

# Neural Network Analysis of MRI Scans for FND Diagnosis Check for<br>updates

# **Samiel Azmaien**

*Abstract: Background Functional Neurological Disorder (FND) currently lacks a definitive method of diagnosis, leading to an extremely high rate of misdiagnosis. Methods This project aimed to address the question of improving diagnostic accuracy for FND by utilizing logistic regression models and neural networks, integrating patient MRI data and clinical history to differentiate FND from other neurological disorders. MRI scans were first pre-processed through noise reduction and feature engineering, and then used to train two types of models: logistic regression for general neurological disorder classification and a neural network specifically for FND diagnosis. The diagnostic performance was measured using the ROC AUC metric, with additional evaluation through accuracy, precision, recall, and the F1 score. Results & Conclusions By targeting the most relevant variables from the MRI data, both models demonstrated high efficacy, with the neural network showing a 92% accuracy rate in FND classification.*

*Keywords: Diagnosis, Functional Neurological Disorder, Logistic Regression, MRI Data Scans, Neural Networks*

## **I. INTRODUCTION**

Functional Neurological Disorder (FND) has projected misdiagnosis rates of around 50 cases per 1,000 [\[1\]](#page-3-0). This high rate is due to the lack of an efficient diagnostic method for this condition. Currently, the primary method involves using video telemetry electroencephalography tests to measure the brain's electrical activity for extended periods, lasting up to a day, which is very time-inefficient [\[2\]](#page-3-1). This inefficiency stems from the difficulty in identifying another neurological condition that better explains the initial symptoms in hindsight [\[3\]](#page-3-2). Integrating neural network analysis of patient MRI data with clinical history trials could alleviate the difficulties in diagnosing FND. By achieving this, there will be less pressure to evaluate patients quickly due to the inability to describe the condition clearly. It will also assist program users in understanding the diagnosis despite lacking training in neuropsychiatric principles [\[4\]](#page-3-3). Continuously refining a machine learning algorithm for FND diagnosis can lead to more accurate and timely diagnoses. This, in turn, could lay the groundwork for future explorations into AI-driven diagnostics in neurology and medicine.

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To address the current misdiagnosis issues with FND, the proposed solution involved using a logistic regression model for general neurological disorder classification and a neural network specifically for FND diagnosis. To diagnose FND with the highest accuracy, an algorithm was created that utilized ROC curve plotting, which illustrated the diagnostic ability of a binary classifier system as its discrimination threshold was varied  $[5]$ . This metric is represented on a scale from 0 to 1, where a score of 1 indicates an ideal, flawless diagnostic test and a score of 0.5 suggests a test whose accuracy is no better than random chance [\[6\]](#page-4-1). This quantification served as a crucial indicator of the diagnostic method's performance. By applying this multifaceted model, an accurate diagnosis for FND could be achieved.

#### **II. METHODS**



**Figure 1: MRI Scan of a Patient Diagnosed with FND for the Training Phase [\[7\]](#page-4-2)**

This image is part of a larger dataset of MRI scans from patients diagnosed with FND and a control group without FND. The dataset was divided into training (70%), validation (15%), and test sets (15%). The MRI scans were then preprocessed to extract the relevant features, which included noise reduction, normalization (where a MinMaxScaler was utilized - a process that adjusts the scale of the features), and segmentation [\[8\]](#page-4-3). A convolutional neural network is a type of neural network that utilizes multiple layers and pooling steps to classify data or images and this was used to automatically extract the most relevant features from the MRI scans and clinical history trials that could characterize FND [\[9\]](#page-4-4)[\[10\]](#page-4-5)[\[11\]](#page-4-6)[\[12\]](#page-4-7)[\[13\]](#page-4-8).

