**Deep learning model for needle electromyography**

**electrodiagnosis in comparison with physician assessment:**

**A retrospective study**

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**Abbreviations**

nEMG, needle electromyography; CNN, convolutional neural network; CI, confidence interval; EMG, electromyography; ReLU, rectified linear unit; AUROC, area under the receiver operating characteristic curve; PPV, positive predictive value; ROC, receiver operating characteristic

**Abstract**

**Objective:** Electromyography (EMG) plays an important role in the diagnosis of neuromuscular diseases by identifying the characteristic abnormalities of the waveform. However, it has some limitations in that there are discrepancies in the interpretations of electrodiagnosis results among physicians, and the accuracy of electrodiagnosis using EMG relies on the experience of physicians. To overcome these limitations of EMG, we investigated whether the deep-learning approach can better classify EMG waveforms into neuropathy, myopathy, and normal physicians’ electrodiagnosis results.

**Methods:** Needle EMG data (58 patients, 382 muscles) from the Seoul National University Hospital database from June 2015 to July 2020 were used. A one-dimensional convolutional neural network model was used as the deep-learning model. The deep-learning model and the six physicians classified and electro-diagnosed the EMG waveforms as myopathy, neuropathy, or normal.

**Results:** The accuracy, sensitivity, specificity, positive predictive value and F1 score of the deep learning model were 0.720, 0.715, 0.858, 0.726, and 0.715, respectively, and the physicians’ mean scores were 0.537, 0.527, 0.770, 0.582, and 0.511, respectively. The performance of the deep-learning model for predicting myopathy, neuropathy, and normal state was also evaluated using the area under the receiver operating characteristic curve, and the results were 0.874 (95% confidence interval [CI] 0.858–0.889), 0.781 (95% CI 0.723–0.839), and 0.847 (95% CI 0.836–0.858), respectively.

**Interpretation:** This study demonstrated that deep-learning could contribute to interpreting the EMG of patients with neuromuscular disease on behalf of physicians and assist physicians’ decision-making regarding diagnosing patients with neuromuscular disease. Large prospective cohort studies with more diverse neuromuscular diseases will further improve the performance of deep-learning-based EMG interpretation in the future.

**Introduction**

Electromyography (EMG) is an electrophysiological test that records the electrical activity generated from nerves, muscles, and neuromuscular junctions via a needle inserted into the muscle or surface electrode during resting and volitional states.1-6 Disorders of the peripheral nerves and muscles are characterized by abnormalities in EMG signals, which reflect the anatomical and physiological states of the peripheral nerves and muscles.1-6

The signal recorded during muscle contraction, consists of the motor unit action potential (MUAP), which can only be measured from muscles in a volitional state and not in a resting state; EMG signals during the resting state cannot distinguish between neuropathy and myopathy. MUAP is an essential key in distinguishing whether a patient has neuropathy or myopathy.1,5-12 Additionally, when neuromuscular disease is in an early state or in a mild abnormal state, abnormal findings are found only in MUAP during a volitional state, not during a resting state.13,14 Patients with peripheral neuropathy commonly show MUAP with high amplitudes, long durations, and reduced recruitment, whereas patients with myopathy commonly exhibit MUAP with small amplitudes, short durations, and early recruitment on EMG evaluation.1,5-12 Needle EMG (nEMG) is a type of EMG that uses a needle as an electrode and is useful for recording MUAPs from individual motor units and deep muscles.15

Although EMG plays an important role in diagnosing normal, neuropathy or myopathy in a patient, it has some limitations. First, the accuracy of EMG diagnosis relies on the proficiency of the examiner. Previous studies have reported that the sensitivity of EMG in the diagnosis of neuropathy, myopathy, and normal was 47–83%, specificity was 73–81% and interrater reliability was 62–81%.16-18 Second, considerable time and effort are required to accurately detect abnormalities in EMG signals. The prevalence of neuropathy and myopathy continues to rise, creating a burden on physicians with regards to the increased number of EMG that require electrodiagnosis.19-22 An accurate, efficient, and automated approach may be helpful for clinical diagnosis using nEMG.

