

Explaining Parkinson's Disease Computational Diagnostic based on Speech Analysis

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Resumo

[TRADUZIR PARA PORTUGUÊS] Parkinson's Disease (PD) is a neurodegenerative disorder that affects the central nervous system. The disease manifests itself in the patient's speech, which usually becomes slurred, monotonic, and breathy. These symptoms provide a powerful biomarker for the detection of PD. The present work will analyse the subject's speech, representing it with a set of acoustic features. With this representation, a Machine Learning (ML) model will be trained to diagnose PD. By performing cross-language tests on this classification model, we will evaluate the hypothesis that a ML classifier can correctly diagnose PD. In addition, this diagnosis will be independent of the language spoken by the subject. Furthermore, we will explore the use of an explainability model, which will "translate" the classification model's diagnosis to a medical professional. The model will provide essential information for the clinician to trust and use this tool. Despite the good results achieved by many ML classification models, their acceptance for the diagnosis of this disease has not yet been achieved. Clinicians are unable to use the model's diagnosis, as it lacks a medical-oriented interpretation. Therefore, this work is motivated by the practical need of a universal, language-independent model that can provide enough human-understandable information to support clinical usage of ML models on PD diagnosis. This project will provide a tool that can boost the transfer of such models from test environments to real-life usage.

Abstract

Parkinson's Disease (PD) is a neurodegenerative disorder that affects the central nervous system. The disease manifests itself in the patient's speech, which usually becomes slurred, monotonic, and breathy. These symptoms provide a powerful biomarker for the detection of PD. The present work will analyse the subject's speech, representing it with a set of acoustic features. With this representation, a Machine Learning (ML) model will be trained to diagnose PD. By performing cross-language tests on this classification model, we will evaluate the hypothesis that a ML classifier can correctly diagnose PD. In addition, this diagnosis will be independent of the language spoken by the subject. Furthermore, we will explore the use of an explainability model, which will "translate" the classification model's diagnosis to a medical professional. The model will provide essential information for the clinician to trust and use this tool. Despite the good results achieved by many ML classification models, their acceptance for the diagnosis of this disease has not yet been achieved. Clinicians are unable to use the model's diagnosis, as it lacks a medical-oriented interpretation. Therefore, this work is motivated by the practical need of a universal, language-independent model that can provide enough human-understandable information to support clinical usage of ML models on PD diagnosis. This project will provide a tool that can boost the transfer of such models from test environments to real-life usage.

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Palavras Chave

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1 Introduction

Neurodegenerative diseases are the most debilitating disorders that ail human kind, and the fourth leading cause of death. Neurodegenerative diseases affect the patient's thinking, movement, cognitive behavior, and memory, causing impairments and disabilities. These diseases include serious disorders like Alzheimer's Disease (AD) and Parkinson's Disease (PD) (Rai et al., 2019).

PD is the second most common neurodegenerative disease. It was estimated that 1% of people over 60 years old are affected with PD (Tysnes & Storstein, 2017). In 2015, more than 6 million people suffered from this disease worldwide. This value is projected to double by 2040, mainly driven by the increase of life expectancy (Dorsey, Sherer, Okun, & Bloem, 2018). One of the consequences of PD is the development of dementia. Almost half the PD patients develop dementia in the first 10 years after diagnosis (Williams-Gray et al., 2013), reaching over 80% after 20 years (Hely, Reid, Adena, Halliday, & Morris, 2008).

Early detection of PD can be critical for the life quality of the patient. Hence, the earlier the diagnostic is made, the earlier the treatment can begin, thus starting to control the evolution of the disease and improving the comfort of the patient. Furthermore, the majority of the treatment costs occur during the later stages of the disease, reinforcing the importance of early diagnosis (Pagan, 2012).

Over the last years, medicine and health care have been a prime focus for Artificial Intelligence (AI) and Machine Learning (ML). Numerous models have been tested to these areas, demonstrating impressive results in early detection of many diseases, among other tasks. However, the majority of these experiments focuses only on maximizing accuracy performance. Hence, a large problem remains unsolved on the real application of the previously referred models, as explainability has yet to become a focus for any of these works (Magesh, Myloth, & Rijo, 2020). Replacing medical decision-making with non-explainable, black-box ML models, can be contravening with the profound ethical responsibilities of clinicians (London, 2019). Consequently, the lack of explainability and interpretability of ML models used in these areas can seriously limit their chances of adoption in real practice (Vellido, 2019). Therefore, the application of explainable models will increase the possibility for medical professionals to understand a model's output, thus increasing the acceptance of AI systems in such tasks (Holzinger, Biemann, Pattichis, & Kell, 2017).

This work aims to improve our ability to automatically and more accurately diagnose PD, by building a classifier that differentiates PD patients from healthy people. This work aims, not only to allow PD to be detected earlier, but also to increase the degree of confidence of the diagnostic, leveraging the accuracy of the classifier close to 100%. Additionally, by using an explainability model, human-understandable explanations for the given classification (Parkinson or Healthy) of each patient will be generated to foster the use of ML models to support PD's diagnosis.

The document is structured as follows. Section 2 describes PD and state-of-the-art methodologies for PD computational diagnosis. Next, section 3 dives into the concept of Explainable Artificial Intelligence (XAI), and reviews multiple approaches developed in this area. Section 4 describes the experimental setup of this work, followed by section 5, which reports the results and their discussion. Finally, Section 7 presents the conclusions and future work.



Parkinson's Disease

PD is a common cause of dementia. It consists of a neurodegenerative disorder that affects the central nervous system. Symptoms begin gradually and worsen over time ([National Institute of Aging, 2017](#)).

Dementia is a syndrome that reduces the cognitive function. It affects a wide range of mental capacities, such as memory, comprehension, calculation, and language. Several factors can contribute to the development of dementia: factors that cannot be changed, such as age (the percentage of people over 60 years old suffering from dementia is estimated to be 5-8% ([World Health Organization, 2020](#))) and genetics (having a family history of dementia increases one's probability of developing some form of dementia), and factors that can be changed, such as a poor diet and lack of exercise, smoking, diabetes, vitamin and nutritional deficiencies, excessive alcohol consumption and cardiovascular risk factors ([Mayo Clinic, 2019](#)).

Dementia impacts the patient physically and psychologically, but also its carers and family (as it creates a high level of dependency for the patient), and society at economic and social levels ([World Health Organization, 2020](#)).

2.1 Symptoms

As previously stated, symptoms begin gradually and worsen over time ([National Institute of Aging, 2017](#)). The most common symptoms include resting tremors (where hands or arms start shaking when resting), bradykinesia (or slowness of movement), muscle stiffness, which results in difficulty in moving and producing facial expressions, postural instability, which reduces the ability to maintain a steady posture, and dystonia, a condition in which patients have involuntary and repetitive muscle movements. In particular, PD also affects speech ability. Slurring and mumbling are observed in PD patients' speech, which is often observed to be monotone and breathy. The speech rate is also affected, as most patients speak slowly, although others speak too fast. Finally, cognitive problems have been associated with the disease, manifested as a difficulty in finding the correct words (which also contributes to slowing the speech) ([Clinic, 2020](#)).