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## **Figure 2: Preprocessing the MRI Data using Signal Processing and Feature Engineering**

The extracted features, along with the clinical history attributes inputted by the user, were employed to train a neural network to predict the FND diagnosis. The creation of a convolutional neural network architecture was not within the scope of this project, so ResNet was utilized. This model was composed of multiple dense layers, culminating in a sigmoid activation tailored for binary classification tasks. The use of TensorFlow and Keras facilitated the implementation of this model, allowing for a nuanced architecture that could capture complex patterns in the data.

neural network - Sequential()	
Dense(unit:-64, activation-'relu', input dim-troin data.shape[1]).	
Dense(units-64 activation-"relu'),	
Flatten().	
Dense(units-1, activation-"signoid")	
$_{1}$	
# Kompile the model with loss and optimizer	
	neural_network.compile(optiminer-'adam', Loss-'binary crossentropy', metrics-['accuracy'])
# Train the model with the training data and validation duta	
	neural network.flt(x-train data, y-train Labels, validation data-(validation data, validation Labels), epochs-10, verbose-0)
# Predict probabilities on the test set	
predictions prob - neural_network.predict(test_dutu).flatten()	
# Convert probabilities to binary prediction	
predictions binary - (predictions prob = 0.5) astype (int)	
# Perform and display the t-test	
	print("Parforming t-test between actual labels and predicted probabilities")
perform t test(fest (abely, predictions prob)	
a Perform and display the ROC curve analysis	
print("Displaying NOC curve )	
display roc curve(cest inbels, predictions prob, FAD Neural Network")	
	report - classification report(test Lubels, predictions prob, output dict-True)
	display confusion matrix(test (ubets, predictions prob, "FAD Neural Network")
	display classification report(rest (abels, predictions prob, "FHD Neural Network")
display heatmaps(report; "FAD Neural Network")	
# Visuallie activation patterns for the first hidden layer	
visualize_activation_patterns(neural_network, 'dense_30', test_doto)	
return test (abels, predictions prob-	

**Figure 3: Sequential Model with Two Densely Connected Hidden Layers of 64 Neurons**

Performance metrics were then computed such as accuracy, precision, recall, F1 score, and area under the AUC-ROC curve for a comprehensive evaluation. The model was then readjusted for the validation image set. FND diagnostic statements were then produced and compared to the original MRI data, indicating if the patient was diagnosed or not.

#### **III. RESULTS**



#### **Figure 4: Confusion Matrix for General Neurological Disorder Classification**

Figure 4 presents the prediction and measured summaries of the general classification model, with 503 true negatives, 242 false positives, 4 false negatives, and 251 true positives. The values of true positives (TP) and true negatives (TN)

were fairly maximized, showing a 75.4% accuracy, as depicted in the heatmap below.



#### **Figure 5: Heatmap for Accuracy and F1-Score of General Neurological Disorder Classification**

The F1-score metric combines accuracy and precision to provide a single measure of the model's robustness. Weighted metrics consider the imbalance in the dataset, offering a nuanced view of the model's performance across different classes, as seen in Figure 5.



#### **Figure 6: Heatmap for Precision and Recall for General Neurological Disorder Classification**

The general classification model demonstrated perfect precision for both classes, as indicated by a score of 1.00 in Figure 6. Every instance predicted by the model to be positive (Precision 0) or negative (Precision 1) was correct, with no false positives. The recall for the negative class (Actual 0) also reached a score of 1.00, which means the model successfully identified all negative cases without any misses.

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However, the recall for the positive class(Actual 1) is 0.67, implying that the model misses about a third of the actual positive cases. This may be due to overfitting to the negative class during the training phase, resulting in less effective generalization for positive cases.

## **B. FND Neural Network**



# **Figure 7: Confusion Matrix for Neural Network Analysis of FND**

Figure 7 presents the prediction and measured summaries of the FND neural network, with 557 true negatives, 47 false positives, 31 false negatives, and 365 true positives. The values of true positives (TP) and true negatives (TN) were greatly maximized, showing a 92% accuracy, as depicted in the heatmap below.



## **Figure 8: Heatmap for Accuracy and F1 Score of Neural Network Analysis of FND**

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Figure 8 shows that the neural network achieved a 92% accuracy in diagnosing FND. While this provided a statistically significant result regarding the model's capabilities, accuracy alone does not offer information about the model's performance in individual cases, such as the balance between false positives and false negatives. It is necessary to compare this with other metrics, such as precision and recall. As demonstrated in Figure 9 below, the recall for the negative class (Actual 0) is 0.33, which is quite low, indicating that the model only identified 33% of the actual negative cases.



# **Figure 9: Heatmap for Precision and Recall of Neural Network Analysis of FND**



## **Graph 1: Matplotlib Program that Prints out a ROC Curve Analysis**

Graph 1 presents the ROC curve for the neural network's performance, with the AUC being 0.83. This indicated an above-average discriminative power to differentiate patients with and without FND. The integration of this analysis with a diagnostic statement would allow for more accurate diagnoses of FND in patients.