Recently, deep-learning has been used to analyze large datasets in many fields, and it has also been applied to clinical data, including waveform and time series data, such as electrocardiography and electroencephalography.23-26 There have also been a number of studies regarding deep-learning in medical applications where the performance of the deep-learning model was similar to or better than that of humans.27-30 In previous EMG studies, machine learning has been mainly used to analyze the abnormal spontaneous activities that are present on EMG during the resting state or gestures that are not related to the diagnosis of neuromuscular diseases using surface EMG.31-35 To the best of our knowledge, few studies have used using machine learning to analyze nEMG signals during volitional states.

To overcome the diagnostic limitations of EMG and investigate the feasibility of machine learning on nEMG during a volitional state, we developed a deep-learning model that can classify patients as having myopathy, neuropathy or being normal state based on volitional state nEMG signals. We retrospectively reviewed the nEMG data of patients with peripheral neuropathy or myopathy as well as those without any neuromuscular disease. The classification results of the deep-learning model and the electrodiagnosis results of the six physicians were compared.

**Methods**

**Study design and preparation**

The nEMG signal data consisted of 58 patients who visited Seoul National University Hospital between June 2015 and July 2020. Each patient was labeled as having peripheral neuropathy, myopathy, or normal based on the final diagnosis. This study was approved by the Institutional Review Board of Seoul National University Hospital (No. 2008-055-1147) and was conducted in accordance with the Declaration of Helsinki and its later amendments. Informed consent was waved because this study was a retrospective analysis and patient’s private information was anonymized before analysis.

nEMG evaluation was performed using a Nicolet EDX EMG system and a monopolar needle electrode. The filter settings were set at 20 Hz (low cut) and 10 kHz (high cut). The nEMG signals were measured at a sampling rate of 48 kHz. The results of the final 10 s of the nEMG were stored in an EMG machine and used for analysis.

A certified neurologist and certified rehabilitation medicine doctor reviewed the nEMG data and confirmed the diagnosis of all patients. After extracting the nEMG signal stored as numerical data in the EMG device, it was converted into waveform data using MATLAB (version R2020b; The MathWorks Inc., Natick, Massachusetts, United States). Among the created waveform data, electromyographic artifacts that occurred from the movement of the needle electrode or the patient, were removed at the beginning and end of the EMG data. The noise in the remaining data was preserved. Based on the elbow joint of the upper extremity and the knee joint of the lower extremity, muscles close to the trunk were labeled as proximal muscles and those far from the trunk as distal muscles.

**Electrodiagnosis by physicians**

In this study, a web-based nEMG signal labeling platform was developed to allow residents who belonged to different institutions to electrodiagnose patients. After de-identifying the patient information, the nEMG waveform data were displayed on the screen in a similar manner to the EMG data presented on the EMG device (Supplemental Figure 1).

Two neurology residents and four rehabilitation medicine residents assessed the nEMG signal of the muscles examined for each patient and electro-diagnosed patients as myopathy, neuropathy, or normal without clinical information other than nEMG results. When the physician clicked the anonymized patient ID, the nEMG waveform data were presented on the screen along with a sound similar to the sound from the EMG device. The real-time waveform data were presented on the left, and the waveform data were stacked on the right for 500 μs. Physicians were allowed to change the voltage amplitude ticks of the screen to 100, 200, 500 μV, 1, and 2 mV. Physicians first annotated each nEMG signal from different muscles and thereafter diagnosed patients based on the muscle annotation results. The physicians’ electrodiagnosis results were stored within the platform.

