Depression increases in patients with Parkinson’s disease according to the increasing severity of the cognitive impairment

Em pacientes com doença de Parkinson, a depressão aumenta de acordo com o aumento da gravidade do comprometimento cognitivo

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ABSTRACT

Objective: To test the hypothesis that severity of cognitive impairment modifies the association between depression and Parkinson's disease (PD). Method: One-phase population-based door-to-door surveys. This is a secondary analysis of 1,451 people aged 65 years and older with cognitive impairment living in defined catchment areas. Depression was estimated according to ICD-10, self-reported PD, disability according to WHODAS-II and cognitive status according to the CSI-D. Results: The mean age of the sample was 79.3 years old and most (69%) were women. Of the total sample, 16.1% had depression and it was significantly higher among participants with PD. There was an increase on the ORs of the association between depression and PD with decreased scores in the cognitive test (Adjusted OR from 0.98 to 8.04). Conclusion: The association between depression and PD increases with the severity of the cognitive impairment.

Keywords: parkinsonism, depression, cognitive impairment, Parkinson's disease.

RESUMO

Objetivo: Testar a hipótese de que a gravidade do prejuízo cognitivo modifica a associação entre depressão e doença de Parkinson (DP).

Método: Estudo populacional através da análise secundária de 1.451 pessoas com idade maior ou igual a 65 anos com prejuízo cognitivo que residiam em áreas de abrangência definidas. A depressão foi estimada de acordo com a CID-10, auto-relato de DP, incapacidade conforme a WHODAS-II e nível cognitivo de acordo com a CSI-D. Resultado: A média de idade foi 79,3 anos, predominaram as mulheres (69%). Do total de indivíduos, 16,1% tinham depressão, significativamente maior entre os participantes com DP. Houve aumento gradativo na OR relativa à associação entre depressão e DP com a diminuição do escore no teste cognitivo (OR ajustado variou de 0,98 a 8,04).

Conclusão: A associação entre depressão e DP parece aumentar com a gravidade do prejuízo cognitivo.

Palavras-chave: parkinsonismo, depressão, prejuízo cognitivo, doença de Parkinson.

Depression and dementia are common non-motor manifestations of Parkinson’s disease (PD). Prevalence estimates vary widely, but most studies have suggested that 40% to 50% of PD patients present clinically relevant depression¹, ² and 30% of them present dementia³.

Cognitive impairment itself increases the risk of depression⁴ and thus it is plausible that the presence of cognitive impairment might influence the association of depression with PD. Most studies investigating the association of depression and PD are based on convenience clinical samples and...
frequently exclude patients with dementia or cognitive impairment. Although both cognitive impairment and depression are frequent comorbidities of PD, little is known on how the association of depression and PD varies according to cognitive impairment.

We estimated the association between depression and PD in a large community sample of 1,451 elderly with cognitive impairment and tested the hypothesis that the strength of the association between depression and PD increases by severity of cognitive impairment even after adjusting for disability, a potential mediator of this association.

METHOD

This study is a secondary analysis of the 10/66 Dementia Research Group population-based surveys. Further details on the protocol can be found elsewhere.

Participants were recruited following informed consent. Studies were approved by local ethical committees as well as by the ethical committee at King’s College London.

Setting and sample: All residents aged 65 years and older from 11 geographically defined catchment areas in India, China, Cuba, Dominican Republic, Venezuela, Mexico, and Peru were interviewed (n=14,960) in one-phase population-based cross-sectional door-to-door surveys. The analysis presented in this study is restricted to those with cognitive impairment (n=1,451). Response rates were 80% and above in all sites. The same protocol was used in all countries allowing fair comparisons between sites.

Measurements

Parkinson’s disease: participants and their informants were asked if a doctor had ever diagnosed the participant with PD. Possible answers were: No, probable or certain. Only those who answered “certainly yes” were considered as “self-reported PD”.

Diagnosis of depression: In order to categorize depressed participants, International Classification of Diseases 10th revision (ICD-10) criteria were used. The ICD-10 divides depressive episodes into mild, moderate and severe, for this analysis we considered any current ICD-10 depressive episode, irrespectively of the severity. These were derived using a computerized algorithm applied to the geriatric mental state (GMS).

Cognitive impairment: the cognitive assessment was based on the “Community Screening Instrument for Dementia” (CSI-D), a 32 items cognitive test assessing naming, comprehension, memory, orientation, apraxia and constructional difficulty. It has shown very good psychometric properties for cross-cultural studies. The CSI-D population normative data for Latin America, India and China for the 10/66 Dementia Research Group has been previously detailed described elsewhere. Objective impairment was defined as a score 2 standard deviations (SDs) below the average obtained with the total sample (n=14,960) and was used to select the sample included in this analysis. Dementia was determined according to the DSM-IV and/or 10/66 dementia diagnosis.

Disability: It was measured using the 12-item interviewer-administered World Health Organization Disability Assessment Schedule II (WHODAS-II), developed by World Health Organization as a culture-fair assessment to use in cross-cultural studies.

Sociodemographic status: Age of all participants was established from participant and informant report and an official ID document; when the information was not clear or there was a discrepancy, an event calendar was used. Level of education and gender were also recorded during the interview.

Statistical analysis

We described the participants’ characteristics in participants with and without PD and estimated the association between depression and PD adjusted for socio-demographic characteristic and disability. Odds ratios (OR) are derived from logistic regression models with 95% confidence intervals (95% confidence interval (CI)). To explore how the association between depression and PD behaved according to levels of cognition we used the same model as before for each levels of cognitive impairment according to quartiles of scores on the CSI-D. In order to further understand how cognition affects the association between depression and Parkinson we repeat the same logistic regression models among people with and we have also conducted the same analysis among participants with and without dementia.

