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Supervised and Unsupervised Deep Learning Approaches for EEG Seizure Prediction

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Abstract

Epilepsy affects more than **50** million people worldwide, making it one of the world's most prevalent neurological diseases. The main symptom of epilepsy is seizures, which occur abruptly and can cause serious injury or death. The ability to predict the occurrence of an epileptic seizure could alleviate many risks and stresses people with epilepsy face. Most of the previous work is focused at seizure detection, we pivot our focus to seizure prediction problem. We formulate the problem of detecting preictal (or pre-seizure) with reference to normal EEG as a precursor to incoming seizure. To this end, we developed several supervised deep learning approaches model to identify preictal EEG from normal EEG. We further develop novel unsupervised deep learning approaches to train the models on only normal EEG, and detecting pre-seizure EEG as an anomalous event. These deep learning models were trained and evaluated on two large EEG seizure datasets in a person-specific manner. We found that both supervised and unsupervised approaches are feasible; however, their performance varies depending on the patient, approach and architecture. This new line of research has the potential to develop therapeutic interventions and save human lives.

Keywords: deep learning, intracranial EEG, seizure prediction, signal processing

1 Introduction

Epilepsy is one of the most prevalent neurological disorders in the world, affecting approximately 1% of the world’s population [1–3]. Epilepsy is characterized by spontaneously occurring seizures, which could lead to bodily injuries, fractures, burns [4], and death in many cases [5]. People with epilepsy are mostly concerned with the fear of incoming seizures [6]. Therefore, there is a dire need to reduce the unpredictability of seizures to reduce the risk of injuries and improve their quality of life.

Electroencephalography (EEG) is normally used to analyze brain activity pertaining to seizures [7]. Brain activity in people with epilepsy can be separated into four states: regular brain activity (interictal), brain activity before the seizure (preictal), brain activity during the seizure (ictal), and brain activity immediately after a seizure (postictal). The preictal state can contain observable physiological changes prior to the onset of a seizure [8] that can be used to predict an incoming seizure. The capability to predict an epileptic seizure could alleviate the risks patients face [9]; it would give patients the time to get help and greatly reduce the risk of injury. However, the biggest challenge in designing seizure prediction approaches is that there is no universally agreed upon preictal period length (PPL). Bandarabadi et al. [10] investigated the optimal PPL for seizure prediction using statistical analysis and found that the optimal PPL varies for each patient and for seizure within each patient [10].

Most of the work in this area is around seizure detection [11], which involves detecting a seizure after its occurrence. Although this is important, contemporary work must aim to predict seizures before their onset as it can save patients’ lives and improve their quality of life. Our main hypothesis is that the correct detection of preictal state against normal brain activity (through supervised or unsupervised approaches) can be a strong indicator of an incoming epileptic seizure. In the supervised setting, a binary classifier can be trained between interictal and preictal periods. Whereas, in the unsupervised setting, a classifier can be trained on only normal EEG (interictal) and preictal state can be identified as an anomaly. Our main contributions are:

- Presented supervised and new unsupervised deep learning approaches to predict epileptic seizures.
- Experimentally determined the PPL and window size, as against heuristics or domain knowledge.
- Performed leave-one-seizure out cross validation for better generalization of results.
- All the experiments were performed in a patient-specific manner to avoid data leakages, overestimation of results and emphasis on individualized outcomes.

Our results showed that the unsupervised approaches were able to obtain comparable results to supervised seizure prediction in many patients. However, across all implementations there was not one clear best-performing model. This paper is an extension of our preliminary work [12] that introduced supervised convolution neural network (CNN) on SWEC-ETHZ dataset [13]. In this paper, we present two new supervised approaches: CNN-Long Short Term Memory (LSTM) and Temporal Convolution Network (TCN), and three new unsupervised approaches (CNN, CNN-LSTM, TCN autoencoders). We developed new seizure prediction baselines for the SWEC-ETHZ dataset [13] and included a new CHB-MIT dataset [14].

2 Related Work

Seizure prediction using supervised machine learning has been used to distinguish the interictal and preictal states [15]. Typical supervised machine learning seizure prediction approaches involve signal pre-processing, extracting and selecting features, followed a classifier [15]. Common signal processing techniques include high-pass, low-pass, or band-pass filtering, as well as artifact removal techniques [15]. Feature extraction is typically done by a bio-signals or epilepsy expert examining a patient’s EEG and deciding appropriate features for separating the preictal and interictal states [15]. These features are often patient-specific and include statistical, non-linear, frequency domain, and time-frequency domain features [15, 16]. Common classifier choices include support vector machines (SVM), k-nearest neighbour and random forest [15].

Machine learning approaches may have limitations in terms of extracting hand-crafted features, which could be sub-optimal and time consuming. Deep learning approaches can overcome some of these challenges by able to learn features from data with little to no pre-processing, generate high-level representations of data, and learn complex functions [17]. An overview of preictal-interictal classification seizure prediction methods (on human subjects) using deep learning is shown in table 1.

Many reviewed deep learning methods performed some type of pre-process the EEG before passing it on to the classifier, typically through filtering [1, 21], artifact removal [35], or time-frequency analysis [21, 22]. Common deep learning architectures used for seizure prediction include CNN [1, 22], LSTM network [29, 32], and feed-forward multilayer perceptron (MLP) [18]. We observed that majority of the studies use CNNs, LSTMs and/or their combinations to benefit from learning spatial and temporal features. The window size (fixed duration data to analyze) and PPL were kept fixed in most of the studies, and they varied even when working on the same dataset and patients. This is an issue in building classifiers to predict seizures because the optimal PPL varies across patients (as concluded by Bandarabadi et al. [10]). Only four of the studies reported experimenting with the PPL [18, 21, 29, 35], while others did not present any rationale for their choice. Some of the studies (e.g., [14]) also found different PPLs sizes, showing that the optimal PPL varies depending on the method’s implementation. These studies show that it is better to determine the PPL empirically at a patient-specific level, rather than using a generic or pre-determined average over a population. We extend the existing supervised methods by obtaining PPL and window size using a leave-one-seizure-out (LOSO) evaluation and introduced new supervised TCN classifier for this task. There is no known work on using unsupervised deep learning for seizure prediction using EEG. For the first time, we introduced three different autoencoder models and studied their performance for this problem.

3 Supervised Seizure Prediction

Preictal-interictal classification for seizure prediction is performed with three different architectures: convolutional neural networks (CNN) (used in our previous work [12]), and two new architectures, i.e., CNN-LSTM, and TCN. We briefly discuss them below.