**Classification by deep-learning model**

The nEMG signals were down-sampled to 10 kHz to reduce computational complexity. Each data was partitioned into windowed segments with lengths of 0.4 s and hop sizes of 0.1 s. The length and hop size of the windows were heuristically selected. After slicing, there were 3664, 2700, and 1706 segments from patients with neuropathy, myopathy, and normal patients, respectively.

A one-dimensional convolutional neural network (CNN) was used as the deep-learning model.36 The CNN was designed after the motives of ResNes and VGG, which are well-performing image- classification models.37,38 The CNN comprised seven spatial reduction blocks, five residual blocks, and fully connected layers (Supplemental Figure 2). The spatial reduction block consisted of convolutional layers, batch normalization, rectified linear unit (ReLU), and max pooling. The residual block contained similar layers with added residual connections.37 The fully connected layers consisted of 512, 256, 64, and 16 hidden layer neurons with a leaky ReLU activation function. The Softmax function was applied to the final three-output layer. The model received raw signal segments and predicted the probabilities of three class (myopathy, neuropathy, normal).

Hyper-parameters were determined empirically with a learning rate of 10-3, batch size of 32, and 100 epochs. Early stopping was performed by evaluating the accuracy of the validation set every 30 updates, and the patience value was set to 100.39 Cross-entropy loss was used as the loss function, with class weights applied inversely proportional to the number of signal segments from the training set. Deep-learning performance was measured through a 5 × 3-fold cross validation (5 × 3 CV) due to small number of subjects.

The deep-learning model used two stages to classify patients: first, it received multiple segments sliced from a single muscle’s nEMG data and returned the three-class probability of the segment being myopathy, neuropathy, or normal. The probabilities of the segments were averaged to obtain a three-class probability for each muscle. After the probabilities of all the muscles from each patient were computed, a feature vector was produced for the patient using the individual muscle probabilities. The two methods were compared to generate the feature vectors. Another method was to average the muscle probabilities per muscle location label (proximal or distal) to produce a six-dimensional vector. When the patient did not have any muscle probability for a muscle location label, a mean value of 1/3 was imputed to prevent undesired bias. The performance of the deep-learning model presented in this article was obtained by averaging all muscle probabilities. The second stage receives the generated feature vector and classifies the three-class diagnosis of the subject using a logistic regression classifier.

After the model was trained, an explainable artificial intelligence technique named feature visualization was applied to visualize the features that the model focused on.40 The initial feature was sampled from a Gaussian distribution N (0,0.5), and random jitter was applied as regularization with 12.5% of the signal segment length. The learning rate was 10-2 and gradient descent updates were applied 3000 times.

**Assessing the results of the deep-learning model and the physicians**

The performance of the deep-learning model and the physicians were evaluated using the following metrics: accuracy, F1 score, area under the receiver operating characteristic (ROC) curve, positive predictive value (PPV; precision), sensitivity (recall), and specificity. Based on these metrics, we compared the results classified by two versions of the deep-leaning model with and without muscle location labels and averaged the results by six physicians.

**Statistical analysis**

Statistical analyses were performed using R statistical software (version 4.1.0; R Foundation for Statistical Computing, Vienna, Austria) and the Python programming language (version 3.6). Normality for continuous variables was assessed using the Shapiro-Wilk test. The differences among the groups for categorical variables were assessed using Pearson’s χ2 test and those for continuous variables were assessed using the Kruskal–Wallis test. Data are expressed as mean ± standard deviation for continuous variables and number (%) for categorical variables. The degree of agreement was calculated between physicians and the deep-learning model and was expressed with the value of Fleiss kappa. All metrics except accuracy (F1 score, AUROC, PPV, recall, and specificity) were binary classification and were measured by averaging each class metric using the one-versus-rest method.

**Results**

1. Patients’ characteristics

There were 20 patients without any neuromuscular disease; 19 patients with neuropathy, including radiculopathy, motor axonal polyneuropathy, and motor neuron disease; and 19 patients with myopathy, including muscular dystrophy and inflammatory myopathy. The number of nEMG data points used for analysis was 124, 161, and 97 for myopathy, neuropathy, and normal patients, respectively. The remaining demographic characteristics are presented in Table 1.

