

Report Biomarkers

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Abstract

This study, which focuses on Parkinson's disease, aims to investigate the significance of left-right lateralization based on dopamine values using DAT-SPECT imaging data and their correlation with clinical symptoms in Parkinson's disease patient (PD) and healthy controls (HC). DAT-SPECT (Dopamine Transporter Single Photon Emission Computed Tomography) is an imaging technique primarily used to visualize and measure the density of dopamine transporters in the brain. Indeed, this technique is particularly useful in the diagnosis and evaluation of neurodegenerative diseases, such as Parkinson's disease and other forms of parkinsonism. The dataset, used to perform the analysis, consists of variables relating to demographic data, DAT-SPECT scores and finally the Neuropsychological (NP) tests which are a combination of clinical, neurological, and imaging test since there is no specific test for the diagnosis of Parkinson's disease. In order to investigate whether the lateralization is related to symptoms, a linear regression approach was used. This analysis verified that the lateralization is highly related to symptoms in PD patients. The lateralization of dopamine function could be useful in the future as a possible biomarker of PD symptoms severity.

ADD RESULTS

1 Introduction

Parkinson's disease is a chronic neurodegenerative disorder of the central nervous system primarily affecting motor control. It is characterized by the progressive loss of nerve cells in the brain region called the substantia nigra, which is responsible for producing a neurotransmitter called dopamine. Dopamine deficiency leads to motor symptoms such as tremors, muscle rigidity, bradykinesia (slowness of movement), and postural instability. In addition to motor symptoms, Parkinson's disease can cause a wide range of non-motor symptoms including sleep problems, depression, anxiety, fatigue, and cognitive difficulties. While the exact cause of Parkinson's disease is not fully understood, it is believed to result from a combination of genetic and environmental factors [1]. Currently, there is no cure for Parkinson's disease, but

there are treatments available to manage symptoms and improve the patients' quality of life. However, studies conducted in recent years have shown that the lateralization of brain dopamine in Parkinson's disease (PD) is a significant and distinctive aspect of the pathology. In particular, during its development, the degeneration of dopaminergic neurons in the substantia nigra is not uniform. Generally, one side of the brain is more affected than the other, leading to marked asymmetry. This marked asymmetry is visible not only in motor symptoms, such as more pronounced tremors on one side of the body and greater use of one hand over the other, but it can also be evident in non-motor disturbances, including cognitive issues for example in the sleep behavior and or sense of smell [5]. In previous studies, the connection between dopamine lateralization and symptoms has already been examined. For example, in [4], it

resulted that the motor symptoms on the less affected side were more correlated to striatal DAT binding.

Therefore, the aim of this study is to further investigate this idea, analyzing a dataset containing information about patients' motor and cognitive symptoms and also their lateralization data obtained from DAT-SPECT scans. The dopamine degeneration is explored in three different Regions of Interests (ROIs) in the brain, divided in right and left: Caudate, Putamen and Putamen Anterior. This analysis aims to investigate the relationship (if present) between the lateralization of dopamine function in these brain areas and the symptoms showed by Parkinson's patients, and relationship (if present) between the lateralization of dopamine function in these brain areas and possible covariates shown by healthy controls. To do so, the group checked also if there was a difference in lateralization between HC and PD.

Agreeing with what found in [4], the group expected to find a strong relation between the dopamine lateralization and the motor symptoms. The lateralization was expected to be significantly different between the two groups of HC and PD. Also the lateralization in different ROIs was expected to be related: a high lateralization in the Caudate was expected if both Putamen and Putamen Anterior were highly lateralized.

2 Material and Methods

2.1 Dataset description

The dataset used in this study was composed of 1556 subjects, of which 256 healthy controls and 1300 PD patients with different levels of symptoms' severity. Not all the variables given were interesting for the study. For example, all the data regarding the MRI acquisitions were discarded, because the analysis focused on the data from the DAT-SPECT scans, which were Striatal Binding Ratios (SBR) of the three ROIs (right and left sides), whether the scan was completed and its quality based on visual interpretation, as well as information regarding the date

of the scan and the injection.