Citation	Window Size (Seconds)	PPL (Minutes)	Pre-processing	DL Architecture
[18]	5	10,20,30,40	Various fixed features	MLP
[19]	10	Unknown	Fractional FT	MLP
[20]	300	50	Handcrafted	CNN
[21]	1	10	CWT	CNN
[22]	30	60	STFT	CNN
[23]	15	60	None	CNN
[24]	5	30	Common spatial pattern	CNN
[25]	30	60	Fast FT	Multi-view CNN
[26]	10	60	MFCC	CNN
[27]	1,2,3,8	10	None	CNN
[12]	5,10,15,30,60	30,60,120	STFT	CNN
[28]	20	30,60	None	CNN
[29]	5	15	Handcrafted	LSTM
[1]	10	30	STFT	CNN + LSTM
[30]	4	60	None	CNN-AE+BiLSTM
[31]	5	60	None	MLP,CNN,CNN+LSTM
[32]	10	30	Image conversion	CNN+LSTM
[33]	30	60	STFT	CNN+SVM
[34]	30	30	None	1DCNN+GRU
[35]	Unknown	60	Image conversion	3DCNN
[36]	4	30	Various fixed features	3DCNN
[37]	28	30	STFT	Convolutional GAN

Table 1: Overview of deep learning EEG seizure prediction methods.

3.1 CNN

The CNN model takes in EEG samples that have been time-frequency transformed using a STFT [22] (see Section 5.3). This helps the model in extracting time and frequency features and puts the data into a suitable format for 2D convolutions [22]. The CNN architecture takes advantage of spatial information in data to learn relevant features. Each sample was converted into a 2D matrix $F \times T$ using a STFT, where F was the number of sample frequencies used and T was the number of segment times used. The matrix was then resized to a 128×128 “image” using bilinear interpolation so that image sizes were consistent regardless of the window size. The time-frequency transform was done independently for each channel, resulting in each sample being of dimensions $C \times 128 \times 128$, where C is the total number of channels. The samples were then passed to the CNN model, which is made up of three convolutional blocks (see Figures 1 and 2), followed by three fully connected layers with ReLU activation functions. Table 2 shows the model hyperparameters used for the CNN.

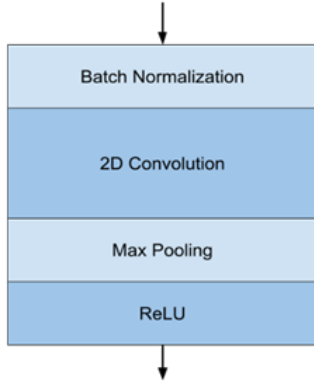
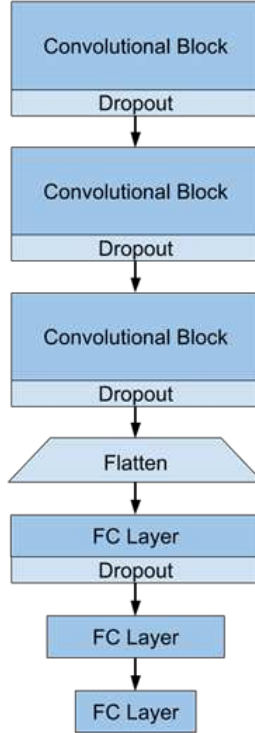


Fig. 1: Convolutional block.



CNN kernel size	5
CNN filter sizes	8, 16, 32
Fully connected sizes	128, 64
Dropout	0.5

Table 2: CNN model hyperparameters.

Fig. 2: CNN architecture with convolution blocks and fully connected (FC) layers.

3.2 CNN-LSTM

The CNN-LSTM architecture takes advantage of both the spatial feature extraction of the CNN along with the LSTM’s propensity to work well with temporal data. The CNN-LSTM model takes in STFT images similar to the CNN model. The input is a consecutive series of images as one sample. The input sequence is divided into smaller sub-sequences, which are independently time-frequency transformed and resized into 64×64 images, leading to a dimensions $C \times n \times 64 \times 64$, where n is the number of sub-sequences in a sample and is equal to the sequence length divided by the sub-sequence length. Each sub-window is passed into a CNN model with two convolutional blocks that outputs a feature vector. Then, each feature vector is concatenated into a sequence and passed into a 2-layer LSTM, whose outputs are passed to a fully connected layer that outputs the final scores. An overview of the CNN-LSTM architecture and hyperparameters are shown in Figure 3 and Table 3a.

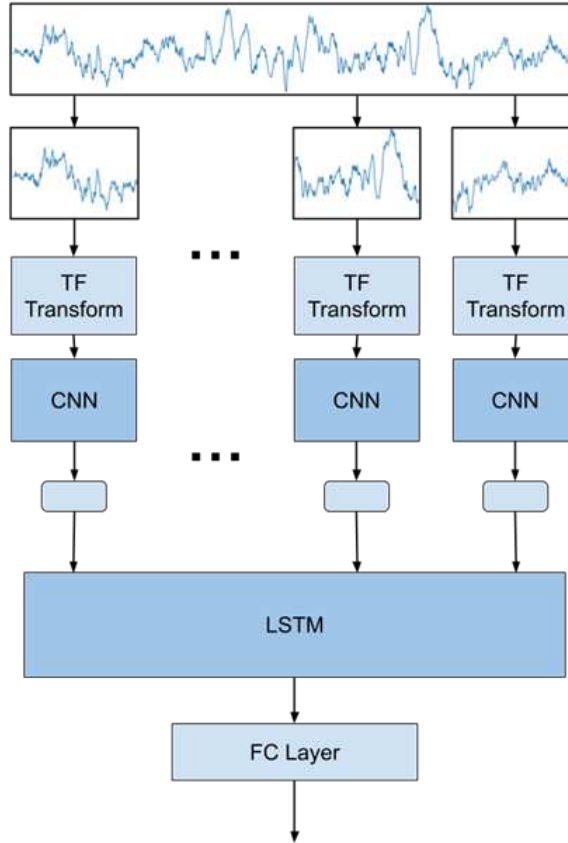


Fig. 3: CNN-LSTM architecture showing time-frequency (TF) transform, CNN layers, LSTM and fully connected (FC) layer.

3.3 TCN

The TCN model takes in scaled sequences of size $C \times S/4$, where S is the sequence length and the sequences were down-sampled by a factor of 4. The TCN model [38] consisted of TCN blocks (see Figure ??). Each TCN block is two consecutive sub-blocks that contain a causal 1D convolution layer with a dilation, a weight normalization layer, a ReLU activation function, and a dropout layer [38]. The TCN blocks have skip connections, where the input to the block is added to the output [38]. The model contained 6 TCN blocks with 32 channels each, followed by a 1D convolution layer, and a fully connected layer. The dilation factor of each block was $2^{(n-1)}$, where n is the layer number. Figure 4 and Table 3b shows the TCN architecture and hyperparameters.