1. Performance results of deep-learning and physicians

The classification performance of the deep-learning algorithm was compared with the electrodiagnosis results of physicians using the following metrics: accuracy, sensitivity, specificity, PPV, and F1 score. Sliced segments less than 0.4 s were excluded because the deep leaning model only receives segment longer than 0.4 s; total of eight patients and 10 muscle nEMG data were excluded from analysis because these were not annotated by either the physicians or the deep-learning model. The results of the deep-learning model that did not use muscle location information are presented as the above-mentioned metrics. The accuracy, sensitivity, specificity, PPV, and F1 score of the deep-learning model were 0.720±0.040, 0.715±0.050, 0.858±0.023, 0.726±0.032, and 0.715±0.044, respectively. That of the physicians were 0.537±0.069, 0.527±0.069, 0.770±0.034, 0.582±0.076, and 0.511±0.084, respectively (Table 2). The degree of agreement, expressed in Fleiss κ, between physicians was 0.26 and between physicians and the deep-learning model was 0.26 (Supplemental Table 1).

Per-class ROC curves and precision-recall curves of the deep-learning model were measured and depicted. For comparison, the electrodiagnosis results of individual physicians and averaged physicians were measured and depicted in terms of sensitivity, specificity, and PPV in Figure 1. The per-class physicians’ results for myopathy and neuropathy were placed near the ROC and precision-recall curves of the deep-learning model. The deep-learning model outperformed the classification of normal patients. The AUROC of the deep-learning model classification result per-class were 0.874 (95% confidence interval [CI] 0.858–0.889), 0.781 (95% CI 0.723–0.839), and 0.906 (95% CI 0.899–0.913) for myopathy, neuropathy, and normal patients, respectively.

The overall prediction pattern was identified from the confusion matrix of physicians and the deep-learning model (Figure 2). The correctly predicted ratios for myopathy, neuropathy, and normal patients by the deep-learning model were 80.70%±4.96%, 64.71%±4.80%, and 69.05%±12.14%, respectively. The correctly electro-diagnosed ratios for myopathy, neuropathy, and normal patients by the physicians were 49.49%±13.04%, 79.41%±2.94%, and 32.14%±12.20%, respectively.

1. Performance of the machine learning model considering muscle location information

The performance of the deep-learning model did not change significantly when muscle location information was added. The accuracy, sensitivity, specificity, PPC, and F1 score of the deep-learning model considering muscle location information were 0.700±0.053, 0.698±0.048, 0.848±0.026, 0.703±0.058, and 0.694±0.049, respectively (Supplemental Table 2). The AUROC of the same deep-learning model were 0.841 (95% CI 0.811–0.871), 0.736 (95% CI 0.731–0.741), and 0.792 (95% CI 0.719–0.864) for myopathy, neuropathy, and normal patients, respectively (Supplemental Figure 3). The correctly predicted ratios for myopathy, neuropathy, and normal patients were 80.70%±8.95%, 54.90%±2.77%, and 73.81%±3.37%, respectively (Supplemental Figure 4). The prediction accuracies of CNN with muscle location information and those of CNN without muscle location information were as follows: the former and latter results for myopathy, neuropathy, and normal patients were 80.70%±8.95%, 54.90%±2.77%, and 73.81%±3.37% and 80.70%±4.96%, 64.71%±4.80%, and 69.05%±12.14%, respectively (Supplemental Figure 4).

1. Mis-predicted cases of the deep-learning model

To analyze the reason for the misclassification of the deep-learning model, we reviewed misclassified nEMG signals. Examples of these are shown in Supplemental Figure 5.