The dataset included a part of demographics data, like age, ethnicity, gender, family members affected by PD, height, weight and dominant hand. A total of 967 and 589 patients were males and females, respectively. The average age in the baseline was 62.69 ± 10.11 years (range [29.3–86.5]).

There was then the data regarding the neuropsychological assessments of the patients using the Movement Disorder Society (MDS)-sponsored revision of the Unified Parkinson's Disease Rating Scale (MDS-UPDRS). The test is divided in 4 parts, concerning different classes of motor and cognitive symptoms, as specified in the PPMI Program Protocols (THE PARKINSON'S PROGRESSION...). Each answer could range from 0 (normal) to 4 (severe), and for each part there was a summing-up variable, that contained the sum of all the scores of that section. These variables were called NP1PTOT, NP1RTOT, NP2TOT, NP3TOT, NP4TOT. Some motor symptoms, especially in the third part, were divided in right and left, for example, the severity of tremor of the right or left upper limb. This was useful to investigate the different relation with the dopamine function lateralization.

Più aggiungere forse grafici con distribuzioni variabili iniziali, tipo grafici lateralizzazione

2.2 Cleaning e preprocessing

The first step was dividing the dataset in patients (PD, SWEDD, Prodromal) and HC.

In the preprocessing of the data, the patterns of missing values were analyzed, to determine the best course of action, through the VIM package in R.

In the HC, as shown in figure 1, GENETICS and FAMILIARITY are mostly empty fields, due to the nature of these variables: for healthy subjects, the chance of having relatives with a PD diagnosis and of having genetics history related to PD are low. On the opposite, for PD patients (figure 2), GENETICS is empty at the TOT %, while FAMILIRITY has the TOT % of

NaNs. The reason for this missingness, is probably that patients didn't know if their relatives had history of Parkinson's, or they didn't take the genetics test.

In the complete dataset, NP4TOT is missing at the 73.9%; nearly all the HC didn't take the fourth part of the test, and also the participation of the PD was really low. This suggests that the fourth part of the test was taken by a few subjects. For this reason, the NP4 variables were not taken into account in the statistical analysis.

For all the test-related variables (NP1PTOT, NP1RTOT, NP2TOT, NP3TOT), both in HC and PD, if a subject is missing a value for one test, also the other tests' results are missing. The pattern of missing data is considered to be MAR (Missingness At Random). For the statistical analysis, all the subjects that

were missing at least one test result weren't considered.

For ETHNICITY, SEX, and HAND (dominant hand of the subject), there aren't missing values in the whole dataset. In the ENROLLAGE variable, the missing values were calculated using the DAT-SCAN date and the birth date of the subject.

Regarding HEIGHT and WEIGHT, the pattern of missing data is MAR, with a total of TOT % of NaNs.

The variable PRIM_DIAG described the primary diagnosis; the group found some subjects with the value of 97, that wasn't explained by the data dictionary, therefore it was removed both from the HC and from the PD.

In figures 3 and 4, all the combinations of missing values of the dataset have been represented, using the VIM package in R.