2.2 Speech Impairments

PD patients exhibit multiple speech impairments, both at acoustic and at language levels. Acoustic parameters of speech, such as the Fundamental Frequency (F0) ([Harel, Cannizzaro, Cohen, Reilly, & Snyder, 2004](#)), pause duration ([Harel et al., 2004](#)) or vowel space time ([Goberman & Elmer, 2005](#)) have been shown to distinguish PD from Healthy Controls (HC).

90% of PD patients are reported to have speech and voice disorders (Froelich, Wróbel, & Porwik, 2015), which show that this biomarker can be an important source of information to detect PD. Instances of incomplete closure of vocal folds along with bowing folds during phonation have been reported (Perez, Ramig, Smith, & Dromey, 1996), leading to noise presence, typically characterized by measures such as Glottal-to-Noise Ratio (GNR), Noise-to-Harmonics Ratio (NHR), Harmonics-to-Noise Ratio (HNR) and Voice Turbulence Index (VTI). An increase in the average values of F0, jitter, and shimmer have also been measured in PD patients.

2.3 *Diagnosis*

To measure the progression of PD, Hoehn and Yahr (H&Y) was proposed (Hoehn & Yahr, 1967). This scale describes the disease through five stages, ranging from stage one, where the patient's symptoms are mild and only manifest on one side of the body, to stage five, where patients are confined to a wheelchair or a bed.

Two decades later, Unified Parkinson's Disease Rating Scale (UPDRS) was created (Fahn & Elton, 1987). It consists on a 50 question test, separated into four sections: (1) Mentation, behavior, and mood, (2) Activities of daily living (ADL), (3) Motor, and (4) Complications (Fish, 2011). Sections 2 and 3 include speech evaluation (section 2 contains questions for self-assessment of speech impairments, while section 3 evaluates free speech from the patient, for an evaluation made by the clinician). The UPDRS test has proven to be very effective in the diagnosis of PD, although favoring moderate and severe impairments. Hence, it may not be ideal to detect mild disease-related signs and symptoms (Goetz et al., 2003). Furthermore, the UPDRS presents some ambiguities (Goetz et al., 2003). In 2003, the Movement Disorder Society (MDS) published a state-of-the-art review on the UPDRS stating the test's inability to measure the severity of PD (Goetz et al., 2008). Furthermore, the heterogeneity of the test (evident on section 4 that uses a mixture of 5-point options with "yes" or "no" questions) hampers a global analysis. Additionally, some redundancy has been found in ADL and motor sections, which increases the time required to perform the test. Lastly, cultural biases have also been identified, in items such as Dressing and Cutting Food/Handling Utensils.

The MDS created the MDS-UPDRS (Goetz et al., 2008), an update on the UPDRS correcting several problems recognized in the original version. Specifically, the response type was homogenized (all questions are answered in a 4-point scale). Some of the redundant items were removed and unrepresented areas in the UPDRS were added. Finally, cultural bias was eliminated. An appendix regarding non-motor problems was also inserted in the MDS-UPDRS.

2.4 *Computational diagnosis*

Over the last years, many experiments have been conducted to diagnose PD using ML models. Such projects have achieved positive results, which are reviewed in this section.

2.4.1 *Speech production tasks*

The most common speech production tasks used for PD classification are:

- productions of a sustained vowel, as there are major variations in glottal noise and tremors in patients with PD (Godino-Llorente, Shattuck-Hufnager, Choi, Moro-Velázquez, & Gómez-García, 2017)
- Diadochokinesia (DDK), which consists of a fast repetition of sounds that imply quick succession of movements with the mouth and tongue (for this task, it is normal to use the pseudo-word /pa-ta-ka/)
- Text-dependent Utterances (TDU)
- Text reading

Several speech production tasks to detect PD were tested (Pompilli et al., 2017) – Sustained vowel phonation (/a/), maximum phonation time (/a/), rapid repetitions of the pseudo-word /pa-ta-ka/, reading of words, sentences and texts, and storytelling guided by visual stimuli. Two approaches were carried out. First, a sentence-level vector was created, with which the classifier achieved accuracies between 55% (with a sustained vowel phonation /a/ production task) and almost 71% (where the speech production task was reading out loud prosodic sentences). Secondly, all sentences were segmented into 4-second segments, with a time shift of 2 seconds. Using the features extracted at a segment level, the classifier achieved accuracies between 58% (with a sustained vowel phonation /a/ production task) and 85% (where the speech production task was reading of prosodic sentences). For this work, the authors used the FraLusoPark dataset (Pinto et al., 2016), which contains audio from 60 PD and 60 HC. The participants were European Portuguese speakers.

A set of 22 acoustic features was extracted from the Parkinson’s Disease Detection Dataset (Little, McSharry, Costello, & Moroz, 2007) and the Parkinson’s Telemonitoring Dataset (Tsanas, Little, McSharry, & Ramig, 2009). The Parkinson’s Disease Detection Dataset includes speech by 23 patients with PD and 8 HC producing sustained vowels. The Parkinson’s Telemonitoring Dataset contains speech from 42 PD patients producing sustained vowels. Using multiple ML classifiers, the system achieved an accuracy of almost 97% using a Gaussian Process Classification (GPC). With this model, the sensitivity reached 88% and the specificity went slightly above 97% (Despotovic, Skovranek, & Schommer, 2020).

To study the relevance of each phonemic group in detecting PD, three datasets were used – GITA (Orozco-Arroyave, Arias-Londoño, Vargas-Bonilla, Gonzalez-Rátiva, & Nöth, 2014), Neurovoz (Moro-Velázquez, Gomez-Garcia, Godino-Llorente, & Dehak, 2019), and CzechPD (Rusz et al., 2013). Neurovoz contains the results for multiple tasks – DDK, TDU and a monologue, based on a picture description – from 47 PD patients and 32 control Spanish Castilian speakers. GITA contains multiple speech production tasks from 50 PD patients and 50 HC Spanish Colombian speakers – DDK, TDU and a monologue. The CzechPD subset considered for this study contains only the DDK task, produced by 20 newly diagnosed and untreated speakers with PD and 14 HC, all Czech speakers. Using a Gaussian Mixture Model - Universal Background Model (GMM-UBM) classifier pre-trained with an auxiliary Spanish Castilian dataset, Albayzin (Moreno et al., 1993), the model yielded an classification accuracy of 94% for the CzechPD dataset, 89% for Neurovoz, and 84% for GITA (Moro-Velázquez et al., 2019).

Sustained vowels and text reading tasks were tested to differentiate PD from HC (Braga, Madureira, Coelho, & Ajith, 2019). The authors use three datasets – Proença (Proença et al., 2014) (containing audio from 22 PD patients in European Portuguese), UCI (Erdogdu Sakar et al., 2013) (with audio from 20 PD and 20 HC) and a dataset created for the purpose of this study

by the authors. The Proença dataset contains word and text reading tasks and the UCI contains results from the sustained vowel task from the patients and healthy controls. The authors tested multiple ML classifiers, such as Neural Networks (NN), Support Vector Machines (SVM) and Random Forests (RF). This work yielded an accuracy of almost 95% with the RF classifier and slightly above 90% with NN (with 4 layers, comprising 7, 7, 6 and 7 neurons, respectively) and SVM.

