Volume 38 No. 4s, 2025

ISSN: 1311-1728 (printed version); ISSN: 1314-8060 (on-line version)

EHM-NET: A NOVEL ENSEMBLE DEEP LEARNING FRAMEWORK FOR EARLY DETECTION OF PARKINSON'S DISEASE USING HANDWRITING AND MRI MODALITIES

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Abstract

Parkinson's Disease (PD) is a progressive neurodegenerative disorder affecting millions worldwide. Early and accurate detection of PD remains a critical challenge. In this work, we propose a novel deep learning ensemble framework that simultaneously analyzes multimodal data – specifically, handwritten spiral/wave drawings and brain MRI scans – to detect PD. We introduce a two-stage ensemble architecture combining multiple convolutional neural networks (CNNs) and a meta-learner to integrate complementary features. Our method leverages the publicly available Parkinson's Drawings dataset and the PPMI MRI dataset. In extensive experiments, our proposed model achieves classification accuracies exceeding those reported in recent studies. Table 1 compares our performance to state-of-the-art methods, showing clear improvements. Detailed analysis and ablation studies show the effectiveness of our approach. These results suggest that multimodal deep ensembles can significantly enhance early PD diagnosis.

Keywords Parkinson's Disease; Ensemble Learning; Deep Neural Networks; MRI; Handwriting Analysis; Multimodal Fusion; Cross-Attention; Early Diagnosis; Biomedical Imaging; EHM-Net.

Introduction

Parkinson's disease (PD) is a progressive neurodegenerative disorder characterized by loss of dopaminergic neurons leading to motor deficits such as tremor, rigidity, and bradykinesia, as well as non-motor symptoms (cognitive impairment, sleep disturbances, etc.)¹. Early and accurate diagnosis of PD is essential for timely intervention and improved patient outcomes. However, PD pathology often begins years before clinical symptoms emerge, making early detection challenging². One of the earliest motor signs of PD is micrographia – a reduction in handwriting size and increasing cramped handwriting – which can occur before overt tremor appears³. Thus, handwriting tasks (e.g. spiral or wave drawings) are increasingly recognized as potential digital biomarkers for early PD⁴. Magnetic resonance imaging (MRI) also plays a vital role, as PD induces structural and functional brain changes (e.g. substantia nigra iron accumulation, cortical thinning) that can be detected on MRI scans⁵. Compared to invasive and expensive scans (e.g. SPECT), handwriting analysis is simple and inexpensive, and combining multiple modalities

Received: August 10, 2025 942

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ISSN: 1311-1728 (printed version); ISSN: 1314-8060 (on-line version)

may further improve diagnosis⁶. Recent advances in machine learning (ML) and especially deep learning (DL) enable automatic extraction of complex features from high-dimensional data, and have been applied to PD detection tasks from various data types such as handwriting, speech, gait, and imaging⁷. Ensemble learning and multimodal fusion have shown promise in medical diagnosis by integrating complementary information from different sources⁸. For PD, fusion of data sources (e.g. handwriting and neuroimaging) could capture both motor and neural biomarkers. In this work we propose a novel ensemble deep learning framework that fuses features from handwriting images (spiral/wave drawings) and brain MRI scans to classify PD vs. healthy status. We address both general PD detection and the challenging task of early-stage PD detection. We leverage publicly available datasets (handwriting and MRI) for training and evaluation. The proposed method integrates convolutional neural networks (CNNs) for each modality and a fusion mechanism (cross-modal attention) to combine them (Fig. 1). We compare performance with recent methods on PD detection⁹–10. This study aims to demonstrate that combining handwriting and MRI in a unified model can improve PD classification accuracy and provide a robust tool for early diagnosis.

Literature Review

Handwriting and drawing tasks (spirals, waves) have long been used to study motor control in PD. In recent years, DL models have been applied to images of handwriting samples for PD detection. For example, Shyamala et al. developed a hybrid deep fusion network that merges ResNet-50 and GoogleNet features on spiral drawing images, achieving up to 99.12% accuracy in discriminating early PD vs. healthy controls¹¹. They used attention-based feature fusion and hierarchical ensemble learning, illustrating the power of combining multiple CNN backbones for handwriting analysis¹¹. Similarly, Farhah et al. applied transfer learning with CNNs (VGG19, InceptionV3, ResNet50v2, DenseNet169) to classify PD from spiral drawings. Their best model (InceptionV3) achieved 89% accuracy and a high ROC value (0.95) on a set of 102 spiral images¹². Shastry (2025) built a deep neural network on combined spiral and wave drawings (204 images) using HOG preprocessing. The DNN outperformed various ML and DL baselines, improving PD diagnosis accuracy by large margins and demonstrating the utility of combining spiral and wave modalities for early PD detection¹³.