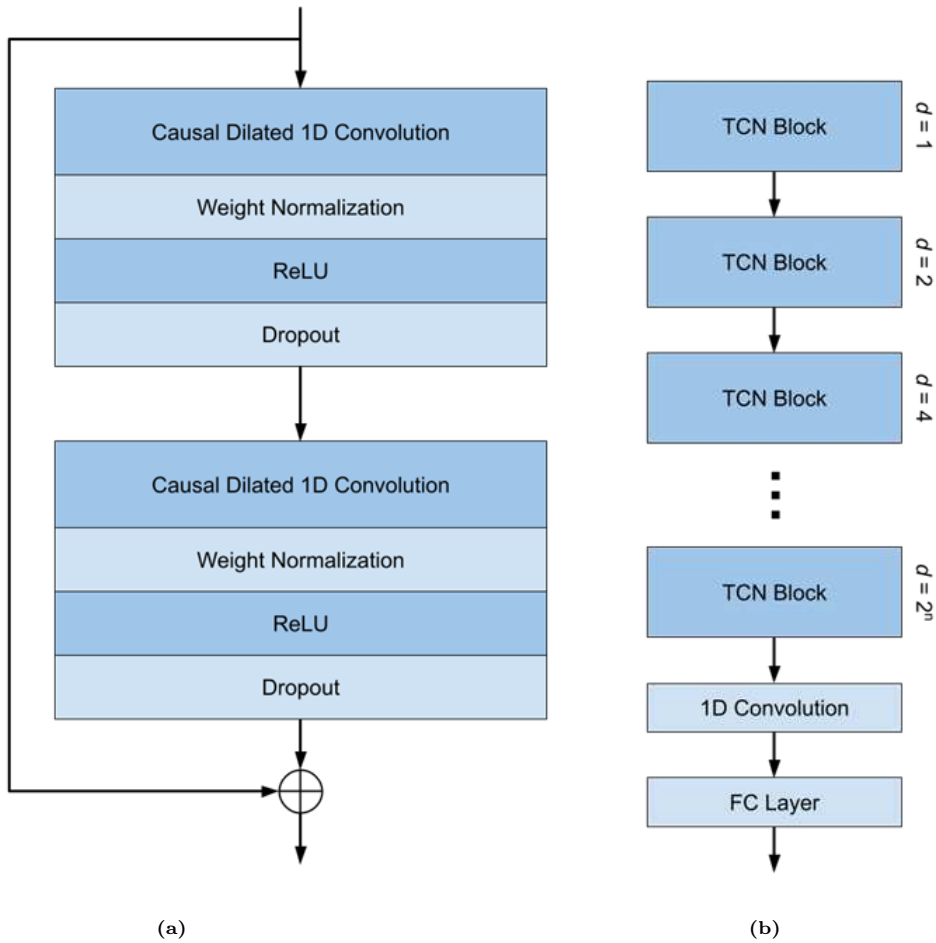


Fig. 4: (a) TCN block description. TCN: temporal convolutional network. ReLU: rectified linear unit. (b) Supervised TCN architecture overview. TCN: temporal convolutional network. FC layer: fully connected layer.

CNN kernel size	5
CNN filter sizes	8, 16
LSTM feature vector size	32
LSTM hidden size	16
Fully connected size	96
Dropout	0.5

(a)

TCN kernel size	5
TCN filter sizes	32, 32, 32, 32, 32, 32
Fully connected size	64
Dropout	0.2

(b)

Table 3: Hyperparameters of (a) CNN-LSTM, and (b) TCN models

4 Unsupervised Seizure Prediction

The reliance on preictal data for supervised seizure prediction methods remains a challenge. Preictal data is typically scarce, and deep learning methods require a considerable amount of data from both classes to work well. Preictal-interictal classification methods cannot be used effectively on patients with little preictal data, and class imbalance still remains an impending problem. An unsupervised approach (anomaly detection) to seizure prediction could remedy these problems. Anomaly detection for seizure prediction would require only interictal (and no preictal data) to train, making it easier to be more accessible to a larger population. Autoencoders (AEs) and its variants are apt to be used within this framework, with reconstruction error used as an anomaly score. To our knowledge, this is the first seizure prediction work that uses unsupervised deep learning approach for epileptic seizure prediction. We implemented the following autoencoder approaches for this task.

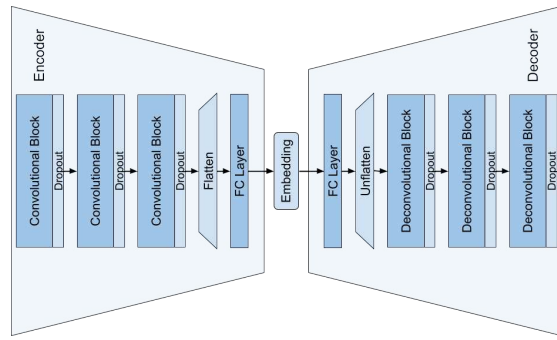
- CNN autoencoder [39]. Similar to the supervised CNN, it takes STFT images as input. The encoder is made up of three convolutional blocks followed by a fully connected layer which generates an embedding state of size 64. The decoder is a mirrored version of the encoder (see Figure 5a).
- CNN-LSTM autoencoder [40]. Similar to the supervised CNN-LSTM, the input sequence was divided into smaller sub-sequences and then an STFT was performed on each sub-sequence. The encoder consisted of an individual CNN encoder for each sub-sequence followed by an LSTM that generated an embedding state of size 64. The decoder has the reverse architecture to the encoder. (see Figure 5b).
- TCN autoencoder [41]. It takes in raw scaled sequences as is the case with the supervised TCN. The encoder was a TCN with three layers, each with 16 channels followed by a 1d convolution and a fully connected layer. The size of the embedding state was 64. The decoder was an exact mirror of the encoder (see Figure 5c).

5 Data Processing

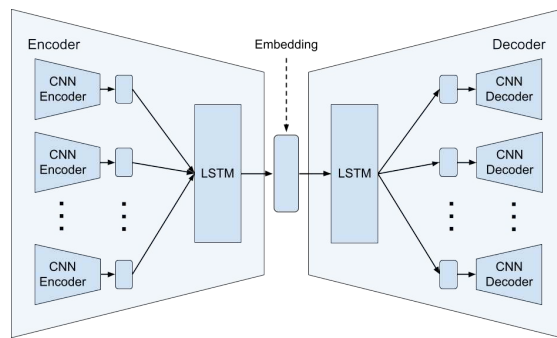
5.1 Datasets

We used two EEG Epilepsy seizure datasets, the Sleep-Wake Epilepsy Centre ETH Zurich (SWEC-ETHZ) dataset [13] and the Children’s Hospital Boston Massachusetts Institute of Technology (CHB-MIT) dataset [14]. Both datasets are publicly available, easy to access, and contain human raw EEG recordings, where no seizure states have been pre-selected. This is important so we can define and experiment with different preictal and interictal regions. The SWEC-ETHZ dataset is an iEEG dataset containing over 2,500 hours of recordings across 18 patients with a sampling rate of either 512Hz or 1024Hz [13]. The CHB-MIT dataset contains scalp EEG recordings from 22 patients sampled at 256Hz with at least 22 EEG electrodes [14].