2.4.2 Feature selection

Multiple acoustic features have been used to attempt to distinguish between PD and HC.

Cases of incomplete vocal folds closure along with folds bowing during phonation were reported (Perez et al., 1996), leading to the presence of noise, that is typically characterized using measures such as NHR, GNR, HNR, and VTI. Some feature values have also been found to increase in PD patients, such as average F0 and jitter (Bang, Min, Sohn, & Cho, 2013) and shimmer (Kent, Vorperian, Kent, & Duffy, 2003).

A set of 5 acoustic features – F0, correlation dimension, HNR, detrended fluctuation analysis and recurrence period density entropy – were selected from a set of 22 acoustic features by using Gaussian processes for regression and classification combined with Automatic Relevance Determination (ARD) (Despotovic et al., 2020). The authors tested multiple ML classifiers (SVM, RF, GPC, among others). The GPC achieved an accuracy of almost 97%, although the model's sensitivity was left on 88% (wrongly classifying 12% of the patients). The specificity reached 97%.

The adequacy of different phonemic groups in identifying PD patients was analyzed (Moro-Velázquez et al., 2019). The work describes the concept of phonemic grouping, which consists of grouping phonemes by their type (such as nasal, fricatives, plosives). Using a GMM-UBM classifier, this work yielded results with accuracies between 77% (using the plosive-nasal-vowel phonemic group) and 94% (with the fricative-nasal phonemic group). The authors extracted Rasta-Perceptual Linear Predictive (Rasta-PLP) (Hermansky, Morgan, Bayya, & Kohn, 1992) and its derivatives, $\Delta + \Delta\Delta$, and labeled them by phonemic group. The focus on the most important sounds has proved that plosive, vowel and fricative segments are the most important for PD detection.

A NN was trained with the VoxCeleb 1 (Chung, Nagrani, & Zisserman, 2017) and 2 (Chung, Nagrani, & Zisserman, 2018) datasets. An affine transformation was applied to the last pooling layer, to retrieve the *x-vectors*, an abstract representation of the input features, which were Mel-frequency cepstral coefficients (MFCC) and its derivatives, $\Delta + \Delta\Delta$. The *x-vectors* are then used as an input to a Probabilistic Linear Discriminant Analysis (PLDA) classifier. The model achieved an accuracy of 90% on TDU production tasks and 79% on DDK production task (repetition of the pseudo-word /pa-ta-ka/) (Moro-Velázquez, Villalba, & Dehak, 2020).

2.4.3 Classification models

Most of the available datasets for this task are very small, considering the usual size for a classification problem. This property made the PD detection difficult. Indeed, complex models are unable to capture the variability of the data from a small dataset, and are therefore unable to correctly simulate and generalize the training set (Andonie, 2010). Therefore, the majority of

the approaches to this problem use traditional machine learning models, such as SVM, RF and K-Nearest Neighbours (KNN), which are able to make accurate predictions training with small datasets. Nevertheless, some experiments have used Multi-Layer Perceptrons (MLP) and other NN architectures, achieving accurate results, in some cases yielding superior performances when compared to other models, such as SVM, and RF (Wan, Liang, Zhang, & Guizani, 2018).

A 114-dimensional feature vector was used as input to a RF. Using acoustic features such as F0, loudness, shimmer, jitter and MFCC, and using 5-fold cross-validation, the classifier achieved an accuracy of 85.1% (Pompilli et al., 2017).

A set of classifiers was used on two PD datasets (Despotovic et al., 2020). The authors extracted the top 5 acoustic features (using ARD) from a set of 22 features. After feature selection, the model achieved an accuracy of almost 97%, using a GPC with Matérns 3/2 and 5/2 as covariance functions. The SVM classifier yielded an accuracy close to 97% as well, whereas the Boosting Classifier (BC) obtained an accuracy around 1% lower, completing the task with close to 96 % accuracy. The RF achieved 96.62% specificity, whereas the model's accuracy almost reached 93%.

From the Naranjo dataset (Naranjo, Pérez, Campos-Roca, & Martín, 2016) 240 recordings were retrieved (Yaman, Ertam, & Tuncer, 2020). From these recordings, 44 acoustic features were extracted. The authors used KNN and SVM classifiers, achieving similar results, yielding accuracies slightly above 91%.

From the Naranjo dataset, a total of 177 acoustic features were retrieved (Yaman et al., 2020). Using the Relief algorithm, the authors selected the 66 more relevant features. Ensemble KNN was compared against Cosine KNN and Gaussian SVM was compared to Quadratic SVM. The Cosine KNN yielded an accuracy slightly above 91%, whereas the Gaussian SVM outperformed the Quadratic SVM, with an accuracy similar to the Cosine KNN (also above 91%).

A total of 2330 acoustic features were extracted from the mPower dataset (Bot et al., 2016) (2268 corresponding to Audio/Visual Emotion and Depression Recognition Challenge (AVEC) 2013 and 62 corresponding to GeMAPS) (Tracy, Özkancab, Atkins, & Ghomi, 2020). With 2023 HC and 246 PD, the authors tested three ML methods to distinguish between PD and HC: L2-regularized Logistic Regression (LR), RF, and gradient-boosted Decision Trees (DT). Because the dataset is heavily biased towards HC ($n = 2023$) compared to PD ($n = 246$), the authors added precision, recall and F1-score to the accuracy as evaluation metrics to compare the performance of each model. The gradient boosted DT achieved the best results, yielding 0.797 for recall, 0.901 precision and an F1-score of 0.836. Similar results were reached with the RF classifier, but with an inferior value for recall (0.693 recall, 0.902 precision and 0.783 for F1-score). The LR achieved the worst results, reaching 0.759 recall, 0.811 precision and 0.784 of F1-score.

A GMM-UBM classifier was trained using one dataset and tested it with three others. The model yielded accuracies between 84% and 94% (Moro-Velázquez et al., 2019).

MLP have also been extensively used for PD classification, having proven their efficacy in performing this task. A 1 hidden layer MLP, used on various sets of acoustic features, was able to classify AD patients with an accuracy of over 92% and HC with an accuracy of almost 91%, surpassing the performance of a KNN model, which yielded accuracies of 90.9% for AD and 87.3% for HC (Lopez-de Ipiña, Solé-Casals, et al., 2014). The Levenberg-Marquardt and Scaled Conjugate Gradient methods were tested as training algorithms for an MLP (Bakar, Tahir, & Yassin, 2010). Using 16 classical acoustic features (such as F0, jitter, shimmer) extracted

from 195 speakers, the authors tested multiple values for the number of hidden units (5, 10, 15, 20, 25) and concluded that the Levenberg-Marquardt outperformed the Scaled Conjugate Gradient, reaching accuracies of over 97% with 25 hidden units, whereas Scaled Conjugate Gradient achieved 79% on 10 hidden units. Using the UCI dataset (Erdogdu Sakar et al., 2013), a set of 23 features was extracted for PD classification (Wan et al., 2018). The authors compared the performance of a Deep Multi-Layer Perceptrons (DMLP), with 5 or 10 hidden layers, with other ML classifiers. The authors reduced the size of the DMLP to 5 hidden layers, using *ReLU* or *softplus* as non-linear activation functions instead of the latter activation function, as these are continuous and can therefore address the vanishing gradient problem that affects Deep Neural Networks (DNN). Results on this experiment concluded that the best performance came from the DMLP using 10 hidden layers, which yielded 80% accuracy, whereas the LR model only reached 77.5% and the KNN could only get to 72.5%. Dropping the size of the DMLP to 5 hidden layers reduced the model's accuracy to 76%, which was still higher than some of the tested models, such as the KNN and RF models.