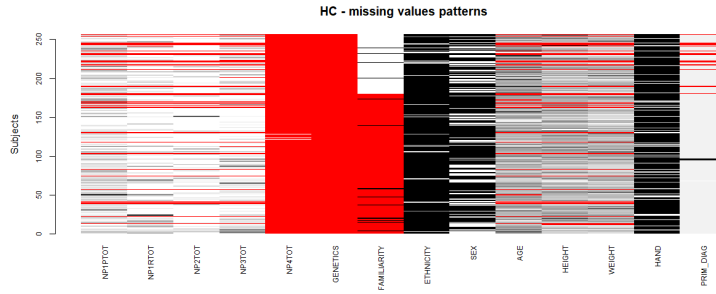


Figure 1: Pattern of missing values in HC

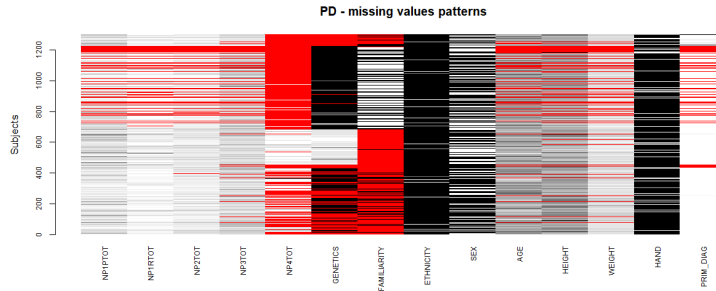


Figure 2: Pattern of missing values in PD

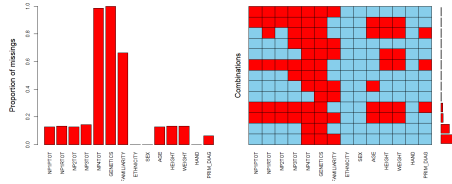


Figure 3: Missing values in HC

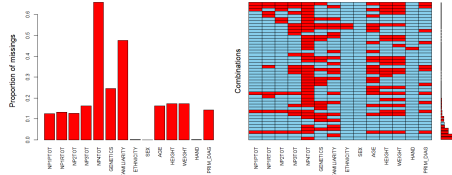


Figure 4: Missing values in PD

Another reason of exclusion from the analysis, was having an incomplete DAT-SCAN, a scan with poor quality, or, just for HC, a MoCA (Montreal Cognitive Assessment - stored in the variable MCATOT), score lower than 26. This test assesses the cognitive impairment, and, under 26, the subjects couldn't be considered healthy.

The preprocessing step ended with the removal of some outliers from ENROLLAGE, HEIGHT and WEIGHT. (DA CONTROL-LARE)

Concerning the demographics data, the two datasets were compared, to see if they had the same distributions. Through the Anova test, the average values of age, height and weight were found identical for the two groups. The proportion of male and female was comparable: in the whole dataset, the proportion of females was 37,85%, while in PD was 38,5% and in HC 34,37%.

The distribution of ETHNICITY was also found similar between the two groups, while FAMILIARITY and HAND were significant different in the two populations when analyzed through a Wilcoxon test. The differences in FAMILIARITY can be explained by the connection between PD patients and the genetic components in the disease: for a PD patient, it's more likely to have some relatives affected by PD, than for an healthy subject.

2.3 Feature extraction

In order to perform the statistical analysis to answer the study questions, the relevant features were extracted.

From the lateralization data, the group used a DAT binding asymmetry index found in [3]: $(\text{right} - \text{left SBR}) / (\text{right} + \text{left SBR})$. The lateralization was considered relevant when the asymmetry index was higher than 20%. This threshold level was adopted from a previous study [2]. In this way, the group identified, for the Caudate, 28 PD patients with a right lateralization and 29 with a left lateralization. For the Putamen, 203 PD had a right lateralization, 2 HC and 150 PD with a left lateralization. For the Putamen Anterior, 113 PD patients were found lateralized on the right side, while 64 on the left. The remaining subjects were not lateralized.

DIVIDE SYMPTOMS - COVARIATES (demographics data)

Through the Pearson pairwise correlation, the group analysed the correlation coefficients of the relevant variables, such as ENROLLAGE, WGTKG (weight - kg), HGTCM (height - cm), and MCATOT, with the lateralization index (figures 5 and 6). For the analysis, were kept only the significant ones, with p-value < 0.05 , and correlation coefficient higher than 0.75. This threshold was chosen in order to have a

	Covariates
HC	NP1ANXS, NP1APAT, NP1COG, NP1HALL, NP1LTHD, NP1PAIN, NP1PTOT NP1PTOT.1, NP1RTOT, NP1URIN, NP2DRES, NP2HOBB, NP2HWRT NP2PTOT, NP2RISE, NP2SALV, NP2SPCH, NP2SWAL, NP2TRMR, NP2TURN NP3BRADY, NP3FACXP, NP3FTAPL, NP3FTAPR, NP3HMOVR, NP3KTRMR NP3PRSPR, NP3PSTBL, NP3PTRML, NP3RIGLU, NP3RIGRL, NP3RIGRU NP3RISNG, NP3RTALU, NP3TTAPL, WGTKG
PD	NP1ANXS, NP1COG, NP1SLPN, NP2DRES, NP2FREZ, NP2HYGN, NP2PTOT NP2RISE, NP2SWAL, NP2WALK, NP3FACXP, NP3GAIT, NP3HMOVR NP3KTRML, NP3PTRML, NP3PTRMR, NP3RIGLL, NP3RTALU, NP3RTARL NP3RTARU, NP3SPCH, NP3TOT, WGTKG

Table 1: Table of the significant covariates for HC and PD

strong correlation (0.6-0.79) and a good trade-off between selecting significant covariates and not having too many of them. With a lower cor-

relation threshold, the number of covariates was too high. Therefore, the table 1 was obtained, listing the significant covariates of HC and PD.

