500
MSE	0.049	0.046	0.056	0.064

To emphasize the purposefulness of the conducted research, it was additionally checked whether there is a chance of using CSI to predict or detect an epileptic seizure. CSI variability was calculated using the Toichi and 1D-CNN algorithms for each user. Then the result obtained for each of the 40 users was averaged. Figure 11 presents the average value of CSI for the Toichi and 1D-CNN algorithms 9 minutes before an epileptic seizure. For CSI calculated using Toichi and 1D-CNN, we can observe a significant increase about a minute before the seizure.

The mean pre-seizure CSI value for 1D-CNN is 2.205 and for the Toichi algorithm is 2.300 (the difference is 4.3%). Higher CSI values determined using the Toichi algorithm than using a 1D-CNN can be observed. Figure 12 shows the statistical distribution of the CSI values calculated using the Toichi algorithm and 1D-CNN. Using both the Toichi and CNN meth-

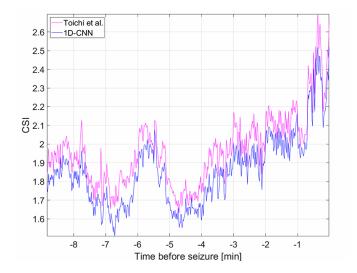


Fig. 11. CSI values averaged for 40 people for the Toichi and 1D-CNN algorithms 9 minutes before the epileptic seizure

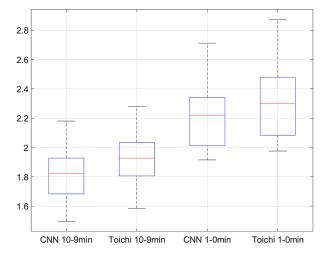


Fig. 12. Boxplot of CSI values calculated using Toichi and 1D-CNN algorithm for 9 to 1 and 1 to 0 minutes before the seizure

ods, it is possible to indicate a statistically significant increase (p > 0.001) of the CSI value a minute before the seizure.

This confirms the assumption that, in general, there is an increase in the CSI before epileptic seizures. However, to assess the effectiveness of such a prediction method, further research is required regarding the CSI variability over a longer period of time. This is outside the scope of this work.

# 5. Conclusions

The proposed structure of a 1D-CNN facilitates an effective approximation of the cardiac sympathetic index. The mean squared error is MSE = 0.046 and the mean absolute percentage error MAPE = 0.097. The proposed solution, based on 1D-CNN, facilitates a better estimation of the Cardiac Sympathetic Index than other regression methods and is more resistant to various types of disturbances. Unfortunately, training a 1D-CNN requires many examples from many people, and the ECG data must be of good quality. The use of a 1D-CNN for the CSI calculations is more reliable for an ECG signal containing disturbances that may occur during the ECG signal registration, especially during an epileptic seizure. The CSI variability indicates that there is a chance to use the ECG signal to predict or detect an epileptic seizure.

# REFERENCES

- [1] World Health Organization (WHO), "Epilepsy", 2019. Accessed: Jul. 10, 2020. [Online]. Available: https://www.who.int/newsroom/fact-sheets/detail/epilepsy
- [2] B. Sommer *et al.*, "Resection of cerebral gangliogliomas causing drug-resistant epilepsy: short- and long-term outcomes using intraoperative MRI and neuronavigation", *Neurosurg. Focus* 38(1), E5 (2015), doi: 10.3171/2014.10.FOCUS14616.
- [3] T. Harnod, C.C.H. Yang, Y.-L. Hsin, P.-J. Wang, K.-R. Shieh, and T.B.J. Kuo, "Heart rate variability in patients with frontal lobe epilepsy", *Seizure* 18(1), 21–25 (2009), doi: 10.1016/j.sei(zure.2008.05.013.
- [4] K. Jansen and L. Lagae, "Cardiac changes in epilepsy", *Seizure* 19(8),455–460 (2010), doi: 10.1016/j.seizure.2010.07.008.
- [5] R. Brotherstone and A. McLellan, "Parasympathetic alteration during sub-clinical seizures", *Seizure* 21(5), 391–398 (2012), doi: 10.1016/j.seizure.2012.03.011.
- [6] A. Van de Vel et al., "Non-EEG seizure detection systems and potential SUDEP prevention: State of the art: Review and update", Seizure 41,141–153 (2016), doi: 10.1016/j.seizure.2016.07.012.
- [7] U.R. Acharya, Y. Hagiwara, and H. Adeli, "Automated seizure prediction", *Epilepsy Behav.* 88, 251–261 (2018), doi: 10.1016/j. yebeh.2018.09.030.
- [8] G. Giannakakis, V. Sakkalis, M. Pediaditis, and M. Tsiknakis, "Methods for Seizure Detection and Prediction: An Overview", in *Modern Electroencephalographic Assessment Techniques: Theory and Applications*, pp. 131–157, V. Sakkalis, Ed. New York, NY: Springer, 2015.
- [9] E. Bou Assi, D.K. Nguyen, S. Rihana, and M. Sawan, "Towards accurate prediction of epileptic seizures: A review", *Biomed. Signal Process. Control* 34, 144–157 (2017), doi: 10.1016/j. bspc.2017.02.001.