2.4.4 Universality

As the goal for this work is to develop a model capable of detecting PD for any patient, universality is an important property for the desired model, which can be achieved with language-independency.

Three distinct datasets, one in Spanish, one in German and one in Czech, were used with a GMM-UBM model to train a semi-language-independent model (Orozco-Arroyave et al., 2016). For each experiment, the model was trained with one dataset and tested with another (adding to the training set subsets of the test set with percentages varying from 10% and 80%). Despite reaching accuracies of 97%, high accuracies are only achieved when large portions of the test language are used to train the model. In a fully language-independent model (where the model is trained using one language and tested with another), the model accuracy only reaches 77% (trained with the German dataset and tested with the Czech dataset).

A GMM-UBM was trained using *corpora* in Spanish Castilian, Spanish Colombian and Czech. Cross-language testing resulted in accuracies of 82% (Moro-Velázquez et al., 2019).

3 Explainability Models

XAI is a field of AI that provides techniques and algorithms able to generate interpretable, intuitive, human-understandable explanations of AI decisions (Das & Rad, 2020).

Explaining the decisions made by a black-box model requires knowledge of its internal operations (Das & Rad, 2020), which makes it impossible to use by end-users who are only focused and interested on getting an accurate result. The very nature of a *black-box* ML/Deep Learning (DL) model is a barrier for their real-life usage (Shrikumar, Greenside, & Kundaje, 2017). For a ML model be used in real life situations, the users must have confidence in it. Two definitions of *trust* must be considered: *trust in the prediction*, where the user trusts a prediction sufficiently such that he is comfortable with performing an action based on it, and *trust in the model*, which gives enough confidence to deploy the model. Thus, in order for such model to be deployed, both definitions must be fulfilled (Ribeiro, Singh, & Guestrin, 2016). This is even more important in critical situations, such as medical diagnosis. To address this limitation in ML and DL, many models have been created to generate explanations for a model's predictions.

Creating human-understandable explanations can also aid in finding erroneous behavior in a model. A peculiar discovery was made in an experiment where Fisher Vector classifiers were used for the image recognition task (Bach, Binder, Montavon, Müller, & Samek, 2016). An interpretability technique called Layer-wise Relevance Propagation (LRP) was applied to explain the predictions of the model. In particular cases, where the input image consisted of a horse, it was found that the model primarily based its decision not on any of the physical traits of the horse, but on a copyright tag present on the bottom left of the image that turned out to be a characteristic of all the horse images used in training. This error certainly highlights the need for interpretability of ML/DL models, especially in the medical field, where such errors can severely impact human lives.

3.1 Explanation

An explanation is a verifiable justification for a model's output or decision (Das & Rad, 2020). There are many kinds of explanations, such as a heat map stressing relevant parts of an image (for example, a DaTscan image in PD detection (Magesh et al., 2020)). Some models, such as Local Interpretable Model-agnostic Explanation (LIME) (Ribeiro et al., 2016), base their explanations on activations or parameters of the black-box models, using simpler surrogate models (Das & Rad, 2020).

3.2 Scope

Explainability models can be subdivided in three large groups, based on the scope of their explanations: local, global or mixed.

3.2.1 Local explanations' models

Locally explainable methods are designed to generate an explanation for the model's decision on a single instance of input data (Das & Rad, 2020). Models that provide local explanations fail to provide a global observation of the model. Their explanations do not provide enough information on the original model computations and do not provide enough detail to understand the model's behavior as a whole (Agarwal, Frosst, Zhang, Caruana, & Hinton, 2020).

The concept of Axiomatic Attributions was proposed (Sundararajan, Taly, & Yan, 2017). Consider a function $F : \mathbb{R}^n \rightarrow [0, 1]$ representing a DNN. Let $x \in \mathbb{R}^n$ be the input, and $x' \in \mathbb{R}^n$ be the baseline input (the black image for image networks, for example). Using a straight line path in \mathbb{R}^n from x' to x , the model computes the gradients along the path, in every point. Integrated gradients are obtained by cumulating these gradients. Specifically, integrated gradients are defined from baseline x' to input x as the path integral of the gradients along a straight line path. For each dimension i , $\frac{\partial F(x)}{\partial x_i}$ defines the gradient along dimension i . integrated Gradients (IG) are then calculated as

$$IG_i(x) = (x_i - x'_i) \times \int_{\alpha=0}^1 \frac{\partial F(x' + \alpha \times (x - x'))}{\partial x_i} d\alpha \quad (3.1)$$

The IG provide a measure of the relative importance of each feature on the model's classification of instance x - the higher the IG of feature i , the higher its importance for the classification.

Randomized Input Sampling for Explanation (RISE) was proposed in 2018. This model is based on random masking to locally understand the most important features (for example, in the case of the image classification problem, RISE will determine the most important pixels for the black-box model's classification) (Petsiuk, Das, & Saenko, 2018).

Consider $f : I \rightarrow \mathbb{R}$ to be the model. For the image classification problem, we consider $\Lambda : \{1, \dots, J\} \times \{1, \dots, W\}$ as the image coordinates and I would map every pixel to its RGB representation ($I = \{I | I : \Lambda \rightarrow \mathbb{R}\}$). f is a classifier that returns the probability of an instance of a certain class be present in the image. Considering a random binary mask $M : \Lambda \rightarrow \{0, 1\}$ following a distribution \mathcal{D} . By masking the image with $I \odot M$ (where \odot represents the element-wise multiplication), we preserve only a subset of the pixels of I . By calculating the confidence score $f(I \odot M)$, we can define the importance of every pixel λ , $S_{I,f}(\lambda)$, $\lambda \in \Lambda$, as the average value of the confidence scores of all masked images where $M(\lambda) = 1$. Mathematically,

$$S_{I,f}(\lambda) = \frac{1}{E[M]} \sum_{m \in M} f(I \odot m) \cdot m(\lambda) \cdot P[M = m] \quad (3.2)$$

3.2.2 Global explanations' models

Understanding the model's behavior on a set of input data points could provide insights on the input features, patterns, and their output correlations, thereby providing transparency of model behavior globally. Various globally explainable methods break down complex deep models into linear counterparts, which are easier to interpret (Das & Rad, 2020).

To generate explanations at class, Concept Activation Vectors (CAV) were proposed, which provide interpretations for a NN's internal state in terms of human-friendly concepts (Kim et al., 2018). The model considers a NN with inputs $x \in \mathbb{R}^n$ and a feed-forward layer l with m neurons. Thus, layer l 's activation can be seen as $f_l : \mathbb{R}^n \rightarrow \mathbb{R}^m$. The user chooses a concept of interest C and creates a series of inputs labeled as *contains concept C* and a series of inputs labeled as *does not contains concept C*. The model then calculates the hyperplane separating the two groups of inputs. The CAV is then defined as the vector normal to this hyperplane.