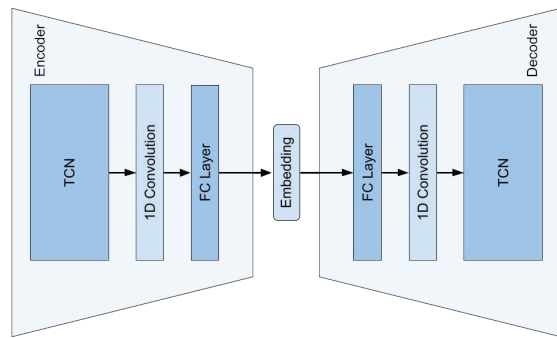
We define a “lead seizure“ as any seizure that occurs at least 30 minutes after a preceding seizure [22]. Only preictal periods from lead seizures were considered because of the lack of interictal and preictal data to train models. Patients that have less than three lead seizures were withheld from the experiments because at least three lead



(a)



(b)



(c)

Fig. 5: Autoencoders: (a) CNN, (b) CNN-LSTM, (c) TCN

seizures were required to perform test partitioning combined with an internal leave-one-seizure-out (LOSO) cross-validation step (see Figure 6b). Six out of the 18 patients in the SWEC-ETHZ dataset were not considered for this work due to this condition. All patients in the CHB-MIT dataset had at least three lead seizures. A description of dataset attributes for all patients used from both the SWEC-ETHZ and CHB-MIT datasets is shown in Tables 4 and 5, respectively.

Patient ID	Hours of data	Seizures	Lead seizures	Electrodes
ID03	158	4	4	64
ID04	41	14	14	32
ID05	110	4	4	128
ID06	146	8	8	32
ID07	69	4	4	75
ID08	144	4	4	61
ID09	41	23	14	48
ID10	42	17	15	32
ID12	191	9	9	56
ID13	104	7	7	64
ID16	177	5	5	34
ID18	205	5	5	42

Table 4: SWEC-ETHZ dataset patient description [13]

5.2 Data Preprocessing

The length and location of the preictal period is defined by the PPL and the intervention time (IT). The IT is the time between the preictal state and the seizure onset. Interictal data is defined as any data that is not preictal, ictal, postictal, and is d distance away from the preictal state, as shown in Figure 6a. The data was divided into samples of a fixed window size, which were labelled as either interictal or preictal. We set $d = 0$ to evaluate the model’s ability to classify interictal and preictal samples in close temporal proximity to actual seizures. The IT was set to 0, increasing it can be a future experiment after generating a baseline. In the SWEC-ETHZ dataset, interictal samples were randomly selected with a down-sampling factor of 8 because interictal data was overly abundant and the classes were significantly imbalanced (patients ID04, ID09, and ID10 used a down-sampling factor of 2 instead because there was less interictal data). The number of preictal samples were artificially increased by using 50% overlapping windows. The size of each sample was $sf \times C$ where s was the window size, f was the sampling rate, and C was the number of EEG electrodes.

The dataset was partitioned into a training set and a testing set using LOSO partitioning. We used the last lead seizure’s preictal data as the test set, while all other preictal data was part of the training set. As shown in Figure 6b, LOSO partitioning is a better way to evaluate a model’s ability to generalize to a new seizure’s preictal

Patient ID	Hours of data	Seizures	Lead seizures	Electrodes
1	41	7	7	22
2	35	3	3	22
3	38	7	7	22
4	156	4	3	22
5	39	5	5	22
6	67	10	7	22
7	67	3	3	22
8	20	5	5	22
9	68	4	3	22
10	50	7	7	22
11	35	3	3	22
12	24	40	11	22
13	33	12	7	22
14	26	8	6	22
15	40	20	14	22
16	19	10	5	22
17	21	3	3	22
18	36	6	5	22
19	30	3	3	22
20	28	8	6	22
21	33	4	4	22
22	31	3	3	22
23	27	7	3	22

Table 5: CHB-MIT dataset patient description [14]

data. Standard test partitioning where samples are randomly assigned to the training or test set may be an overestimation of the actual performance of the classifier.

5.3 Time-Frequency Transform

We transformed the EEG data from a time-series input into the time-frequency domain [42, 43] using short-time Fourier transform (STFT). It converts a one-dimensional time-series signal into a two-dimensional matrix of values with axes of time and frequency [44]. The STFT splits the signal into a series of smaller sequences and then performing Fourier transforms on each one individually, providing a way to see changes in the frequency domain at various points in time [45]. In CNN based models used in the work, an STFT was used to pre-process the input before passing samples to the model. Other time-frequency analysis methods such as the continuous wavelet transform [21] and phase-amplitude coupling [46] were experimented with in our preliminary work but did not provide better results.

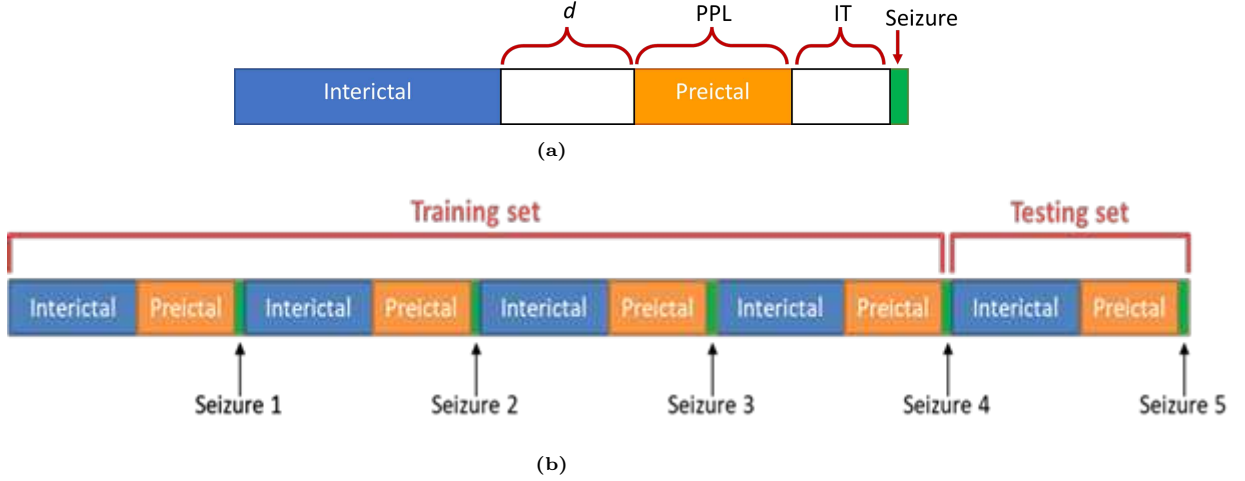


Fig. 6: (a) Labelling of the preictal and interictal periods with parameters. (b) Simplified visualization of LOSO test partitioning by withholding the last seizure.

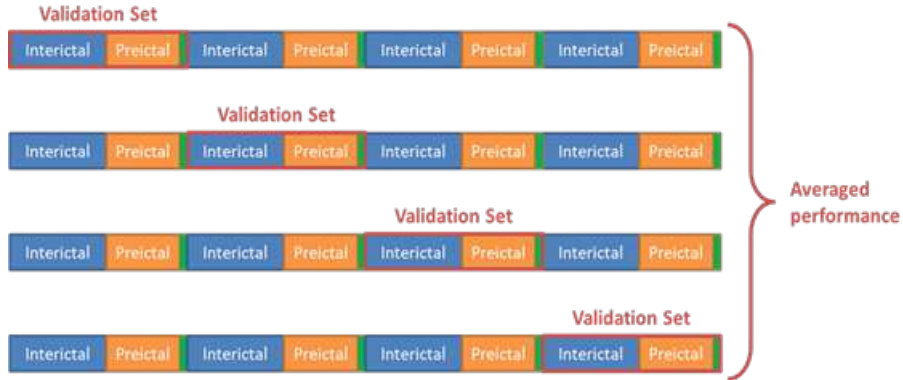


Fig. 7: LOSO cross-validation example with four seizures. One seizure is used for validation while the others are used for model training.