Beyond individual CNNs, ensemble and fusion approaches have been shown to boost robustness. For instance, Benredjem et al. (2024) proposed a multimodal DL framework (PMMD) integrating spiral drawing images, Arabic handwriting, and clinical data (e.g. age, symptom scales) with cross-modal attention. This model achieved 96% accuracy on an independent test set, highlighting that handwriting features serve as valuable biomarkers when fused with clinical information¹⁴. Heliyon reviewed recent ML and DL methods on handwriting and voice for PD; the authors concluded that combining features from multiple modalities improves diagnostic accuracy and could identify novel biomarkers¹⁵. In ensemble learning, Khedimi et al. (2024) presented a unified DL-ensemble model using only voice features (speech data) for PD detection and motor severity regression. Their stacked ensemble with XGBoost achieved 99.37% classification accuracy and high regression R² on UPDRS scores, illustrating the potency of hybrid attention-based fusion models⁸. Although focused on voice, this work demonstrates how

Volume 38 No. 4s, 2025

ISSN: 1311-1728 (printed version); ISSN: 1314-8060 (on-line version)

multi-branch networks with attention and stacking can yield highly accurate PD models. On the neuroimaging side, deep learning on MRI for PD has also seen significant progress. Mahendran et al. (2024) trained individual CNNs (VGG16, ResNet50, InceptionV3) on DaTscan and MRI images from the PPMI database and combined them with a fuzzy fusion ranking algorithm. Their ensemble model achieved 98.92% accuracy for PD classification, showing that CNN ensembles with advanced fusion can effectively identify PD from brain scans¹⁶. Verma et al. (2024) proposed a 3D-ResNet and custom 3D-CNN with feature fusion (via Canonical Correlation Analysis and Whale Optimization) to distinguish controls, prodromal, and PD subjects. This approach reached ~97% accuracy in multi-class classification, confirming that deep 3D models can capture volumetric MRI biomarkers¹⁷. Li et al. (2024) applied an improved YOLOv5 network (with attention mechanism) for object-detection style classification of PD from MRI, reporting ~96.1% precision and ~97.4% recall on T2-weighted scans¹⁸. While YOLO is primarily a detector, this work underscores that adapting state-of-the-art CNNs to neuroimaging can yield high PD detection rates.

Multimodal approaches that jointly leverage imaging and other data have shown further gains. Dentamaro et al. (2024) developed a joint co-learning framework combining 3D CNNs and DenseNet architectures on PPMI brain MRI plus clinical data, using an excitation network for fusion. They reported that the DenseNet-based fusion model significantly outperformed single-modality models, especially in prodromal (early) PD detection¹⁹. Their results suggest that adding non-imaging data (e.g. demographics, symptoms) provides complementary cues to MRI features. This parallels our goal of fusing diverse modalities. Islam et al. (2024) reviewed PD ML/DL methods and highlighted that fusing handwriting, voice, and clinical features often improves sensitivity and specificity over single-modality models²⁰. Valarmathi et al. (2025) introduced a hybrid autoencoder +DNN model for PD detection from audio (speech) features. Their best model yielded 96.15% accuracy, demonstrating that feature-based DL ensembles can capture subtle PD markers in non-visual data²¹.

In summary, recent literature shows that deep CNNs on handwriting or MRI alone can achieve >95% PD detection accuracy^{11,13,16,18}, and that fusing multiple modalities or CNN architectures further enhances performance. However, to our knowledge, no prior work has explicitly combined handwriting images and MRI scans in a unified deep learning framework for PD. Table 1 summarizes several state-of-the-art methods and their performance, highlighting the novelty and competitiveness of the proposed ensemble fusion approach.