- [10] G. Giannakakis, M. Tsiknakis, and P. Vorgia, "Focal epileptic seizures anticipation based on patterns of heart rate variability parameters", *Computer Methods and Programs in Biomedicine* 178, 123–133 (2019), doi: 10.1016/j.cmpb.2019.05.032.
- [11] M. Kotas, "Projective filtering of time-aligned beats for foetal ECG extraction", Bull. Pol. Acad. Sci. Tech. Sci. 55(4), 331–339 2007.
- [12] K. Lewenstein, M. Jamroży, and T. Leyko, "The use of recurrence plots and beat recordings in chronic heart failure detection", *Bull. Pol. Acad. Sci. Tech. Sci.* 64(2), 339–345 (2016).
- [13] J. Jarczewski, A. Furgała, A. Winiarska, M. Kaczmarczyk, and A. Poniatowski, "Cardiovascular response to different types of acute stress stimulations", *Folia Medica Cracoviensia* 59(4), 95–110 (2019).
- [14] J. Jeppesen, S. Beniczky, P. Johansen, P. Sidenius, and A. Fuglsang-Frederiksen, "Comparing maximum autonomic activity of psychogenic non-epileptic seizures and epileptic seizures using heart rate variability", *Seizure* 37, 13–19 (2016), doi: 10.1016/j. seizure.2016.02.005.
- [15] J. Jeppesen, S. Beniczky, P. Johansen, P. Sidenius, and A. Fuglsang-Frederiksen, "Detection of epileptic seizures with a modified heart rate variability algorithm based on Lorenz plot", *Seizure* 24, 1–7 (2015), doi: 10.1016/j.seizure.2014.11.004.
- [16] J. Jeppesen, S. Beniczky, P. Johansen, P. Sidenius, and A. Fuglsang-Frederiksen, "Using Lorenz plot and Cardiac Sympathetic Index of heart rate variability for detecting seizures for patients with epilepsy", in 2014 36th Annual International Conference of the IEEE Engineering in Medicine and Biology Society, 2014, pp. 4563–4566, doi: 10.1109/EMBC.2014.6944639.
- [17] F. Fürbass, S. Kampusch, E. Kaniusas, J. Koren, S. Pirker, R. Hopfengärtner, H. Stefan, T. Kluge, C. Baumgartner, "Automatic multimodal detection for long-term seizure documentation in epilepsy", *Clinical Neurophysiology* 128(8), 1466–1472 (2017), doi: 10.1016/j.clinph.2017.05.013.
- [18] M. Toichi, T. Sugiura, T. Murai, and A. Sengoku, "A new method of assessing cardiac autonomic function and its comparison with spectral analysis and coefficient of variation of R-R interval", *J. Auton. Nerv. Syst.* 62(1–2), 79–84 (1997), doi: 10.1016/s0165-1838(96)00112-9.
- [19] J. Pan and W.J. Tompkins, "A Real-Time QRS Detection Algorithm", *IEEE Transactions on Biomedical Engineering* BME-32(3), 230–236 (1985), doi: 10.1109/TBME.1985.325532.
- [20] S. Rezaei, S. Moharreri, S. Ghiasi, and S. Parvaneh, "Diagnosis of sleep apnea by evaluating points distribution in poincare plot of RR intervals", in 2017 Computing in Cardiology (CinC), 2017, pp. 1–4, doi: 10.22489/CinC.2017.158-398.
- [21] S. Kiranyaz, O. Avci, O. Abdeljaber, T. Ince, M. Gabbouj, and D. J. Inman, "1D Convolutional Neural Networks and Applications: A Survey", arXiv:1905.03554 [cs, eess], May 2019, Accessed: Jul. 10, 2020. [Online]. Available: http://arxiv.org/ abs/1905.03554.
- [22] T. Poggio and Q. Liao, "Theory I: Deep networks and the curse of dimensionality", Bull. Pol. Acad. Sci. Tech. Sci. 66(6), 761–773 (2018).
- [23] J. Kurek, B. Świderski, S. Osowski, M. Kruk, and W. Barhoumi, "Deep learning versus classical neural approach to mammogram recognition", *Bull. Pol. Acad. Sci. Tech. Sci.* Vol. 66(6), 831–840 2018, doi: 10.24425/bpas.2018.125930.
- [24] Y. Zhang and Z. Wang, "Research on intelligent algorithm for detecting ECG R waves", in 2015 IEEE 5th International Conference on Electronics Information and Emergency Communication, 2015, pp. 47–50, doi: 10.1109/ICEIEC.2015.7284484.