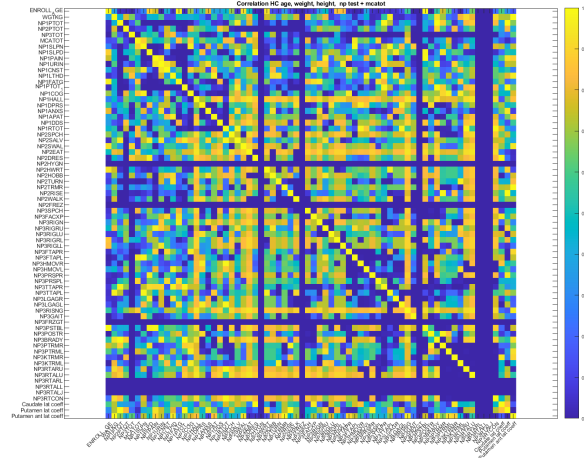


Figure 5: Correlation matrix with the lateralization index - HC

In the correlation matrix of the HC in figure 5, the rows and columns with correlation coefficients all equal to -1 are due to the NaN values.

CONFRONTO PER VARIABILI DIS-CRETE

2.4 Statistical methods

At first, the group compared HC and PD populations in terms of lateralization, using an

Anova test. The Anova test was chosen because [forse Wilcoxon]?????????????

Doing so, the group investigated the presence of a difference between the lateralization in HC with the one in PD. To perform the analysis, the lateralization was taken in absolute value. The choice of using the absolute value came from the interest on the amount of lateralization, not its direction.

As hypothesized at the start of the study, the

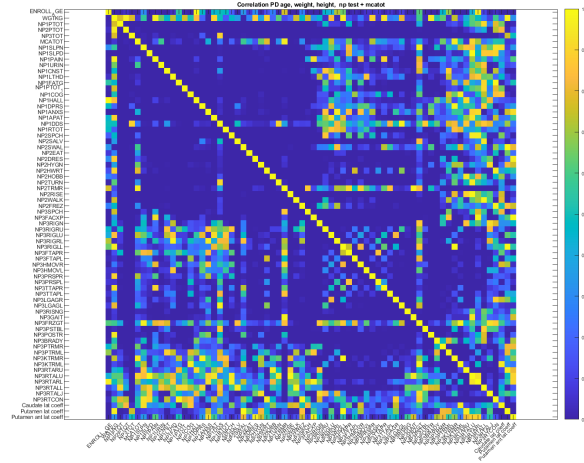


Figure 6: Correlation matrix with the lateralization index - PD

analysis confirmed that there was a significant difference (p-value < 0.01) in the lateralization of HC and PD in all the ROIs.

As parameter, the adjusted R^2 was used to get the true value of the fit.

ALTRI METODI USATI

2.4.1 Linear regression

Using the covariates found for PD and HC through the correlation matrix, the group performed a linear fit through the MATLAB function `fitlm` (the lateralization was not taken in absolute value, in order to have a relation with its direction).

3 Results

As an answer to the first research question, which was to investigate the relationship between dopamine lateralization function and symptoms, the group found the following statistical results.