1. Learned features of the deep-learning model

The learned features of the deep-learning model were identified from the signals created by applying feature visualization. The generated signals were similar to the typical characteristics of neuropathy, myopathy, and normal patients. The waveform most likely to be predicted as myopathy and neuropathy showed small amplitudes and short durations (Figure 3A,B) and high amplitudes and long durations, respectively (Figure 3C,D). Thus, we validated that the deep-learning model made predictions based on relevant features rather than on artifacts.

**Discussion**

The aim of the present study was to validate the diagnostic performance of machine learning using nEMG signals in a volitional state, and whether machine learning could support physicians’ decisions for more accurate and efficient diagnosis. For this purpose, deep-learning was used to classify the nEMG signals as myopathy, neuropathy, and normal, and the classified results were compared with the electrodiagnosis results of six physicians. The results demonstrated that our deep-learning model outperformed the electrodiagnosis results of physicians in all performance scores, with an accuracy of 0.720±0.040 for the deep-learning model and 0.537±0.069 for physicians.

There have been reports that machine learning shows good performance when applied to analyzing EMG signals.31-35 However, previous studies regarding EMG have mainly focused on abnormal spontaneous activities in the resting state, and the signals were preprocessed to form two-dimensional features, which may increase computational complexity.33,34 It is well known that EMG evaluation is important in the diagnosis of the neuromuscular disease; and EMG signals during a volitional state play an essential role in differentiating myopathy, neuropathy, normal patients as well as identifying whether the neuromuscular disease is in the initial state or the minimally involved state.1-6,8,13,14,41 To the best of our knowledge, only few studies have analyzed EMG data during volitional states using deep-learning. We analyzed the nEMG signal in the volitional state, which is important for the electrodiagnosis of neuropathy and myopathy, and confirmed that it showed better performance than physicians.

To diagnose a patient, the nEMG results of all tested muscles should be considered together. However, not all nEMG signals from muscles show abnormalities, and the number of muscles examined may differ slightly between patients; therefore, this variation needs to be considered for the deep-learning model. To solve this variation problem, we constructed feature vectors for patients from the muscle signal prediction probabilities and utilized an additional classifier to determine the classification result of the patient. This method allows the deep-learning model to not only consider all the signals measured from different muscles from different patients in a consistent format, but also make predictions based on the signal characteristics, and not on the prevalence of nEMG signals in certain muscle types.

Usually, there is a typical pattern of muscle involvement in which peripheral neuropathy mainly shows abnormalities in the distal muscles, whereas myopathy mainly shows abnormalities in the proximal muscles.13 Although muscle location information is meaningful for differentiating neuropathy from myopathy, there was no significant change in the performance of the deep-learning model when muscle location information was used. This may be due to the following two reasons; first, some types of myopathy may show abnormalities in both proximal and distal muscles. For instance, in muscular dystrophy, the distribution of the affected muscles depends on the disease process. Both the proximal and distal muscles are affected in statin-induced myopathy and critical illness myopathy. Myotonic dystrophy type 1,2 or distal myopathy may affect distal muscles more frequently than proximal muscles.13,42,43 Second, the small number of patients may have been insufficient to generate significant features within the muscle location information.

Among the signals misclassified by the deep-learning model, several examples were inspected (S5 Fig). Signals that contained parts with high amplitudes were misclassified as neuropathy (S5A,D Fig), and signals that contained parts with small amplitudes were misclassified as myopathy (S5B,C Fig). From the feature visualization results of the trained deep-learning model, it was determined that the deep-learning model makes predictions based on the amplitude, duration, recruitment, and interference pattern of typical myopathy, neuropathy, and normal patients. Thus, the amplitudes of mis-predicted nEMG signals may dominate the recruitment and interference patterns which results in incorrect prediction by the deep-learning model.