- [25] M. Kołodziej, A. Majkowski, P. Tarnowski, R.J. Rak, and A. Rysz, "Implementation of 1DConvolutional Neural Network for Cardiac Sympathetic Index Estimation", presented at the 2020 IEEE 21st International Conference on Computational Problems of Electrical Engineering (CPEE), 2020.
- [26] K.M. Gaikwad and M.S. Chavan, "Removal of high frequency noise from ECG signal using digital IIR butterworth filter", in 2014 IEEE Global Conference on Wireless Computing Networking (GCWCN), 2014, pp. 121–124, doi: 10.1109/GCWCN.2014.7030861.
- [27] M. Kołodziej, A. Majkowski, and R.J. Rak, "A new method of feature extraction from EEG signal for brain-computer interface design", *Prz. Elektrotechniczny* 9, 35–38 (2010).
- [28] K. Hayase, K. Hayashi, and T. Sawa, "Hierarchical Poincaré analysis for anaesthesia monitoring", J. Clin. Monit. Comput. 34, 1321–1330 (2020), doi: 10.1007/s10877-019-00447-0.
- [29] J. Niehoff, M. Matzkies, F. Nguemo, J. Hescheler, and M. Reppel, "The Effect of Antiarrhythmic Drugs on the Beat Rate Variability of Human Embryonic and Human Induced Pluripotent Stem Cell Derived Cardiomyocytes", Sci. Rep. 9(1), 14106 (2019), doi: 10.1038/s41598-019-50557-7.
- [30] M.M. Platiša, T. Bojić, S. Mazić, and A. Kalauzi, "Generalized Poincaré plots analysis of heart period dynamics in different physiological conditions: Trained vs. untrained men", *PLoS ONE* 14(7), e0219281 (2019), doi: 10.1371/journal.pone.0219281.
- [31] P. Fontana, N.R.A. Martins, M. Camenzind, M. Boesch, F. Baty, O.D. Schoch, M.H. Brutsche, R.M. Rossi, and S. Annaheim, "Applicability of a Textile ECG-Belt for Unattended Sleep Apnoea Monitoring in a Home Setting", *Sensors (Basel)* 19(15), 3367 (2019), doi: 10.3390/s19153367.
- [32] T. Schmidt, S. Wulff, K.-M. Braumann, and R. Reer, "Determination of the Maximal Lactate Steady State by HRV in Overweight and Obese Subjects", *Sports Med. Int. Open* 3(2), E58–E64 (2019), doi: 10.1055/a-0883-5473.

- [33] J. Piskorski and P. Guzik, "Geometry of the Poincaré plot of RR intervals and its asymmetry in healthy adults", *Physiol. Meas*. 28(3), 287–300 (2007), doi: 10.1088/0967-3334/28/3/005.
- [34] Ö. Yıldırım, P. Pławiak, R.-S. Tan, and U. R. Acharya, "Arrhythmia detection using deep convolutional neural network with long duration ECG signals", *Computers in Biology and Medicine* 102, 411–420, (2018), doi: 10.1016/j.compbiomed.2018.09.009.
- [35] B. Zhao, H. Lu, S. Chen, J. Liu, and D. Wu, "Convolutional neural networks for time series classification", *Journal of Sys*tems Engineering and Electronics 28(1), 162–169 (2017), doi: 10.21629/JSEE.2017.01.18.
- [36] V. Lebedev and V. Lempitsky, "Speeding-up convolutional neural networks: A survey", Bull. Pol. Acad. Sci. Tech. Sci. 66(6), 799–810 (2018), doi: 10.24425/BPAS.2018.125927.
- [37] M. Grochowski, A. Kwasigroch, and A. Mikołajczyk, "Selected technical issues of deep neural networks for image classification purposes", *Bull. Pol. Acad. Sci. Tech. Sci.* 67(2), 363–376 (2019).
- [38] X. Glorot, A. Bordes, and Y. Bengio, "Deep Sparse Rectifier Neural Networks", in *Proceedings of the Fourteenth Interna*tional Conference on Artificial Intelligence and Statistics, 2011, 315–323, [Online]. Available: http://proceedings.mlr.press/v15/ glorot11a.html.
- [39] A. Krizhevsky, I. Sutskever, and G.E. Hinton, "ImageNet classification with deep convolutional neural networks", *Commun. ACM* 60(6), 84–90 (2017), doi: 10.1145/3065386.
- [40] Y. Gal and Z. Ghahramani, "A Theoretically Grounded Application of Dropout in Recurrent Neural Networks", in *Advances in Neural Information Processing Systems* 29, pp.1019–1027, Eds. D.D. Lee, M. Sugiyama, U.V. Luxburg, I. Guyon, and R. Garnett, Curran Associates, Inc., 2016.
- [41] S. Albawi, T.A. Mohammed, and S. Al-Zawi, "Understanding of a convolutional neural network", in 2017 International Conference on Engineering and Technology (ICET), 2017, pp. 1–6, doi: 10.1109/ICEngTechnol.2017.8308186.