Common interpretability methods (such as saliency maps), calculate the derivatives for the *logit* in terms of the input features. With this approach, these methods are able to measure sensitivity in the set of input features. When combining CAVs and directional derivatives, the model can gauge sensitivity of ML predictions in directional input changes of the concept C , at activation layer l .

In 2020, the concept of Neural Additive Models (NAM) was proposed (Agarwal et al., 2020). The explanations are created by shape functions, relative to each input feature. To parameterize these functions, a NN is created for each function. With this architecture, the model is able to create an exact representation of how NAMs compute a prediction, thus creating an explanation of the model's global behavior.

Consider $\mathcal{D} = \{(x^{(i)}, y^{(i)})\}_{i=1}^N$ as the training set, with N instances, where x is the input vector and y is the target vector. The proposed model was trained using the following loss function:

$$\mathcal{L}(\Theta) = E_{x,y \sim \mathcal{D}}[l(x, y; \Theta) + \lambda_1 \eta(x; \theta)] + \lambda_2 \gamma(\Theta) \quad (3.3)$$

where $\eta(x, \Theta) = \frac{1}{K} \sum_x \sum_k (f_k^\Theta(x_k))^2$ is the output penalty, $\gamma(\Theta)$ is the weighted decay and f_k^Θ represents the k^{th} feature network.

The authors use the cross-entropy loss for binary classification as the task-dependent loss function $l(x, y; \Theta)$, which, considering $p_\Theta(x) = \sigma(\beta^\Theta + \sum_{k=1}^K k_k^\beta(x_k))$, yields

$$l(x, y; \Theta) = (\beta^\Theta + \sum_{k=1}^K f_k^\Theta(x_k) - y)^2 \quad (3.4)$$

where β^Θ defines the parameters to be calculated.

3.2.3 Mixed models

To combine the advantages of the local and global explanations' models, mixed models provide explanations that are able to locally interpret decisions, while also allowing to understand the behavior of the model as a whole.

Similarly to RISE, LRP allow to understand which pixels of the image contribute the most to the model's decision (Lapuschkin et al., 2015). This model, created for DNN architectures, redistributes the relevance of each neuron at the last layer of the network to pixel-wise scores (R_i^l) using the rule

$$R_i^{(l)} = \sum_j \frac{z_{ij} \sum_{i'} z_{i'j}^{(l+1)}}{R_j}, z_{ij} = x_i^{(l)} w_{ij}^{(l,l+1)} \quad (3.5)$$

where i is the i -th neuron in layer l , \sum_j iterates through all the upper-layer neurons to which neuron i contributes. This result can be assessed using a visualization tool, such as a heat map, to explain the model's classification.

LIME is an algorithm that uses *local interpretable representations* of the classification data to generate an output that can be interpreted by humans (Ribeiro et al., 2016). We define $x \in \mathbb{R}^d$ as the original representation of the instance to be explained and $x' \in \{0, 1\}^{d'}$, a binary vector and its interpretable representation. Let $g \in G$, where G is the set of models that can present a interpretable output to the user. We also denote $\Omega(g)$ as a measure g 's explanation complexity and $f : \mathbb{R}^d \rightarrow \mathbb{R}$ as the model to be interpreted. $f(x)$ will be the probability or binary indicator that x belongs to a particular class. Let $\pi_x(z)$ be a proximity measure of distance between x and an instance z to define around x . Lastly, we define $\mathcal{L}(f, g, \pi_x)$ as a measure of how unfaithful g is approximating f in the space defined by π_x . As we want to maximize interpretability while keeping local fidelity, the explanation can be defined as:

$$\xi(x) = \underset{g \in G}{\operatorname{argmin}} \mathcal{L}(f, g, \pi_x) + \Omega(g) \quad (3.6)$$

The algorithm creates samples around x' , weighted by π_x . Considering a perturbed sample $z' \in \{0, 1\}^{d'}$ containing a fraction of the non-zero elements of x' , the original representation $z \in \mathbb{R}^d$ is obtained, so the value $f(z)$ can be calculated. Considering \mathcal{Z} as the set of all perturbed z' with the label $f(z)$, equation 3.6 is used to calculate the explanation.

DeepLIFT (DeepLIFT) was presented as a method to understand the output of a NN by backpropagating the neurons' contributions to every feature of the input (Shrikumar et al., 2017). To assign contribution scores C , DeepLIFT compares the activation of each neuron t to its reference activation value t^0 , using the summation-to-delta property:

$$\sum_{i=1}^n C_{\Delta x_i \Delta t} = \Delta t \quad (3.7)$$

where $\Delta t = t - t_0$ and $C_{\Delta x_i \Delta t}$ is a measure of the difference from the reference value t^0 attributed to the neuron x_i . The reference values t^0 are calculated by defining a set of reference input values $x_1^0, x_2^0, \dots, x_n^0$ for a given neuron, resulting in $t^0 = f(x_1^0, x_2^0, \dots, x_n^0)$, where f is the activation function of the neuron. The choice of reference input values is highly context-dependent. For example, for the MNIST (LeCun & Cortes, 2010), the reference values were set to 0 (which represents the background color of the images, black). For DNA classification tasks, the references were defined based on the expected frequency of each of the elements on the DNA's alphabet (A, C, G, T). This creates a limitation on some applications, as defining reference values may be difficult.

3.3 *Parkinson's Disease diagnosis*

As stated in section 1, ML models used for sensitive tasks, such as detection of PD, lack the ability to generate an explanation to be interpreted by the medical professionals that need to establish a diagnosis. These models, called black-box models (Holzinger et al., 2017), take an input and return as an output a classification, which cannot be interpreted by a medical professional. This problem difficulties the acceptance of these models for such tasks, as the risk of decision-making based on the results of a black-box system raises numerous ethical concerns (Chen et al., 2020).

Image-based explanations were generated for a black-box model (the VGG16 convolutional neural network) on a dataset of SPECT DaTSCAN images of the brain (Magesh et al., 2020). The authors retrieved a 2-dimensional section of the 3-dimensional image, trained, and tested the *black-box* model, which yielded an accuracy of 95.2%, a specificity of almost 91%, a sensitivity of 97.5% and a precision of 95.2%. After the classification, the authors generated a color map over the input images to highlight the regions of interest (the pixels with larger weights for the classification process). This showed that the most interesting regions of the brain for this task were the *putamen* and the *caudate*, confirming the medical background information described, providing trust in the model, as it could be easily interpreted by a medical professional.

Explainability models have been applied to many other medical tasks, such as breast cancer detection (Pfob et al., 2020), identification of individuals with high-risk of depressive disorder (Choi, Shim, Jeong, & Jo, 2020), and early detection of COVID-19 (Punn & Agarwal, 2020).

This area remains almost unexplored for the task of early detection of PD and, to the best of our knowledge, no work has combined explainability algorithms with acoustic-based models for this task.

4

Experimental Setup

Methodology for this work is described in this section. First, the *corpora* used in this work are described, followed by the approaches to be taken will be described (feature selection, classification model, explanation generation model, and multi-language tests). Finally, the evaluation proceedings are presented. Figure 4.1 shows the pipeline for the system’s architecture.