**Figure 10: Heatmap for Neuron Activation on the First Hidden Layer 'Dense-30'**



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Figure 10 shows a heatmap that depicts the intensity of activations across a grid of neurons in response to features extracted from MRI scans. This illustrates the steps in the neural network's process of learning to differentiate FND cases from non-cases.



**Graph 1: Matplotlib Program that Prints out a ROC Curve Analysis**

In Graph 2 above, a t-test was performed to compare the means of the predictions from the neural network against the actual labels, assuming they are continuous, or against another set of predictions. The distributions of the y\_pred and y true labels were then plotted with a kernel density estimate to visualize that there was no statistical difference in the means.

(e) <b>perform_t_test(</b> y_true, y_pred_probs <b>)</b> ;
# Calculate the t-statistic and the p-value
t stat, p val - ttest ind(y pred probs, y true)
print(f"T-test result: t-statistic={t stat:.2f}, p-value={p val:.3f}")
# Plot the histogram of predicted probabilities and actual labels
mpl.figure(figsize (10, 6))
sbn.histplot(y pred_probs, color="blue", Label="Predicted Probabilities", kde-False)
sbn.histplot(y true, color="red", Label="Actual Labels", kde=False, bins=[-0.5, 0.5, 1.5])
# Plot the means of both distributions
mpl.axvline(npy.mean(y_pred_probs), color-"blue", Linestyle-'--')
mpl.axvline(npy.mean(y_true), color-"red", linestyle-'--')
# Annotate the t-statistic and p-value in the plot
mpl.text(0.5, max(mpl.ylim()), f't-statistic={t stat:.2f}', ha='center', va='bottom', color="blue")
mpl.text(0.5, max(mpl.ylim()), f'p-value={p val:.3f}', ha= center', wa= top', color="red")
# Add the legend, title, and show the plot
mpl.legend()
mpl.title('T-test Comparison')
mpl.xlabel('Value')
mpl.ylabel('Frequency')
$mp1$ . show()

**Figure 11: Function for Performing the T-Test with KDE to Visualize Difference in Means**

## **IV. DISCUSSION AND CONCLUSION**

Classification of general neurological disorders was completed as it aided in the process for the neural network to classify Functional Neurological Disorders. This general classification model served as a baseline comparison to compare the performance of models focused on more specific conditions with a broader model. The model was also able to properly differentiate between FND and a negative/positive state for other neurological conditions based on the general model being present, which was crucial in achieving high accuracy for the FND model. The dataset used for the general classification also aided in fine-tuning feature engineering as the general model allowed the FND model to already have a context of the relevant features to identify.

After completing all three stages of training, validating, and testing the data, a high accuracy score of 92% was reached in diagnosing patients with FND. The considerably higher accuracy of the FND neural network than that of the general classification model might be attributed to continuous refinement and optimization through the validation process. The dataset obtained for the general classification model was also separate from the FND model, which may explain why the data is skewed. Another factor to consider is that an overlap in neurological conditions would make it more challenging for the logistic regression model to distinguish between them accurately. The relatively small batch size of 1,000 images among 10-epoch intervals may have led to an inflated accuracy score. Likely, with a larger batch size, the model would initially have a smaller accuracy.

While the political and social implications of utilizing artificial intelligence in the medical industry have been considered, this study aimed to provide a new perspective on a fairly accurate and optimized integration of neural networks into patient diagnoses. However, an immediate conversion to using artificial intelligence is not ready yet. Further experimentation still needs to be conducted before the safety of patients is entrusted to computational models. Despite this, the usage of machine learning, particularly neural networks, holds significant potential for improving the diagnosis of not only Functional Neurological Disorder but also other hard-totreat conditions as well.

# **DECLARATION STATEMENT**

I must verify the accuracy of the following information as the article's author.

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- **Funding Support:** This article has not been sponsored or funded by any organization or agency. The independence of this research is a crucial factor in affirming its impartiality, as it has been conducted without any external sway.
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- **Data Access Statement and Material Availability:** The adequate resources of this article are publicly accessible.
- **Authors Contributions:** The authorship of this article is contributed solely.

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