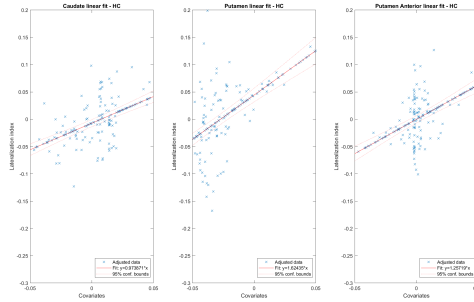


Figure 7: Linear fit - HC

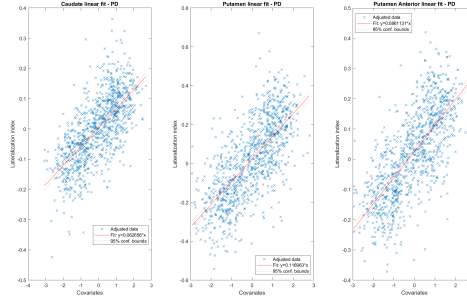


Figure 8: Linear fit - PD

In figures 7 and 8, the linear fits are shown for HC and PD. The graphs contain on the x-axis the covariates, while on the y-axis the lateralization index, obtained in the section of Feature Extraction. Each figure shows the fit for

the three different ROIs, Caudate, Putamen and Putamen Anterior.

The numerical results of these linear fits are shown in the tables below, with R^2 (adjusted) and p-value for each ROI.

CAUDATE	HC	PD
R^2 (adjusted)	0.021899285491766	0.441924423381424
p-value	0.398349585769054	1.950516939619888e-20
PUTAMEN	HC	PD
R^2 (adjusted)	0.027100337935025	0.481441593928543
p-value	0.375829445576761	1.749319862864852e-24
PUTAMEN ANT	HC	PD
R^2 (adjusted)	-0.048170586150496	0.511431631643848
p-value	0.705855414513558	6.121704939031713e-28

The second research question was whether a relationship between dopamine lateralization function and possible covariates shown by healthy controls was present. To answer this question, the following results were found.

- A clear and concise description of the statistical results providing answers to the research questions
- A sensitivity analysis of the results to covariates, group matching and data quality (e.g. missing data, data miss balance)

4 Discussion

As shown in figures 7 and 8, and from the values of R^2 and p-value, the group understood

that the linear fit wasn't appropriate to describe the lateralization of HC, because there wasn't any significant lateralization, nor any symptoms. The image of the fit is used to show that the results are as expected. On the other hand, for PD patients, the values of R^2 and p-value are highly significant, showing that the lateralization is related to the symptoms, as hypothesized.

HC differenza significativa con PD grazie all'indice j_{20}

- Direct answers to the research questions
- An overview of the limitations of the study
- A list of possible suggestions to improve the study in case someone will repeat it in future

References

- [1] Janice M. Beitz. “Parkinson’s disease: a review.” eng. In: *Frontiers in bioscience (Scholar edition)* 6.1 (Jan. 2014). Place: Singapore, pp. 65–74. ISSN: 1945-0524 1945-0516. DOI: 10.2741/s415.
- [2] Eleonora Fiorenzato et al. “Asymmetric Dopamine Transporter Loss Affects Cognitive and Motor Progression in Parkinson’s Disease”. en. In: *Movement Disorders* 36.10 (Oct. 2021), pp. 2303–2313. ISSN: 0885-3185, 1531-8257. DOI: 10.1002/mds.28682. URL: <https://movementdisorders.onlinelibrary.wiley.com/doi/10.1002/mds.28682> (visited on 05/31/2024).
- [3] Valtteri Kaasinen. “Ipsilateral deficits of dopaminergic neurotransmission in Parkinson’s disease”. en. In: *Annals of Clinical and Translational Neurology* 3.1 (Jan. 2016), pp. 21–26. ISSN: 2328-9503, 2328-9503. DOI: 10.1002/acn3.268. URL: <https://onlinelibrary.wiley.com/doi/10.1002/acn3.268> (visited on 05/31/2024).
- [4] Walter Pirker. “Correlation of dopamine transporter imaging with parkinsonian motor handicap: How close is it?” en. In: *Movement Disorders* 18.S7 (Oct. 2003), S43–S51. ISSN: 0885-3185, 1531-8257. DOI: 10.1002/mds.10579. URL: <https://onlinelibrary.wiley.com/doi/10.1002/mds.10579> (visited on 05/31/2024).
- [5] P. Riederer et al. “Lateralisation in Parkinson disease”. en. In: *Cell and Tissue Research* 373.1 (July 2018), pp. 297–312. ISSN: 0302-766X, 1432-0878. DOI: 10.1007/s00441-018-2832-z. URL: <http://link.springer.com/10.1007/s00441-018-2832-z> (visited on 05/31/2024).