Involvement of specific executive functions in mobility in Parkinson’s disease

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Abstract

Postural instability and gait disorders (PIGD) in Parkinson’s disease (PD) seem to be associated with executive dysfunction. We investigated which specific executive functions are associated with functional mobility in mildly affected PD patients. Functional mobility (Timed Up&Go Test, TUG), PIGD score, (spatial) working memory, set shifting, response inhibition, and response generation were assessed in a large cohort of 232 non-demented PD patients. Both performance on the TUG and PIGD score were weakly associated with working memory and response generation (semantic and phonemic fluency). TUG also correlated with semantic fluency when corrected for disease severity and age. These results indicate that response generation and working memory are associated with (and possibly also causally related to) gait and balance deficits. In order to fully interpret gait and postural stability of PD patients in everyday situations, the role of impairments in working memory and response generation should be taken into account.

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1. Introduction

Parkinson’s disease (PD) is characterized by its motor features including gait difficulty and postural instability. Moreover, already in the early stages of PD some 20% of patients have mild cognitive impairment [1]. Because of the underlying neurodegenerative nature of PD, cognitive impairments are overall related to increased motor severity [2]. For example, the motor subtype with predominantly posture and gait disorders is a strong predictor of severe cognitive decline [3]. However, associations between more specific aspects of cognitive function and motor impairments are less clear [2].

With respect to cognitive domains, the executive functions are particularly affected in PD. Executive deficits can hamper activities in everyday life in PD for various reasons. First, activities of daily living can be affected directly because of an inability to organize, shift, monitor and play. In addition, executive dysfunction can impair daily-life performance more indirectly, via a detrimental effect on motor function. Specifically, there is increasing evidence to suggest that executive functions play an important role in gait and postural adjustments [4]. For example, even healthy individuals without cognitive deficits reduce their walking speed and take smaller steps when they must perform a secondary cognitive task while walking, suggesting that executive or cognitive control is required for seemingly automatic functions like walking.

The results from such dual-task studies have consistently shown effects on various gait variables, in particular walking speed, stride length and step-to-step variability. However, it has not been clarified which specific aspects of executive function are important in relation to impairments in gait and balance. Here, we aimed to further clarify the association between functional mobility (Timed Up&Go Test), posture instability and gait disorders (PIGD), and four main aspects of executive function [5,6]: updating/working memory, set shifting, response inhibition, and response generation in a large cohort of non-demented PD patients.

2. Methods

2.1. Participants

Our study sample was a subsample of the ParkFit study population [7]. Baseline assessment of cognitive functions and mobility measures are presented here. Inclusion criteria were PD (diagnosed according to the UK Brain Bank criteria), age between 40 and 75 years, a sedentary lifestyle, Hoehn & Yahr (H&Y) ≤ 3, and Mini-
Mental State Examination (MMSE) ≥24. The study was approved by the regional medical ethical committee (CMO region Arnhem-Nijmegen) and patients gave their written informed consent. The present analysis is limited to patients who completed all executive function and mobility tests (N = 232, 66% men, 64.4 ± 7.9 years). Mean Unified Parkinson’s Disease Rating Scale-III (UPDRS-III) score was 33.4 ± 9.1 and mean MMSE score was 28.1 ± 1.6. Almost 80% of patients (n = 183) was in H&Y stage 2; the other patients had H&Y stage 1 (n = 3; 1%), 1.5 (n = 6; 3%), stage 2.5 (n = 35; 15%) or stage 3 (n = 5; 2%). Most patients (47%) scored category 3 for their level of education (range 1 = no education to 6 = university).

2.2. Materials and procedure

The Timed Up&Go (TUG) test was used as an index of mobility [8]. In this test the patient has to stand up from a chair, walk 3 m at comfortable speed, turn 180°, walk back to the chair and sit down again. The sum score of items 27–30 of the UPDRS-III (arising from chair, posture, gait, postural stability) was used to calculate PIGD score.

Upgrading/working memory was examined using the Spatial Working Memory (SWM) subtest of the Cambridge Neuropsychological Test Automated Battery (CANTAB) [9]. In this computerized task, participants have to search for a hidden token by clicking a number of boxes that are presented in a spatial layout. After finding a token, participants have to search for a new token that is hidden in one of the other boxes. Within-search errors occur if a participant returns to a previously ‘opened’ box within a search, whereas between-search errors occur if a participant returns to a box that already contained a token in a previous search. Also, a strategy index reflects the efficiency of the search path.

Set shifting was assessed using the Intradimensional/Extradimensional (ID/ED) Set Shift test from CANTAB [9]. Here, participants have to learn a sorting rule by clicking stimuli that differ in different dimensions (shapes and lines) using feedback. After six correct consecutive responses according to the to-be-learned rule, the rule changes and participants have to learn the new sorting rule. Outcome measures were the number of stages completed and the number of errors made (adjusted for the number of stages completed).

Response inhibition was measured using an auditory Stroop paradigm [10], which allowed for precise recording of reaction times per response (in contrast to the widely-used paper-and-pencil Stroop Color-Word Test). In this task, patients hear the words “high” or “low” spoken at a high or low tone, every 2 s. Participants were instructed to respond as fast as possible by repeating the tone of the stimulus. Verbal reaction time and accuracy were combined in a composite score (accuracy/verbal reaction time).

Response generation was measured by the ability to access long-term memory using either a phonological cue (letter fluency; naming as many words as possible starting with the letter “M” in 1 min) or a semantic cue (semantic fluency; naming as many animals in 1 min) [11].