6 Experimental Setting and Results

A grid search was performed to find the optimal window size and PPL for each patient. We ran the model with varying window size (5, 10, 15, 30, 60 seconds) and PPL (30, 60, 120 minutes) values. We used an internal LOSO cross-validation to tune the parameters without looking at test data. This was done by dividing the training set into folds where each fold was a different seizure’s preictal and interictal data. One fold was the validation set while the others were used for training. Each fold in the set was used as the validation set once, and the performance across all runs in a patient was averaged. An example of the cross-validation method used is shown in Figure 7. The

area under the Receiver Operating Characteristic curve (AUC ROC) [47] was used as a performance metric for hyperparameter tuning. The test set was completely withheld from this process. All the models were trained using an NVIDIA V100S-PCIe GPU with 32GB memory. A class-weighted (class weights vary per patient) cross-entropy loss function was used with the Adam optimizer and was trained for 100 epochs with a batch size of 128 and a learning rate of 0.0001. All implementations were done in the PyTorch framework [48]. After the final parameters for a model were set, it was evaluated on the test set using the AUC ROC and precision-recall curve (AUC PR). AUC PR is more appropriate for imbalanced classification problems [49].

6.1 Supervised Prediction

Hyperparameter tuning results using the supervised CNN are shown in Tables 6 and 7 for the SWEC-ETHZ and CHB-MIT datasets, respectively. The window size and PPL obtained using cross-validations as well as AUC ROC vary considerably across different patients in both datasets. More than half of the patients in each dataset show AUC ROC values greater than 0.7. In the SWEC-ETHZ dataset, six of the patients had a test AUC ROC at least 0.1 lower than their validation AUC ROC while in the CHB-MIT dataset it was eight patients. This is consistent with Bandarabadi et al. [10] that the optimal preictal period for seizure prediction varies even on seizures within the same patient. The best way to account for this problem is to train and test on as many lead seizures’ preictal data as possible.

Patient ID	Window size	PPL	Validation AUC ROC	Test AUC ROC
ID03	30	1800	0.793	0.939
ID04	60	3600	0.708	0.509
ID05	60	7200	0.953	0.918
ID06	60	7200	0.704	0.948
ID07	30	7200	0.722	0.713
ID08	15	7200	0.722	0.454
ID09	30	7200	0.901	0.944
ID10	60	3600	0.807	0.574
ID12	60	1800	0.981	0.798
ID13	60	1800	0.721	0.499
ID16	10	1800	0.719	0.423
ID18	15	1800	0.832	0.850

Table 6: Validation and test results for preictal-interictal classification with optimized hyperparameters on the SWEC-ETHZ dataset.

6.1.1 Comparison with Fixed Parameters

We implemented a preictal-interictal classification model with a fixed window size of 30 seconds and PPL of 1 hour to compare to our tuned hyperparameter model. The model architecture is a CNN identical to the optimized parameter implementation.

Patient ID	Window size	PPL	Validation AUC ROC	Test AUC ROC
1	60	1800	0.987	0.997
2	5	7200	0.718	0.982
3	10	1800	0.853	1.000
4	5	7200	0.376	0.649
5	5	3600	0.821	0.828
6	10	3600	0.616	0.858
7	5	7200	0.744	0.133
8	10	1800	0.999	0.379
9	10	7200	0.789	0.788
10	5	1800	0.732	0.686
11	30	3600	0.890	0.978
12	60	7200	0.917	0.549
13	5	3600	0.973	0.898
14	5	1800	0.817	0.139
15	30	7200	0.824	0.470
16	60	1800	0.686	0.688
17	5	7200	0.933	0.565
18	60	1800	0.750	0.820
19	30	7200	1.000	0.966
20	5	1800	0.983	0.898
21	5	7200	0.807	0.721
22	5	7200	0.770	0.397
23	30	1800	1.000	0.614

Table 7: Validation and test results for preictal-interictal classification with optimized hyperparameters on the CHB-MIT dataset.

This was done to explore the benefits of optimizing hyperparameters for seizure prediction. Figures 8 show the comparison of the two methods on the SWEC-ETHZ and CHB-MIT dataset. In general for the SWEC-ETHZ dataset, the optimized hyperparameter implementation performed slightly better than the fixed parameter. In patient ID09, the optimized hyperparameter implementation performed much better than the fixed parameter implementation. For patient ID09, the hyperparameter tuning found a window size of 30 seconds and a PPL of 2 hours. It is likely that there was additional preictal information in the extra hour of data not used in the fixed parameter implementation. For the CHB-MIT dataset, most patients had similar results for both the fixed and optimized hyperparameter implementations. There were a few patients (ID 5, 16, 17, 18) that had much better results with the optimized model. However, there were also patients (ID 9, 22, 23) who performed better with a fixed hyperparameter implementation. For these patients, the last seizure’s optimal hyperparameters were likely different from the optimal hyperparameters for the preceding seizures in the patient’s dataset. Figures 10 and 11 show the comparison between the optimized and fixed hyperparameter implementations for the SWEC-ETHZ and CHB-MIT datasets

respectively using AUC PR instead. It can be observed that the optimized implementation generally performs better on the SWEC-ETHZ dataset in both metrics and that the difference is marginal in the CHB-MIT dataset. These experiments indicate that hyperparameter tuning can potentially improve the performance in comparison to fixed parameters.

Comparison of Preictal-Interictal Classification With Optimized vs Fixed Parameters on the SWEC Database

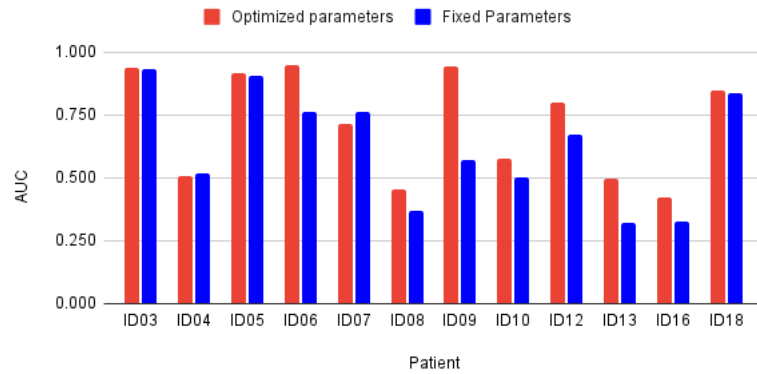


Fig. 8: AUC ROC Comparison of CNN models using optimized hyperparameters vs fixed hyperparameters on the SWEC-ETHZ dataset.