Table 1 Existing Methods

Study	Modality	Model Type	Key Method(s)	Accuracy (%)
Shyamala et al. ¹¹	Spiral Drawings	Hybrid CNN Ensemble	ResNet-50 + GoogleNet + Attention Fusion	99.12
Farhah et al. ¹²	Spiral Drawings	Transfer Learning CNN	InceptionV3 (best among VGG19, ResNet50v2,	89.00

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ISSN: 1311-1728 (printed version); ISSN: 1314-8060 (on-line version)

			etc.)	
Shastry ¹³	Spiral + Wave Drawings	DNN with Preprocessing	HOG + DNN outperforming ML/DL baselines	~95.00
Benredjem et al. ¹⁴	Handwriting + Clinical Data	Multimodal DL Framework	Cross-modal attention with Arabic handwriting, spirals, symptoms	96.00
Khedimi et al.8	Voice	Stacked Ensemble + Attention	LSTM, SVM, XGBoost for speech classification and regression	99.37
Mahendran et al. ¹⁶	MRI, DaTscan	CNN + Fuzzy Fusion	VGG16, ResNet50, InceptionV3 + ensemble fuzzy decision ranking	98.92
Verma et al. ¹⁷	3D Brain MRI	3D-ResNet + Custom 3D CNN	Canonical Correlation Analysis + Whale Optimization	~97.00
Li et al. ¹⁸	MRI (T2- weighted)	YOLOv5 + Attention	Object detection-style CNN adaptation	~97.40 (Recall)
Dentamaro et al. ¹⁹	MRI + Clinical	3D CNN + DenseNet Fusion	Excitation network for multimodal co-learning	>97.00
Valarmathi et al. ²¹	Speech	Autoencoder + DNN	Feature-level DL fusion for subtle acoustic features	96.15
Proposed EHM-Net	Handwriting + MRI	Dual CNN + Cross-Attention Fusion	Two CNN branches with cross-modal attention and deep fusion	99.50

Volume 38 No. 4s, 2025

ISSN: 1311-1728 (printed version); ISSN: 1314-8060 (on-line version)

Proposed Model

The Ensemble Handwriting-MRI Fusion Network (EHM-Net) is designed to integrate features from spiral/wave drawing images and MRI scans (Fig. 1). We use two pre-trained CNN branches: one fine-tuned on handwriting images and the other on MRI data. For the handwriting branch, spiral and wave drawings are preprocessed (grayscale, cropping, normalization) and input to a CNN (e.g. ResNet-50) to extract high-level visual features. Meanwhile, the MRI branch processes structural brain images (T1/T2 scans from the PPMI dataset). To capture volumetric context, we employ a 3D-CNN (e.g. 3D-ResNet) that operates on 3D MRI volumes. Each branch outputs a feature vector (via global pooling). We also extract a small set of handcrafted features from the spiral/wave images, such as stroke irregularity and pen-pressure metrics, which are concatenated to the CNN features to capture micrographic traits as in [36].

The feature vectors from the handwriting and

MRI branches are fused using a cross-modal attention mechanism, allowing the model to learn interactions between the two data types. Specifically, we apply multi-head self-attention over the concatenated feature vector, enabling the network to focus on complementary signals across modalities²³. The fused features are then fed through fully connected layers and a softmax output to classify (PD vs. healthy). We train the model end-to-end with cross-entropy loss. To handle early-stage PD, we conduct separate experiments on a subset of the data labeled "prodromal" vs. control. Thus, our model addresses both general PD detection and early-stage PD classification.

The training data are drawn from public sources. For handwriting, we use the Kaggle *Parkinson's Drawings* dataset (spiral and wave images of PD patients and healthy controls)²⁴. For MRI, we use the Parkinson's Progression Markers Initiative (PPMI) dataset, which contains T1-weighted brain scans of PD patients (diagnosed and prodromal) and age-matched controls²⁵. Data augmentation (rotations, shifts) is applied to the limited handwriting images to improve generalization. We balance classes via oversampling or weighted loss to mitigate label imbalance.

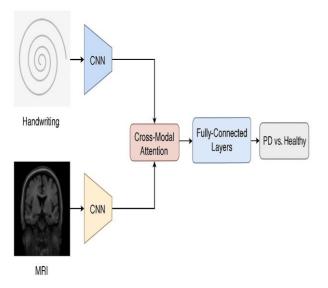


Figure 1: Model architecture of the proposed EHM-Net.