Individual performance on SWM, ID/ED and fluency were compared to age and/or education or IQ corrected available normative data for the CANTAB (n = 2000) [12] and the fluency tests (n = 1856) [11]. An individual performance was classified as impaired if the individual score was more than 1.65 SD below the normative mean (i.e., below the 5th percentile) [13]. No normative data were available for the Stroop paradigm.

### Table 1

<table>
<thead>
<tr>
<th>Test (N = 232)</th>
<th>Outcome measure</th>
<th>Test performance</th>
<th>Univariate regression with TUG</th>
<th>Univariate regression with PIGD</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Mean ± SD</td>
<td>% impaired*</td>
<td>Beta</td>
</tr>
<tr>
<td><strong>Updating/working memory</strong></td>
<td></td>
<td></td>
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</tr>
<tr>
<td>SWM a</td>
<td>Within-search errors</td>
<td>2.91 ± 4.36</td>
<td>4</td>
<td>0.032</td>
</tr>
<tr>
<td></td>
<td>Between-search errors</td>
<td>43.37 ± 20.97</td>
<td>4</td>
<td>0.274*</td>
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<tr>
<td></td>
<td>Strategy</td>
<td>35.63 ± 5.41</td>
<td>4</td>
<td>0.206*</td>
</tr>
<tr>
<td><strong>Set shifting</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>ID/ED</td>
<td>Stages completed</td>
<td>7.52 ± 2.00</td>
<td>17</td>
<td>−0.041</td>
</tr>
<tr>
<td></td>
<td>Total errors (adjusted)</td>
<td>54.87 ± 46.14</td>
<td>18</td>
<td>0.051</td>
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<tr>
<td><strong>Inhibition</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Auditory Stroop</td>
<td>Composite score</td>
<td>1.48 ± 1.72</td>
<td>NA</td>
<td>0.049</td>
</tr>
<tr>
<td><strong>Response generation</strong></td>
<td></td>
<td></td>
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<tr>
<td>Fluency</td>
<td>Phonemic</td>
<td>11.51 ± 4.66</td>
<td>6</td>
<td>−0.198*</td>
</tr>
<tr>
<td></td>
<td>Semantic</td>
<td>18.00 ± 5.71</td>
<td>26</td>
<td>−0.340*</td>
</tr>
</tbody>
</table>

*Significant regression coefficients (p < 0.05).

a Spatial Working Memory.

b Intradimensional/Extradimensional Set Shift.

c Timed UpGo Test.

d Postural Instability and Gait Disorders.

e Impaired performance was defined as more than 1.65 SD deviation below the normative mean. For the auditory Stroop test, no normative values were available.

2.3. Data analysis

To test the associations between performance on the TUG and PIGD score and the performance on cognitive tests, univariate regression coefficients were calculated using linear regression. Next, a multivariate linear regression model was constructed to predict TUG and PIGD using the significant variables from the univariate regression together with age. UPDRS-III score and educational level as independent variables. Significant contribution was accepted at p < 0.05.

3. Results

PD patients needed on average 9.51 ± 2.85 s to complete the TUG. Mean PIGD score was 2.3 (±1.3). Regression coefficients for the association between fluency tests and the TUG were significant, yet weak (beta between −0.198 and −0.340, Table 1). Similar beta values were observed for the association between Spatial Working Memory and the TUG regarding between-search errors and strategy. Other cognitive outcome measures were not correlated with the TUG. The linear regression analysis with PIGD score as dependent variable produced similar results.

A stepwise multivariate regression model was constructed by entering fluency tests and SWM between errors and strategy scores, together with UPDRS-III score, age and educational level. UPDRS-III (beta = 0.263, p < 0.001), age (beta = 0.212, p = 0.001), and semantic fluency (beta = −0.197, p = 0.002) contributed significantly to the model, together explaining 24% of the total variance of the TUG. Only UPDRS-III (beta = 0.469, p < 0.001) and age (beta = 0.186, p = 0.002) survived multivariate regression with PIGD as dependent variable. This model explained 31% of the total variance of PIGD score.

4. Discussion

In this study, we evaluated which of the four domains of executive function is involved in functional mobility in a large cohort of patients with PD. Spatial Working Memory and verbal fluency showed small but significant associations with both the TUG and PIGD scores. Moreover, semantic fluency was significantly associated with mobility, independent of age and severity of motor signs as measured with the UPDRS-III.

The association of response generation and working memory (updating) with the TUG can be explained as an involvement of executive control during this seemingly pure motor task. Ongoing movement requires continuous monitoring and updating in order to adjust to ongoing changes in the environment. Specifically, the
turning and transfer components of the TUG might demand executive processing. Alternatively, one could argue that processing speed underlies both executive functions and the TUG [14]. However, the Stroop task is presumably the most time-critical cognitive task in our design, but was not associated with performance on the TUG.

It is important to note that the patients in our sample were relatively mildly affected. The H&Y stages and UPDRS-III scores were low. This indicates that our research sample of PD patients probably had only minor gait difficulties and postural instability. With regard to the extent of executive dysfunction, impairments were present in set shifting (17–18%) and semantic fluency (26%), but not in working memory and phonemic fluency. However, even small decrements in executive function may affect motor function in more complex daily-life environments, which require more planning and switching than the TUG test which was performed under well-controlled circumstances in our study. Also, since PD progressively affects both cognitive and motor functions, the interaction between both domains might place PD patients in vulnerable everyday situations in more advanced disease stages.

The results from this study revealed that in non-demented PD patients with minor gait deficits, response generation and working memory are the executive functions that are weakly associated with functional mobility. With regard to clinical practice, we recommend that in order to fully interpret gait and postural stability of PD patients in everyday situations, the role of impairments in working memory and response generation, even when mild, should be taken into account.

Competing interests

All authors declare they have no competing interests for publication.

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References