Comparison of Preictal-Interictal Classification With Optimized vs Fixed Parameters on the CHB-MIT Database

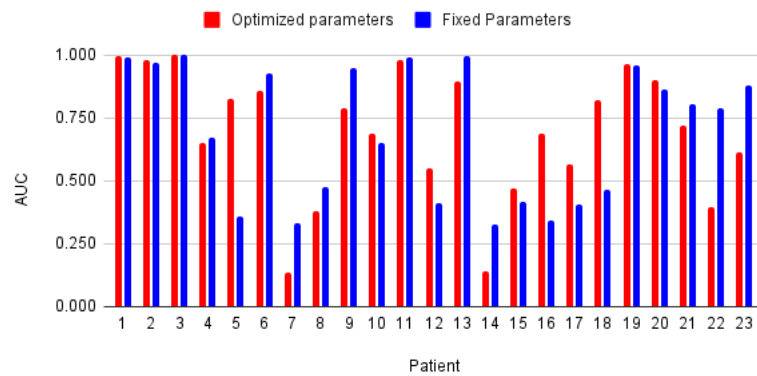


Fig. 9: AUC ROC Comparison of CNN models using optimized hyperparameters vs fixed hyperparameters on the CHB-MIT dataset.

Comparison of Preictal-Interictal Classification With Optimized vs Fixed Parameters on the SWEC Database

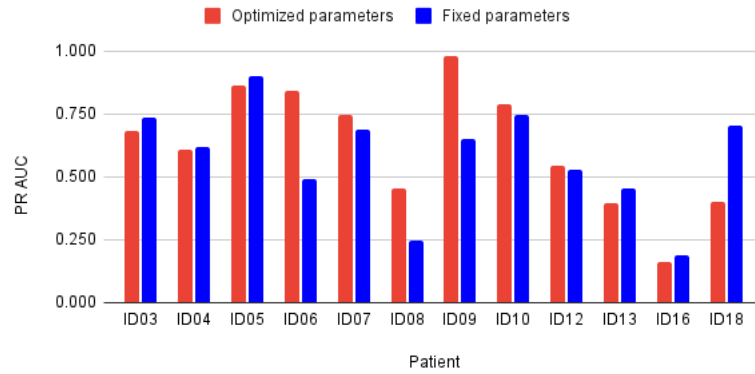


Fig. 10: PR AUC Comparison of CNN models using optimized hyperparameters vs fixed hyperparameters on the SWEC-ETHZ dataset.

Comparison of Preictal-Interictal Classification With Optimized vs Fixed Parameters on the CHB-MIT Database

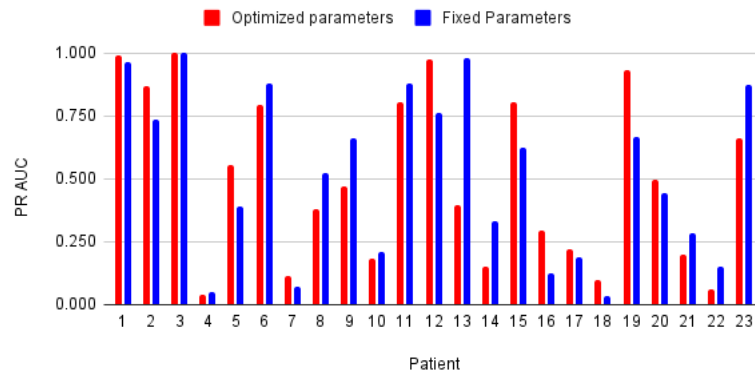


Fig. 11: PR AUC Comparison of CNN models using optimized hyperparameters vs fixed hyperparameters on the CHB-MIT dataset.

6.1.2 Comparison with Other Architectures

Using a fixed hyperparameter implementation, CNN-LSTM and TCN models were trained using a window size of 30 seconds and a PPL of 1 hour. A comparison of the AUC PR for all models is shown in Figures 12 and 13. In the SWEC-ETHZ dataset,

the CNN, CNN-LSTM and TCN were the best performing model for 3, 5 and 4 patients. The CNN and CNN-LSTM performed comparably well, and in each patient, the results were fairly similar. The TCN results was more variable, performing well on some patients that the other models performed poorly. In the CHB-MIT dataset, the CNN, CNN-LSTM and TCN were the best performing model for 7, 6 and 10 patients. The TCN model performed much better in the CHB-MIT dataset compared to the SWEC-ETHZ dataset. Overall, the CHB-MIT results were very variable with PR AUC values, varying considerably even within the same patient.

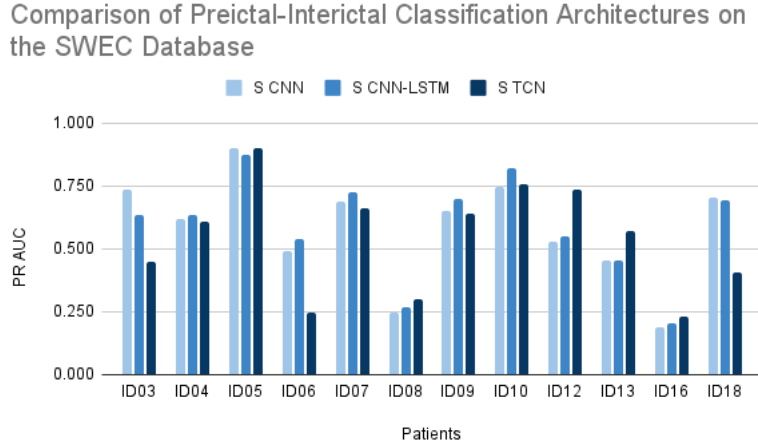


Fig. 12: Comparison of CNN, CNN-LSTM, and TCN implementations for preictal-interictal classification on the SWEC-ETHZ dataset.

6.2 Unsupervised Prediction

In the unsupervised approach, the training set only contained interictal data. For these experiments, the hyperparameters were fixed, with a window size of 30 seconds and a PPL of 60 minutes. The models were trained for 500 epochs with a batch size of 128 and a learning rate of 0.0005. After training, the models were evaluated on the test set that contained both interictal and preictal samples. Both AUC ROC and PR AUC were used to evaluate performance.

6.2.1 Comparison of Architectures

Figures 14, 15a, and 15b show the anomaly detection seizure prediction AUC PR results for the CNN, CNN-LSTM, and TCN AEs on the SWEC-ETHZ dataset and CHB-MIT dataset, respectively. We also show the supervised CNN with fixed hyperparameters for comparison. It can be observed that the performance varies significantly across different architectures and patients. For the SWEC-ETHZ dataset, the CNN AE

Comparison of Preictal-Interictal Classification Architectures on the CHB-MIT Database

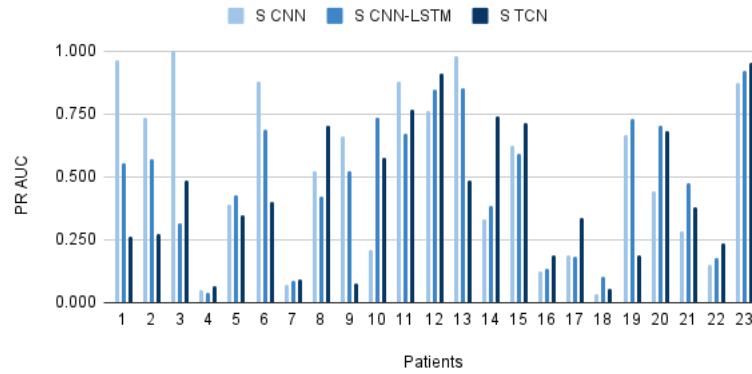


Fig. 13: Comparison of CNN, CNN-LSTM, and TCN implementations for preictal-interictal classification on the CHB-MIT dataset.

performed the worst across most patients while the CNN-LSTM and TCN AEs performed relatively better, and even surpassed the supervised implementation in some patients. In the CHB-MIT dataset, the results vary even more with no clear winner.