4.1 Corpus Description

As described in section 3.3, most datasets available for this task are very small, thus being unfitted to train neural models (Andonie, 2010). Nevertheless, a small number of common speech production tasks are contained by the datasets. Thus, the differences between some of these datasets are only the model of the microphone used for recording and the language of the test subjects. Hence, some works have combined several datasets (Braga et al., 2019), (Despotovic et al., 2020), (Moro-Velázquez et al., 2019), (Moro-Velázquez et al., 2020) to use different datasets for training and testing, or to mix instances from different datasets in the training and/or testing sets (Orozco-Arroyave et al., 2016), all proving to be accurate in the PD classification task.

In this study, we will use 3 datasets for training and testing the model – FraLusoPark (Pinto et al., 2016), GITA (Orozco-Arroyave et al., 2014), and Mobile Device Voice Recordings at King’s College London (MDVR.KCL) (Jaeger, Trivedi, & Stadtschnitzer, 2019).

The FraLusoPark dataset is composed by speech from 120 patients, half of which are native French speakers and the other half are European Portuguese speakers. The dataset also contains 120 healthy participants as a control group (with the same distribution between French and European Portuguese speakers as the PD participants). Each group of PD patients is divided into three subgroups, based on the number of years since diagnostic: 20 early stage patients (who have been diagnosed less than 3 years before, and present no motor fluctuations), 20 medium stage patients (with a diagnostic made 4 to 9 years before the study, or less than 3 years, and experiencing motor fluctuations), and 20 advanced stage patients, diagnosed over 10 years ago. The patients are recorded twice for every speech production task, BEFORE (at least 12 hours after medication) and AFTER medication (at least 1 hour after medication). FraLusoPark participants were asked to perform a set of speech production tasks: sustain the vowel *a* at a steady pitch, maximum phonation time of the vowel *a* on a single breath, DDK (repetition of the pseudo-word *pa-ta-ka* for a fast rate during 30 seconds), reading aloud 10 words and 10 sentences, formed by adapting part of section V.2 of the Frenchay Dysarthria Assessment of Intelligibility (FDA-2), reading of a short text (adapted to French and European Portuguese), storytelling by guided visual stimuli, reading a collection of sentences with specific language-dependent prosodic properties and free conversation for 3 minutes. In the scope of this study, we will only consider the Portuguese speakers of this dataset. These recordings total 62 minutes and 45 seconds for PD patients’ and 51 minutes and 2 seconds for HC participants’.

The GITA dataset contains recordings of 50 PD patients and 50 HC, evenly distributed between genders. For the PD group, the average age is 62.2 ± 11.2 years for male participants and 60.1 ± 7.8 for female participants. Considering the HC group, the average age is 61.2 ± 11.3 years for male participants and 60.7 ± 7.7 for female participants. Multiple stages of disease progression are considered in this study (time since diagnostic ranges between 0.4 - 20 years for male patients and 1 - 41 years for female patients). All the participants are Colombian Spanish native speakers. Recordings of the PD patients were made no more than 3 hours after the morning medication. Different speech production tasks were performed to examine phonation, articulation and prosody. To analyze phonation, participants were asked to sustain the five Spanish vowels and to repeat the same five vowels, but alternating the tone between low and high. Regarding articulation, a DDK evaluation was performed with the pseudo-words /pa-ta-ka/, /pa-ka-ta/ and /pe-ta-ka/. Finally, for the evaluation of prosody, both PD patients and HC were asked to repeat a series of sentences with different levels of complexity, to read a dialogue between a doctor and a patient (this text contained the complete set of Spanish sounds), to read sentences with a strong emphasis on a set of words and freely speak about their daily routine. These recordings total 15 minutes and 31 seconds for PD patients' and 14 minutes and 41 seconds for HC participants'.

Lastly, the MDVR_KCL dataset was recorded in the context of phone calls, but recorded in an acoustically controlled environment. The dataset contains 16 participants with PD (11 male and 4 female) and 21 HC (3 male and 18 female), totaling 37 English speakers. The PD group contains patients from all the stages of the disease (early, mid and late stages) according to the Hoehn and Yahr scale **hoehn_yahr**. The participants were asked to read a text ("The north wind and the sun" or "Tech. Engin. Computer applications in geography snippet"). Additionally, the text executor started a spontaneous conversation with the participant about various topics.

To homogenize the datasets, only the text-reading tasks will be considered. This yields a total of 131 HC and 125 PD speakers of European Portuguese, Colombian Spanish, and English.

4.2 Data Processing

The original audio files contained full interviews of each test subject, therefore needing segmentation in order to remove unwanted audio fragments. Firstly, silence between speech segments were removed. Next, sounds produced by the subject that were not considered as speech were also delete. Finally, any audio segments containing speech from other participants in the interview (interviewers) were eliminated.

After data processing, FralusoPark recordings totaled XX minutes and YY seconds for PD patients and XX minutes and YY seconds for HC participants. Gita recordings yielded a total of ZZZ minutes and AA seconds for PD subjects and XX minutes and YY seconds for HC participants. Finally, MDVR_KCL recordings totaled XX minutes and YY seconds for PD patients and XX minutes and YY seconds for HC subjects.

4.3 Feature Extraction

In order to extract the features previously mentioned, the openSMILE (Eyben, Wöllmer, & Schuller, 2010) tool was used. To extract the complete set of features, four configurations were

used: *MFCC12_0_D_A.conf* for MFCC's, *PLP_0_D_A.conf* to extract Perceptual Linear Predictive (PLP)'s and *prosodyAcf2.conf*, for prosody features (F0 and HNR).

OpenSMILE was configured to use a sliding window of *25ms* with a frame step of *10ms*. After the extracting, each subject was represented by a list of frames, each one described by a list of features.

4.4 Classification Experiments

Three distinct experiments have been conducted during the present work. First, to create a baseline, a classification model was trained with each train set individually and tested with the corresponding test set (i.e., both from the same database). This first approach allowed to score the classification model without considering multiple languages. Secondly, the same model was trained to evaluate its performance as a semi language-independent classifier. For this, the model was trained using one complete dataset and with a fraction of another dataset (for this work, this fraction represented 90% of the number of subjects in the dataset), thus combining two languages in the same train set. The model was then trained with the remaining percentage (10%) of the second dataset. All the combinations between the three datasets were tested. By testing this semi language-independent version, it was possible to evaluate an intermediate step between a language-dependent and a language-independent classification model, shedding light into the model's sensitivity to the language. Lastly, a completely language-independent model was trained with each one of the datasets. For this last experiment, each model was trained with one dataset and tested with a different one, thus allowing to evaluate the model's ability to diagnose a patient who speaks in a different language used to train the model.

For these experiments, the scikit-learn (Pedregosa et al., 2011) implementation of a MLP was used. Two different architectures were tested to evaluate the ability of the two to learn from the train data. The first architecture contains one entry layer with N neurons (where N is the number of input features), a fully-connected hidden layer with $N + 1$ neurons and an output layer with 1 neuron, whose value represents the probability of the test subject to be classified as PD. **Refer the papers where they say this model is great.** The second architecture also contains an input layer with N neurons, two fully-connected hidden layers, comprising 200 neurons each and, similarly to the first architecture, an output layer with 1 neuron, also representing the probability of the train subject to be diagnosed with PD. For there experiments, the breaking point between HC and PD diagnostic was set to 0.5 (50%).