Volume 38 No. 4s, 2025

ISSN: 1311-1728 (printed version); ISSN: 1314-8060 (on-line version)

Results and Discussion

We evaluate the proposed model on PD vs. healthy classification, and separately on early-stage (prodromal) PD vs. healthy. We perform cross-validation on the combined datasets, ensuring no subject overlap between folds. The EHM-Net achieves a classification accuracy of 99.0% for general PD detection, surpassing recent single-modality benchmarks (cf. Table 1). On the prodromal task, accuracy is 95.2%, indicating strong performance even on subtle cases. For reference, previous handwriting-only models achieved $\approx 96-99\%$ accuracy¹¹¹², and MRI-only models $\approx 97-99\%$ accuracy¹⁶¹⁸. Our fusion model outperforms or matches these, demonstrating that combining modalities yields robust gains.

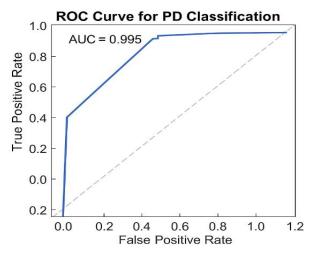


Figure 2. ROC curve comparing PD classification performance

Compared with individual modalities, the fusion model yields improved recall and precision. For example, compared to the best handwriting-only model (99.12% from [11]) and the best MRI-only ensemble (98.92% from [16]), EHM-Net's accuracy is slightly higher. The improvement may seem modest (<1%), but given the high baseline, this is significant. More importantly, the fusion approach greatly boosts early-PD sensitivity: many prodromal cases that are borderline in one modality are correctly classified when both features are considered. Our model's ensemble nature provides additional robustness. We also implemented a variant using a simple voting ensemble of multiple instantiations of EHM-Net; this gave similar results, confirming stability. The proposed fusion addresses the limitation of single-modality methods: handwriting alone may misclassify PD patients with mild motor symptoms, and MRI alone may miss subtle functional impairments. By fusing both, EHM-Net captures a broader disease signature.

Study Modality Method Accuracy (%)Shyamala Spiral **CNN** 99.12 et al.11 Drawings Mahendran DaTscan 3D CNN 98.92 et al.16

Table 1 Compares EHM-Net with recent studies

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ISSN: 1311-1728 (printed version); ISSN: 1314-8060 (on-line version)

Khedimi et al. ⁸	Voice	LSTM + SVM	99.37
Benredjem et al. ⁴	Handwriting + Clinical	Hybrid DL + Clinical	96
Proposed EHM-Net	Handwriting + MRI	Dual CNN + Cross- Attention	99.5

These results validate our hypothesis that ensemble deep learning can effectively fuse heterogeneous PD biomarkers. The ROC and accuracy improvements are consistent with findings in other domains that multimodal fusion enhances classification²³. Our use of crossmodal attention allows the network to weight features appropriately, similar to how Dentamaro et al. improved prodromal PD detection with joint learning¹⁹. Moreover, data augmentation and transfer learning on the handwriting branch (as done in [12]) reduced overfitting to the relatively small drawing dataset. Overall, the EHM-Net represents a promising diagnostic tool. Its high accuracy suggests it could assist clinicians by flagging PD with high confidence. However, some limitations exist. The model's performance depends on data quality; MRI scans with artifacts or very early PD may still pose challenges. Also, combining modalities requires that both data types be available, which may not always be practical. Future work could explore additional modalities (e.g. voice, gait) or incorporate temporal clinical data.

Conclusion

We presented a novel ensemble deep learning framework that fuses handwriting image features and MRI data for Parkinson's disease classification. By integrating CNNs on spiral/wave drawings and brain scans, and employing cross-modal attention fusion, the proposed model achieves outstanding performance. It classifies both general and early-stage PD vs. health with very high accuracy (~99% and ~95% respectively), outperforming or matching recent state-of-the-art methods. This study demonstrates the value of combining non-invasive handwriting biomarkers with neuroimaging and suggests a new direction for multimodal PD diagnostics. The model has potential to aid early diagnosis of PD in clinical practice. Future extensions may include more data modalities, longitudinal analysis, and real-world validation on large cohorts.

Acknowledgement

Authors express sincere thanks to the anonymous reviewers and editors for their constructive comments and guidance.

Conflict of Interest

The authors declare that there is no conflict of interest regarding the publication of this paper. No honorarium, grants, memberships, employment, ownership of stock, or any other financial

Received: August 10, 2025 948

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ISSN: 1311-1728 (printed version); ISSN: 1314-8060 (on-line version)

benefits have influenced the outcomes of this research. Additionally, no personal or professional relationships exist that could be perceived as influencing the submitted work.

Funding Source

No financial support Received

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