Comparison of Anomaly Detection Early Fusion Architectures on the SWEC-ETHZ Database

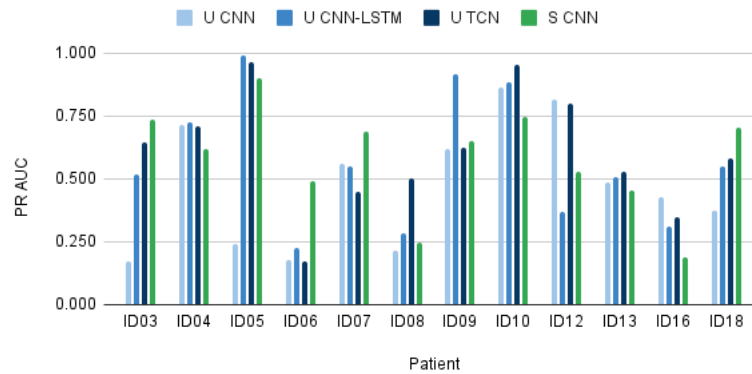
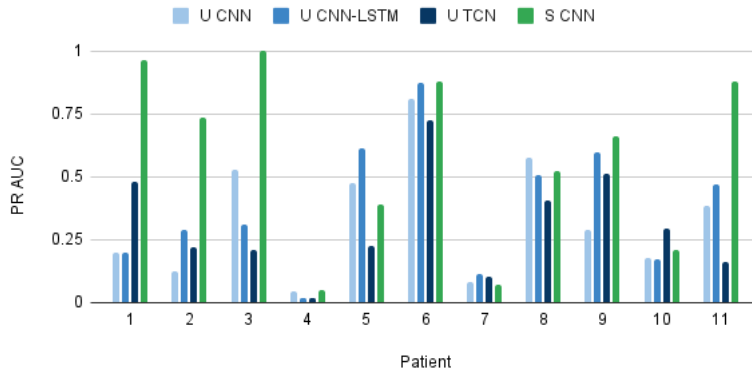


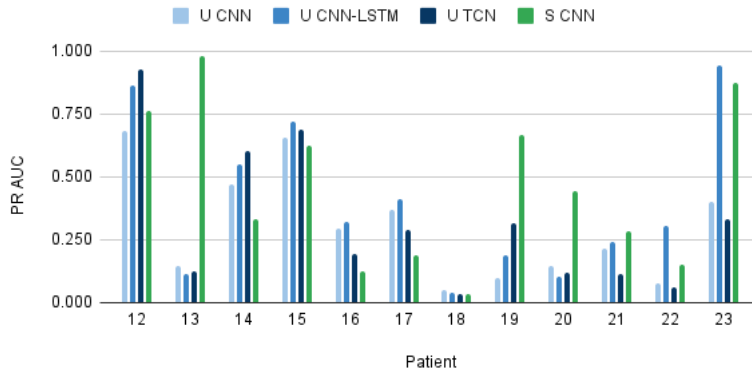
Fig. 14: Comparison of unsupervised seizure prediction using different model architectures on the SWEC-ETHZ dataset. U: unsupervised, S: supervised.

Comparison of Anomaly Detection Early Fusion Architectures on the CHB-MIT Database (Patient 1 to 11)



(a)

Comparison of Anomaly Detection Early Fusion Architectures on the CHB-MIT Database (Patient 12 to 23)



(b)

Fig. 15: Comparison of unsupervised seizure prediction using different model architectures on the CHB-MIT dataset (a) patients 1 to 11, (b) patients 12 to 23. U: unsupervised, S: supervised.

6.3 Best Implementations

Tables 8 and 9 show the best-performing implementation (from all experiments with supervised and unsupervised approaches) for each patient in the SWEC-ETHZ and CHB-MIT datasets and its corresponding AUC PR. For SWEC-ETHZ dataset, an unsupervised approach was the best-performing implementation for 7 out of 12 patients. For the CHB-MIT dataset, for 16 out of 23 patients, supervised approaches

performed better. In particular, the supervised CNN performed the best for 8 patients – the most of any model. Figure 16 shows that using the CNN-LSTM was the most effective for the most patients with best performance in 16 of the 35 patients.

Patient ID	Best Implementation	PR AUC
ID03	Supervised CNN	0.735
ID04	Unsupervised CNN-LSTM	0.727
ID05	Unsupervised CNN-LSTM	0.991
ID06	Supervised CNN-LSTM	0.537
ID07	Supervised CNN-LSTM	0.726
ID08	Unsupervised TCN	0.501
ID09	Unsupervised CNN-LSTM	0.918
ID10	Unsupervised TCN	0.955
ID12	Unsupervised CNN	0.817
ID13	Supervised TCN	0.573
ID16	Unsupervised CNN	0.430
ID18	Supervised CNN	0.702

Table 8: Best-performing implementation for each patient in the SWEC-ETHZ dataset.

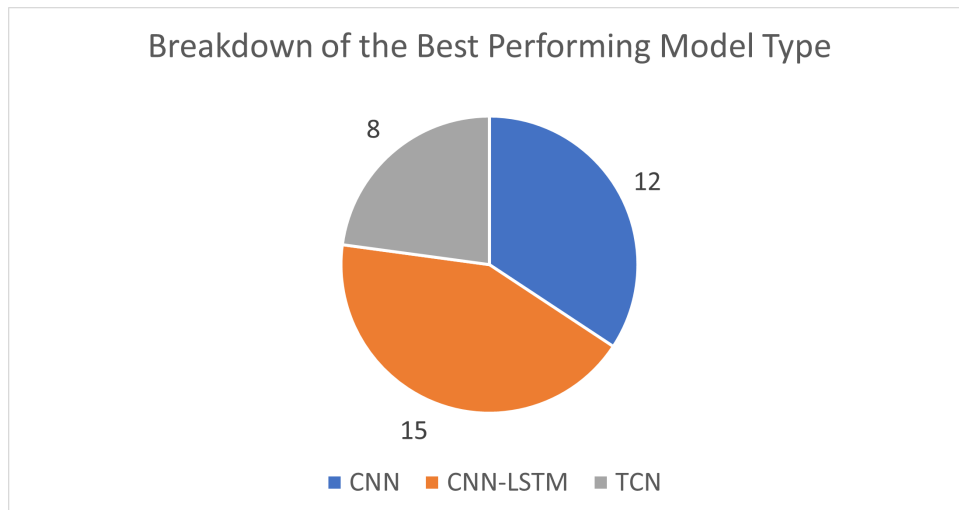


Fig. 16: Number of times the best performing model was of each architecture type (CNN, CNN-LSTM, TCN).