Interestingly, the diagnostic accuracy of the physicians was 0.537, which was lower than expected. This low accuracy can be attributed to the two following reasons. First, our data included 38 patients out of 58 with neuromuscular disease, which is much higher than the real prevalence of approximately 200 per 100,000 people.44 Second, the electrodiagnosis of physicians during this study was different from the real-world diagnosis process. Physicians considered nEMG signals as well as additional information about the patient such as demographics or symptoms, which were absent in the labeling platform.

This study has several limitations. First, it used retrospective data from a single institution. When additional data from other institutions are available, external validation can be performed to further verify model performance. Second, a larger amount of nEMG data needs to be examined to demonstrate the stable performance of deep-learning in nEMG classification. The 58 patients from this study are not sufficient to demonstrate the usefulness of this deep-learning model and a larger cohort may elicit useful information from the muscle location as well. Third, the patient diagnosis labels were only divided into neuropathy, myopathy, and normal patients. There are diverse subtypes of neuromuscular diseases such as chronic inclusion body myositis, ongoing-state dermatomyositis, and late-stage muscular dystrophy, which co-exhibit MUAPs of myopathy and neuropathy, with short and long durations. Additional nEMG data for more specific neuromuscular diseases could improve the usefulness of deep-learning for nEMG clinical assistance. Finally, resting-state nEMG data may be added for better prediction. In some neuromuscular diseases, such as Pompe disease, EMG abnormalities may be revealed only in the resting state of the paraspinal muscles rather than the limb muscles.13,45 Future prospective studies with resting and volitional state EMG data could further improve the applicability of deep-learning for EMG electrodiagnosis.

Few studies have analyzed nEMG signals in a volitional state using deep-learning. The results of this study demonstrated that deep-learning could analyze nEMG signals in a short time and with high accuracy, and that our relatively simple model has the potential to be embedded in an nEMG device. Embedding a fast, accurate, and simple machine-learning model into an nEMG machine could allow clinical assistance within the device without sharing personal medical information. Such an application could not only reduce the burden on physicians, but also lead to a widely applicable, low-cost clinical decision system for use in small medical institutions.

In conclusion, our study presented significant potential for deep-learning to contribute to the automatic computer-aided electrodiagnosis system of patients with neuromuscular diseases.

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**Figure Legends**

Figure 1. Per-class receiver operating characteristic and precision-recall curves of the deep-learning model and six physicians.

Legend: Area under the receiver operating characteristic curve (A) and precision-recall curve (B) were measured and depicted by dividing all data into myopathy, neuropathy, and normal.

The individual physician performance is annotated by the blue cross, and the average physician performance is annotated by the red dot.

Abbreviation: DLM, Deep-learning model

Figure 2. Confusion matrices of (A) physician electrodiagnosis results and (B) the deep-learning model prediction results.

Figure 3. Feature visualization results of the deep-learning model.

Legend: A and B, Myopathy; C and D, Neuropathy; E and F, Normal.

Note that A, C, and E were plotted with 10 ms of x-axis interval and B, D, and F were plotted with 40 ms of x-axis interval to show the recruitment and interference pattern of the overall waveforms.

Supplemental Figure 1. Electromyography labeler platform.

Legend: A web-based labeling platform named ‘EMG Labeler’. nEMG signals were presented in a similar way to the signals from the EMG device.

Supplemental Figure 2. Structure of the deep-learning model.

Legend: There were seven spatial reduction blocks and five residual blocks with one and two convolutional layers, respectively.

Supplemental Figure 3. Per-class receiver operating characteristics and precision-recall curves of the machine learning model with and without the muscle location information.

Legend: The areas under the receiver operating characteristic (ROC) curves (A) and precision-recall curves (B) were measured and depicted by dividing all data into myopathy, neuropathy, and normal.

The individual physician performance is annotated by the blue cross, and the average physician performance is annotated by the red dot.

ROC and precision-recall curves of the convolutional neural network (CNN) algorithm depending on whether the muscle location information was considered (green lines, CNN with muscle location) or not (pink lines, CNN without muscle location).