In order to find the best model configuration, the experiments were repeated testing multiple values for *alpha* (10^{-3} , 10^{-2} , 10^{-1}), maximum number of iterations (1000, 2000, 5000) and solver for weight optimization (lbfgs, sgd, adam).

As described in section 4.3, each patient is represented by a set of feature lists, extracted using openSMILE, using a sliding window of *25ms* with a frame step of *10ms*. When classifying each patient of each test set, the resulting classification is obtained by averaging the model's output for each of the patient's frames.

4.5 Explanations

After the classification experiments, explanations were generated for each individual of the test set (with all models). As the objective of this work was to generate an explanation for each

diagnostic individually, a mixed model was used. By select a mixed model, it is possible to extend this work to generate global explanations, thus allowing a better understanding on the generic impact of each acoustic feature on the diagnostic.

The selected model was LIME. It yielded results that were confirmed by the bibliography on explaining PD diagnostics with SPECT DaTSCAN images of the brain, and this work aimed to verify if the same performance can be achieved using acoustic features.

To explain the diagnostic of one subject, the *explain_instance* method from the *LimeTabularExplainer* class was used with each feature list (each representing a time frame) used to describe it. Next, two operations were performed. First, the classification model's output was averaged between all time frames, creating a final classification probability for the subject. Secondly, each feature weight (which LIME calculated) was also averaged, thus creating a final weight for each feature (from which the top 5 features with larger contribution to the classification were selected and displayed). To this report, a list of values considered to be normal for each one of the features was added. Finally, to assist the interpretation of the report by the medical professional, a small description of each feature was also added. The complete list of normal values and descriptions can be found on tables 1 and 2, respectively.

4.6 Model Evaluation

In order to evaluate the classification model's performance, multiple metrics have been selected, namely:

- *accuracy* (which allows to evaluate the percentage of subjects correctly diagnosed)
- *precision* (that yields the fraction of subjects diagnosed with PD that were correctly classified)
- *recall* (that evaluates the percentage of PD subjects that were correctly diagnosed)
- *F1-score* (that allows to evaluate precision and recall in the same metric)
- *Specificity* (which assess the ratio of subjects classified as HC that were correctly diagnosed)

These metrics shed light on the performance of the models, which allows to determine the best parameters and architecture. Furthermore, recall allows to evaluate the percentage of subjects from the PD group that were correctly diagnosed, which, combined with specificity (that evaluates the number of subjects from the PD group wrongly diagnosed), provides information for medical professionals that provide trust in the model.

To assess LIME's performance, average values for each one of the features were extracted from the bibliography (see [text](#)) and are shown along with the values in each explanation, in order to compare the each subject's feature values with the it's range for a healthy individual. This comparison will allow to evaluate the model's ability to find abnormal values (or their absence) and select those features as justifications for a certain classification.

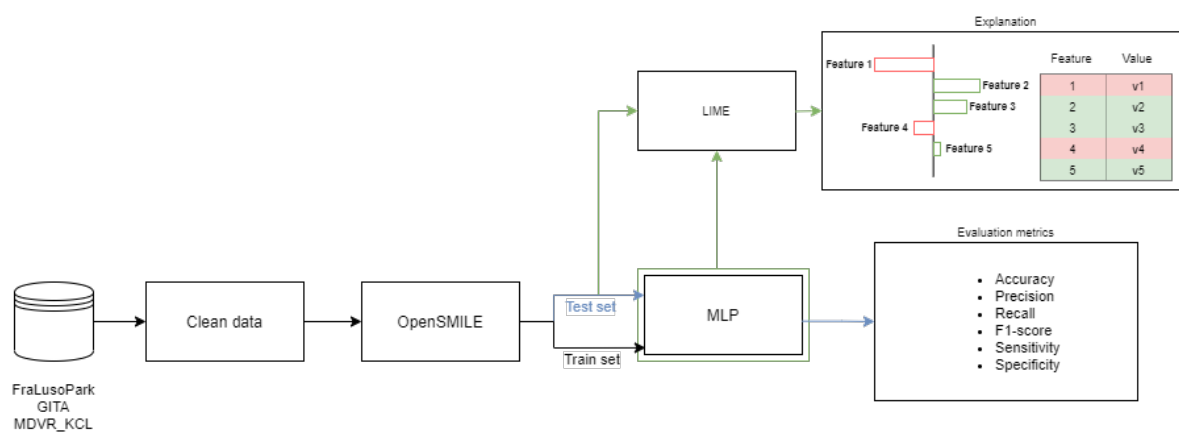


Figure 4.1: Pipeline of the proposed model.

5

Results and discussion

5.1 *Classification Experiments*

For this work, three types of experiments have been conducted, each using two different architectures, as described in the previous chapter. Results are shown in tables 5.1 and 5.2 (for the baseline experiments), 5.3 and 5.4 (for the semi-independent experiments), and 5.1 and 5.6 (for the language independent experiments). These tables show the top five MLP parameter configurations for each of the experiments. Tables 5.1, 5.3 and 5.5 present the results for architecture 1, whereas tables 5.2, 5.4 and 5.6 communicate the results for architecture 2.

5.1.1 Baseline experiments

Tables 5.1 and 5.2 show that both architecture 1 and 2 of the MLP yielded an accuracy of 90% with the best parameterization.

- Avaliar cada up dos parametros

The distribution between MLP solvers (adam and lbfgs) on the top 5 model configurations for architecture 1 is well balanced, whereas 4 out of the 5 (or 80%) of the best model configurations on architecture 2 use the adam solver.

- Avaliar as outras métricas

Architecture 1 yielded precision values between 0.75 and 1, meaning that 75% to 100% of the patients labeled as PD by the models were correctly classified. Architecture 2 produced slightly worst results regarding precision, achieving values between 67% and 100%. Recall values (which evaluates how the percentage of PD patients were correctly classified) were similar between the two architectures Architecture 1 produced recall values between 71% and 100%, whereas architecture 2 achieved values between 67% and 100%. Using the specificity metric (which allows to evaluate the percentage of HC patients that were correctly classified) to compare the two architectures, architecture 2 outperformed architecture 1 by a small margin, producing a range of values between 75% and 100%, whereas architecture 1 produced a range of values between 80% and 100%. Finally, comparing both architectures using the F1-score metric, it is possible to see a significantly higher performance with architecture 2, which produced a maximum of almost 91%, compared with architecture 1 that returned a maximum of 85%. We can conclude that there are no significant differences between the two architectures.