Patient ID	Best Implementation	PR AUC
1	Supervised CNN	0.966
2	Supervised CNN	0.737
3	Supervised CNN	1.000
4	Supervised TCN	0.064
5	Unsupervised CNN-LSTM	0.612
6	Supervised CNN	0.878
7	Unsupervised CNN-LSTM	0.115
8	Supervised TCN	0.704
9	Supervised CNN	0.660
10	Supervised CNN-LSTM	0.733
11	Supervised CNN	0.878
12	Unsupervised TCN	0.928
13	Supervised CNN	0.979
14	Supervised TCN	0.742
15	Unsupervised CNN-LSTM	0.720
16	Unsupervised CNN-LSTM	0.319
17	Unsupervised CNN-LSTM	0.414
18	Supervised CNN-LSTM	0.104
19	Supervised CNN-LSTM	0.733
20	Supervised CNN-LSTM	0.706
21	Supervised CNN-LSTM	0.477
22	Unsupervised CNN-LSTM	0.308
23	Supervised TCN	0.953

Table 9: Best-performing implementation for each patient in the CHB-MIT dataset.

6.4 Discussion

We found that it is important to tune the window size and PPL to maximize performance. Preictal-interictal classification performed slightly better in both datasets when using an optimized hyperparameter implementation. However, in the CHB-MIT dataset, this difference was marginal. This is likely because of the size of the dataset. The CHB-MIT dataset has less data per patient compared to the SWEC-ETHZ dataset, so it is harder to properly tune hyperparameters that will generalize to new seizures. The CNN and CNN-LSTM architectures performed similarly in most experiments. This is likely because both use time-frequency transforms followed by two-dimensional convolutions for spatial feature extraction. Even though the architectures are not exactly the same, it is likely that both are capturing similar underlying patterns in the data. The TCN performed fairly well and was able to get good results in some patients when the other two models failed. Although there was not one consistently high-performing model, it is encouraging that different architectures were able to perform well for different patients in different datasets.

The prediction results vary considerably across datasets, patients, and implementations. This demonstrates the variable nature of preictal and interictal data. To account

for this, it is important to have as many lead seizures data in a patient as possible since preictal data is typically scarce. A limitation of our work is that a patient requires three lead seizures in their data to work with this method. It may not always be feasible for a patient’s data to have at least three lead seizures, especially considering the difficulty of data acquisition.

Anomaly detection seizure prediction performance varied significantly across different architectures. Although supervised preictal-interictal classification performed better overall, there were many patients where an unsupervised approach was the best implementation. Additionally, in the SWEC-ETHZ dataset, an unsupervised approach was the best implementation for the majority of patients. This is likely because the SWEC-ETHZ dataset had a much larger recording duration and interictal-preictal ratio. Anomaly detection seizure prediction shows promise, and it may not be necessary to have access to substantial preictal data to predict a seizure.

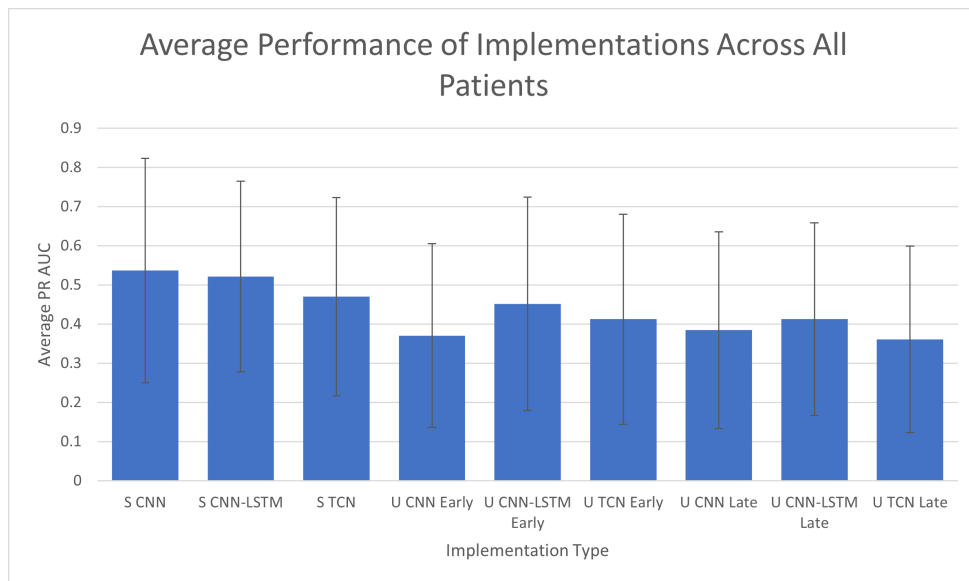


Fig. 17: Average performance of each implementation. S: supervised. U: unsupervised.

Figure 17 shows the average performance in terms of AUC PR across all patients. It can be observed that the supervised CNN and the supervised CNN-LSTM performed the best on average. However, the difference in performance across models is not large, and with a large standard deviation, it is impossible to make a statistical claim on the best performing model. In general, it can be observed that the supervised approaches performed better than the unsupervised approaches with results varying across individual patients. Our results also showed the potential of using unsupervised approaches for seizure prediction. A major advantage is that it only uses unlabelled normal EEG data, which is easier to acquire and is not dependent on an expert to annotate.

7 Conclusions and Future Directions

We developed several supervised approaches and introduced new unsupervised deep learning approaches for predicting epileptic seizures. In each approach, the main goal was to identify a preictal state (either as a class or anomaly) to predict the onset of an incoming seizure. We accounted for the variability of EEG and the preictal period by tuning the window size and PPL using a grid search. We trained personalized models and tuned hyper-parameter using LOSO approach for better generalization of results. This method has achieved good results on more than half of the patients. We experimented with different supervised and unsupervised deep learning architectures on two large EEG datasets. Our results vary across different implementations depending on the patient. The advantage of unsupervised methods is that they do not require preictal data to train, alleviating the challenges around data acquisition, and effort and time spent in labelling. We found that in many cases, an unsupervised approach was able to get similar or even better performance than a supervised approach; however, there was no single best performing model. Our extensive experiments show the feasibility of supervised and unsupervised deep learning approaches for seizure prediction. However, the amount of preictal data per patient appears to be a crucial factor in training generalized models.

A future extension would be to experiment with a larger range for the hyper-parameters. These parameters can also vary across implementations, so optimized hyperparameter implementations with the CNN-LSTM or TCN architecture as the base could be valuable. Another extension would be to try different signal processing methods and advanced CNN and sequential models, including Resnet and Transformers. A breakthrough in reducing intervention time before the onset of seizure would lead to development of therapeutic interventions that can empower epilepsy patients to live without the fear or adversarial outcomes.

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