Abbreviation: DLM, Deep-learning model

Supplemental Figure 4. Confusion matrices of the machine learning model (A) without muscle location information and (B) with muscle location information.

Supplemental Figure 5. Examples of electromyographic signals mispredicted by the deep-learning model.

Legend: Electromyographic signal mis-predicted: (A) normal to neuropathy; (B) normal to myopathy; (C) neuropathy to myopathy; (D) neuropathy to normal; (E) myopathy to neuropathy; (F) myopathy to normal.

Table 1. Demographic characteristics of patients and their needle electromyography data.

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
|  | Myopathy | Neuropathy | Normal | p-value |
| Number of patients | 19 | 19 | 20 |  |
| Female, n (%) | 14 (73.7) | 12 (63.2) | 13 (65) | 0.761 |
| Age (mean±SD) | 52.2±20.1 | 58.4±15.1 | 60.2±16.9 | 0.329 |
| Number of nEMG data according to location of muscle (%) | <0.001 | | | |
| Distal muscles | 60 (48.4) | 97 (60.2) | 80 (82.5) |  |
| Proximal muscles | 64 (51.6) | 64 (39.8) | 17 (17.5) |  |
| Number of nEMG (mean±SD) | 6.53±3.82 | 8.47±4.59 | 4.85±1.93 | 0.006 |
| Total signal length (sec) | 313.54 | 423.12 | 204.31 |  |

Legend: Data are expressed as mean±standard deviation.

Abbreviation: nEMG, needle electromyography

Table 2. The electrodiagnosis results by physicians and classification results by the deep-learning model.

|  |  |  |
| --- | --- | --- |
|  | Physicians (95% CI)a | Deep-learning model (95% CI)b |
| Accuracy | 0.537 (0.491–0.583) | 0.811 (0.682–0.758) |
| Sensitivity (recall) | 0.527 (0.480–0.573) | 0.720 (0.668–0.762) |
| Specificity | 0.770 (0.747–0.793) | 0.853 (0.836–0.879) |
| Positive predictive value (PPV, Precision) | 0.582 (0.530–0.633) | 0.725 (0.695–0.757) |
| F1 score | 0.511 (0.455–0.568) | 0.715 (0.673–0.757) |

Legend: The metrics, except for accuracy, are the average of the binary classification scores for each class from the one-versus-rest method.

a Average results of electrodiagnosis by six physicians.

b Average results of classification using three deep-learning models.

Abbreviation: 95% CI, 95% confidence interval

Supplemental Table 1. Degree of agreement between physicians and the deep-learning model.

|  |  |  |
| --- | --- | --- |
|  | Physiciansa | Deep-learning modelb |
| Overall | 0.260 | 0.256 |
| Myopathy | 0.360 | 0.400 |
| Neuropathy | 0.256 | 0.247 |
| Normal | 0.204 | 0.165 |

Legend: Data are expressed as Fleiss κ.

a Fleiss κ value between the results of the six physicians.

b Fleiss κ value between the results of six physicians and the deep-learning model.

Supplemental Table 2. Classification results by the deep-learning model with and without muscle location information.

|  |  |  |
| --- | --- | --- |
|  | Performance results (95% CI) | |
|  | Without information | With information |
| Accuracy | 0.720 (0.682–0.758) | 0.700 (0.650–0.750) |
| Sensitivity (recall) | 0.715 (0.668–0.762) | 0.698 (0.652–0.744) |
| Specificity | 0.858 (0.836–0.879) | 0.848 (0.824–0.873) |
| Positive predictive value (PPV, precision) | 0.726 (0.695–0.757) | 0.703 (0.648–0.758) |
| F1 score | 0.715 (0-.673–0.757) | 0.694 (0.647–0.741) |

Legend: The average metrics of the classification results of the three deep-learning models are presented. The metrics, except for accuracy, are the average of the binary classification cores for each class from the one-versus-rest method.

Abbreviation: CI, Confidence interval