RESULTS

A total of 1,451 participants were classified as having cognitive impairment and were included in this analysis. The sample mean age was 79.3 (standard deviation (SD) 0.2) years old and most part of participants were women (69%). Overall, educational levels were low and 61.1% did not complete primary school. Levels of disability were high, 38.3% of the whole sample and 54.8% of PD patients reported 15 or more days loss due to disability in the last month. Table 1 shows participants’ characteristics according to the presence of PD. Groups with and without PD did not significantly differed regarding their mean age, gender, mean scores on cognitive tests and levels of education, but PD patients had much higher levels of disability. A total of 234 (16.1%) individuals fulfilled criteria for depressive episode according to ICD-10 and 42 (2.9%) reported having PD.
Depression was significantly more frequent among participants with PD, 28.5% (12) versus non-PD subjects 15.8% (222). Crude OR was 2.18 (95%CI 1.10-4.32) and after adjusting for age, gender, disability and years of education the association of PD with depression remained significant (OR 2.19, 95%CI 1.08-4.45).

Table 2 shows the association of PD with depression by quartiles of cognitive status (crude and adjusted for age, gender, level of education and functional disability). When divided by quartiles of cognitive status, PD was significantly associated with depression only among those most severely impaired (adjusted OR 8.04 95%CI 1.57-41.15). There was a clear increase in the OR values with the decrease in cognitive test scores (from 8.04 among the most impaired to 0.98 to the least impaired). We have also looked this association among participants with and participants without dementia, which shows a much stronger association between depression and PD among people with dementia (adjusted OR=4.44, 95%CI 1.80-10.93) than among people with cognitive impairment but without dementia (adjusted OR=0.69, 95%CI 0.18-2.55).

DISCUSSION

We studied the relationship between PD and depression in a community sample of 1,451 elderly subjects with cognitive impairment from seven different low and middle-income countries. We found a strong association between depression and PD, which was stronger among those with worse cognitive impairment and among those with dementia.

In our sample around one third of those with PD were depressed. There was a strong positive association of PD and depression even after adjusting for age, gender, levels of education and disability. In a recent systematic review, Reijnders et al. (2008) found that estimates on the prevalence of depression in PD vary according to study methodology, being lower in community based compared to clinical samples. These authors found a mean prevalence of 8.1% for studies assessing major depression in PD patients from the general population. We used a structured interview in a community sample of cognitive impaired elderly and found a much higher prevalence (28.5%), which could be explained by the fact that we had a cognitively impaired sample and we used ICD-10 for diagnosing depressive episode.

Small previous studies suggested that the association of depression and PD is stronger for those with cognitive impairment, but none looked at this association according to the severity of the cognitive impairment in a community sample. When we divided our sample by quartiles of levels of cognitive impairment we found that the association of depression with PD increased with increased cognitive impairment. Cognitive impairment itself increases the risk for depression irrespectively of other

<table>
<thead>
<tr>
<th>Impairment (by Quartiles)</th>
<th>OR (95% CI)</th>
<th>Multi-variable* adjusted OR (95% CI)</th>
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</thead>
<tbody>
<tr>
<td>Most impaired (1st quartile)</td>
<td>5.40 (1.24-23.54)</td>
<td>8.04 (1.57-41.15)</td>
</tr>
<tr>
<td>Intermediate Impaired (2nd quartile)</td>
<td>2.77 (0.65-11.90)</td>
<td>2.14 (0.45-10.26)</td>
</tr>
<tr>
<td>Less impaired (3rd quartile)</td>
<td>1.61 (0.42-6.12)</td>
<td>1.42 (0.33-6.15)</td>
</tr>
<tr>
<td>Least impaired (4th quartile)</td>
<td>1.32 (0.36-4.87)</td>
<td>0.98 (0.24-4.02)</td>
</tr>
<tr>
<td>With Dementia</td>
<td>3.78 (1.62-8.83)</td>
<td>4.44 (1.80-10.93)</td>
</tr>
<tr>
<td>Without Dementia</td>
<td>0.88 (0.24-3.03)</td>
<td>0.69 (0.18-2.55)</td>
</tr>
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OR: odds ratio; *adjusted by age, gender, level of education and functional disability.
diseases. It might be that either depression or cognitive decline are symptoms of a more severe presentation of PD or that the presence of depression leads to cognitive deterioration.

PD is the second most common neurodegenerative disorders after Alzheimer disease. Nevertheless overall, PD is a low prevalent condition and only few studies are based in community samples. One strength point of our study is that we used data from community-based surveys in seven countries. However, yet our sample included only 42 participants with PD, which brings some uncertainty to our estimates. Another limitation to be considered is that we used self-report to identify participants with PD, the same protocol was used in all sites. Also, diagnosis of PD is difficult to be ascertained and misdiagnosis is common even among specialists. Self-reported PD has been validated against post mortem neuropathological examination in a large prospective community based study, showing that self-reported PD is a good predictor of PD. Moreover the prevalence of PD we found is very similar to PD prevalence among other populations of older adults.

The cognitive deficits found in PD depressed patients can be reversible with remission of depressive symptoms, however large prospective studies are needed to test if treating depression in PD would reduce the likelihood of developing dementia. Our findings show that the strength of the association of depression with PD increases with the severity of cognitive impairment and is also stronger among people with dementia. It is difficult to disentangle how much of this association is due to the bad performance of people with depression in the cognitive tests, however it is still plausible to think that the prevalence of depression among people with PD might be underestimated if people with impaired cognition or dementia are excluded.

References