Semi independent

- Obtivemos Z com um modelo semi independente

- A performance baixou, manteve-se/não se manteve ao nível do state-of-the art

- Avaliar as outras métricas

- Resultados semelhantes à baseline provam que podemos retrainar um modelo com um novo pequeno dataset e o modelo fica preparado para uma classificação noutra idioma

dataset	solver	alpha	iterations	accuracy	precision	recall	specificity	f1-score
gita	adam	0.0001	5000	0.9	0.75	1.0	0.857	0.857
gita	lbfgs	0.0001	2000	0.9	0.75	1.0	0.857	0.857
gita	adam	0.001	2000	0.8	1.0	0.75	1.0	0.857
gita	lbfgs	0.01	5000	0.8	0.833	0.833	0.75	0.833
gita	lbfgs	0.01	2000	0.8	1.0	0.714	1.0	0.833

Table 5.1: Baseline experiment results using architecture 1.

dataset	solver	alpha	iterations	accuracy	precision	recall	specificity	f1-score
gita	adam	0.001	5000	0.9	0.8	1.0	0.833	0.889
gita	lbfgs	0.001	5000	0.9	1.0	0.833	1.0	0.909
gita	adam	0.0001	5000	0.8	0.8	0.8	0.8	0.8
gita	adam	0.01	5000	0.8	0.667	0.667	0.857	0.667
gita	adam	0.001	2000	0.8	1.0	0.778	1.0	0.875

Table 5.2: Baseline experiment result using architecture 2.

Independent

- Com o modelo 100% independente obtivemos XXX de performance
- A performance atingida foi semelhante à do estado da arte
- Avaliar as outras métricas

- Avaliar correlação dos parametros da rede com os resultados Figures 1 and 2 compare the different values of each parameter of the MLP with the accuracy yielded by the model.

- Overall, o modelo X/Y apresentou uma performance superior ao outro

- Overall de quantos doentes fomos capazes de diagnosticar

5.2 Language Independency

-

dataset	solver	alpha	iterations	accuracy	precision	recall	specificity	f1-score
mdvr_kcl + gita	adam	0.001	2000	0.9	0.857	1.0	0.75	0.923
fralusopark + gita	lbfgs	0.0001	5000	0.9	0.875	1.0	0.667	0.933
gita + fralusopark	adam	0.0001	2000	0.857	0.833	1.0	0.5	0.909
gita + fralusopark	adam	0.01	5000	0.857	0.889	0.889	0.8	0.889
gita + fralusopark	lbfgs	0.001	2000	0.857	1.0	0.75	1.0	0.857

Table 5.3: Semi independent experiment result using architecture 1.

dataset	solver	alpha	iterations	accuracy	precision	recall	specificity	f1-score
mdvr_kcl + gita	lbfgs	0.001	5000	0.9	1.0	0.8	1.0	0.889
fralusopark + gita	lbfgs	0.0001	2000	0.8	0.75	0.75	0.833	0.75
mdvr_kcl + gita	adam	0.0001	5000	0.8	1.0	0.667	1.0	0.8
mdvr_kcl + gita	adam	0.001	5000	0.8	0.6	1.0	0.714	0.75
mdvr_kcl + gita	lbfgs	0.0001	5000	0.8	0.6	1.0	0.714	0.75

Table 5.4: Semi independent experiment result using architecture 2.

dataset	solver	alpha	iterations	accuracy	precision	recall	specificity	f1-score
gita	lbfgs	0.001	5000	0.67	0.644	0.76	0.58	0.697
gita	lbfgs	0.01	2000	0.65	0.612	0.82	0.48	0.701
gita	lbfgs	0.001	2000	0.65	0.615	0.8	0.5	0.696
gita	lbfgs	0.0001	2000	0.63	0.592	0.84	0.42	0.694
gita	adam	0.0001	5000	0.63	0.6	0.78	0.48	0.678

Table 5.5: Independent experiment result using architecture 1.

dataset	solver	alpha	iterations	accuracy	precision	recall	specificity	f1-score
gita	lbfgs	0.01	5000	0.66	0.614	0.86	0.46	0.717
gita	lbfgs	0.0001	2000	0.63	0.589	0.86	0.4	0.699
gita	lbfgs	0.0001	5000	0.62	0.579	0.88	0.36	0.698
gita	lbfgs	0.001	2000	0.6	0.571	0.8	0.4	0.667
fralusopark	lbfgs	0.0001	5000	0.586	0.586	0.773	0.369	0.667

Table 5.6: Independent experiment result using architecture 2.

5.3 *Explanations*

-

6

Conclusions

Future work: - Add meta features, such as gender, that condition the normal value of a feature (such as F0) - Extend this work to generate global explanations at the same time - Test simpler features, such as log-filter bank (instead of MFCC) and compare accuracies. These would be easier to explain

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I Appendices

Glossary

- AD** Alzheimer's Disease. 1, 7
- AI** Artificial Intelligence. 1, 9
- ARD** Automatic Relevance Determination. 6, 7
- BC** Boosting Classifier. 7
- CAV** Concept Activation Vectors. 11
- DDK** Diadochokinesia. 5, 6, 15, 16
- DeepLIFT** DeepLIFT. 12
- DL** Deep Learning. 9
- DMLP** Deep Multi-Layer Perceptrons. 8
- DNN** Deep Neural Networks. 8, 10, 12
- DT** Decision Trees. 7
- F0** Fundamental Frequency. 3, 4, 6, 7, 17
- GMM-UBM** Gaussian Mixture Model - Universal Background Model. 5–8
- GNR** Glottal-to-Noise Ratio. 4, 6
- GPC** Gaussian Process Classification. 5–7
- H&Y** Hoehn and Yahr. 4
- HC** Healthy Controls. 3, 5–7, 15–18
- HNR** Harmonics-to-Noise Ratio. 4, 6, 17
- IG** integrated Gradients. 10
- KNN** K-Nearest Neighbours. 7, 8
- LIME** Local Interpretable Model-agnostic Explanation. 9, 12
- LR** Logistic Regression. 7, 8
- LRP** Layer-wise Relevance Propagation. 9, 12
- MDS** Movement Disorder Society. 4
- MFCC** Mel-frequency cepstral coefficients. 6, 7
- ML** Machine Learning. 1, 4–9, 11, 13
- MLP** Multi-Layer Perceptrons. 7, 17
- NAM** Neural Additive Models. 11
- NHR** Noise-to-Harmonics Ratio. 4, 6
- NN** Neural Networks. 6, 7, 11, 12
- PD** Parkinson's Disease. i, 1–9, 13, 15–18

PLDA Probabilistic Linear Discriminant Analysis. 6

PLP Perceptual Linear Predictive. 17

Rasta-PLP Rasta-Perceptual Linear Predictive. 6

RF Random Forests. 6–8

RISE Randomized Input Sampling for Explanation. 10, 12

SVM Support Vector Machines. 6, 7

TDU Text-dependent Utterances. 5, 6

UPDRS Unified Parkinson’s Disease Rating Scale. 4

VTI Voice Turbulence Index. 4, 6

XAI Explainable Artificial Intelligence. 2, 9

	Male	Female
F0 (Hz)	105-160	175-245
Jitter (%)	< 1.04	
Shimmer (%)	< 3.81	
HNR (dB)	< 20 (/a/, /i/), < 40 (/u/)	

Table 1: Feature values for healthy subjects.

Description	
F0	The number of open/close cycles of the glottis.
Jitter	Measures the frequency variation between cycles. Affected by lack of control on the vocal cords vibration.
Shimmer	Measures the amplitude variation between cycles.
HNR	The ratio between periodic (associated with normal speed production) and non-periodic (associated with noise) speech components.
MFCC	Features that approximate to our perception of the audio quality
PLP	Features that approximate to our perception of the audio quality

Table 2: Acoustic features description.