http://dx.doi.org/10.1590/1980-57642018dn13-010008

Montreal Cognitive Assessment scale in patients with Parkinson Disease with normal scores in the Mini-Mental State Examination

Krisly Arguedas Vásquez¹, Erick Miranda Valverde², Daniel Valerio Aguilar², Henri-Jacques Hernández Gabarain³

ABSTRACT. Several screening tests have been used for cognitive evaluation in Parkinson's disease (PD). **Objective:** To evaluate the usefulness of the Montreal Cognitive Assessment (MoCA) in patients with Parkinson's disease and no cognitive impairment complaints. **Methods:** A total of 40 PD patients with no complaints of cognitive problems were included. Patients were selected using the Mini-Mental State Examination (MMSE) and the MoCA was then administered. **Results:** 80% of patients exhibited Mild Cognitive Impairment (MCI) according to the MoCA. Statistically significant differences in visuospatial, attention and delayed recall functions were evident between the normal and abnormal MoCA groups. **Conclusion:** The study results suggest that MoCA may be a good screening test in patients with PD who do not present cognitive complaints. **Key words:** Parkinson's disease, mild cognitive impairment, cognitive impairment, Montreal Cognitive Assessment.

DETECÇÃO DE COMPROMETIMENTO COGNITIVO DE ACORDO COM A ESCALA MONTREAL COGNITIVE ASSESSMENT EM PACIENTES COM DOENÇA DE PARKINSON IDIOPÁTICA COM COGNIÇÃO NORMAL DE ACORDO COM O ESCORE NO MINIEXAME DO ESTADO MENTAL

RESUMO: Vários testes de triagem foram utilizados para avaliação cognitiva na doença de Parkinson (DP). **Objetivo:** Avaliar a utilidade da Avaliação Cognitiva de Montreal (MoCA) em pacientes com doença de Parkinson sem queixa de comprometimento cognitivo. **Métodos:** Um total de 40 pacientes com TP sem queixas de problemas cognitivos foram admitidos e com o Estado de Exame do Estado Mental Mini (MEEM) foram selecionados e receberam o MoCA. **Resultados:** 80% apresentaram dados de Comprometimento Cognitivo Leve (ICM) segundo o MoCA, sendo as funções visoespaciais, atenção e memória atrasada aquelas que apresentaram diferenças estatisticamente significantes entre os grupos MoCA normal e anormal. **Conclusão:** Este estudo sugere que o MoCA pode ser um bom teste de triagem em pacientes com DP que não apresentam queixas cognitivas.

Palavras-chave: doença de Parkinson, comprometimento cognitivo leve, comprometimento cognitivo, Montreal Cognitive Assessment.

Parkinson's disease (PD) is the second most common neurodegenerative disease after Alzheimer's disease (AD). As many as 20-30% of patients with Parkinson's disease (PD) have cognitive deficits within the range of Mild Cognitive Impairment (MCI) at the time of diagnosis, a condition that is very impor-

tant to detect since it is associated with an increased risk of developing dementia. ^{1,2} The typical profile of cognitive impairment in PD involves impairment in executive functions, attention, visuospatial and subcortical memory functions, recall, language preservation, and praxis. ^{3,4} Given this scenario, it is impor-

This study was conducted at the Hospital Carlos Luis Valverde Vega, Caja Costarricense de Seguro Social, Alajuela, Costa Rica.

'Geriatrician-Gerontologist Hospital Carlos Luis Valverde Vega, Caja Costarricense de Seguro Social, Alajuela, Costa Rica. 'Geriatrician-Gerontologist Memory Clinic, National Hospital of Geriatrics and Gerontology, Caja Costarricense de Seguro Social, San José, Costa Rica. 'Physician Neurologist Memory Clinic, National Hospital of Geriatrics and Gerontology, CAja Costarricense de Seguro Social, San José, Costa Rica.

Erick Miranda Valverde. Memory Clinic, National Geriatric and Gerontology Hospital – San José Costa – Rica. E-mail: emirandaval@gmail.com

Disclosure: The authors report no conflicts of interest.

Received August 15, 2017. Accepted in final form October 16, 2017.



tant to screen patients with PD because of the risk of developing dementia associated with PD (PDD).

Currently, several screening tests have been recommended for the evaluation of cognitive impairment in PD, such as the Scale for Outcomes of Parkinson's Disease Cognition (SCOPA-COG), Mini-Mental Parkinson (MMP), Parkinson Neuropsychometric Dementia Assessment (PANDA), Parkinson Disease Dementia-Short Screen (PDD-Short Screen) and the Montreal Cognitive Assessment (MoCA).5 This last test was originally designed for the evaluation of mild cognitive impairment associated with AD, and evaluates memory, executive functions and verbal fluency among others, and can be applied in a short period of time. It has been used for the cognitive evaluation of the patients with PD to identify the presence of cognitive deficit when the MMSE score is normal. This test has a maximum value of 30 points and can be administered in 10 minutes. At least four studies have been conducted demonstrating that the MoCA in patients with PD is sensitive to small cognitive changes. The MoCA has shown a good testretest effect, low inter-evaluator variability, sensitivity of 82% and specificity of 75% with respect to neuropsychological tests for detecting MCI and Dementia using a 26-point cut-off.^{6,7} MoCA has proven the most suitable instrument for screening Mild Cognitive Impairment in PD (PD-MCI) and PDD in clinical trials according to the Parkinson Study Group (PSG) Cognitive/Psychiatric Working Group.

The objective of this research was to evaluate the usefulness of the MoCA in the detection of cognitive impairment in PD patients without cognitive complaint and with normal MMSE score.

METHODS

A total of 40 PD patients were selected from Neurology outpatient consultations at the National Hospital of Geriatrics and Gerontology. Disease diagnosis was according to the criteria of the United Kingdom Parkinson's Disease Society Brain Bank Diagnostic Criteria for Parkinson's Disease. Patients with disease duration <10 years, no cognitive complaints and an MMSE^{8,9} score ≥24 points were included. During a 12-month period, all patients underwent a complete anamnesis, a neuroimaging study consisting of Computed Axial Tomography scan, general biochemistry workup consisting of hemogram, renal function, electrolytes, renal and hepatic function tests, thyroid function, VDRL (Venereal disease research laboratory), levels of B12 vitamin and folic acid, in order to exclude other causes that might explain the cognitive deterioration. In addition, participants completed the Clock Drawing Test (scored according to Cacho et al.), ^{10,11} Barthel's Scale of Activities of Daily Living ¹² (BADL), Lawton's Scale of Instrumental Activities of Daily Living ¹² (IADL), the Yesavage Geriatric Depression Scale ¹³ (GDS) 15-point version, the Hoehn and Yahr scale, ^{14,15} as well as the MoCA test. Cases with a previous diagnosis of mild cognitive impairment, dementia, or psychiatric illness including depression, an advanced medical condition or severe sensory deficit were excluded.

In all cases, the different scales were applied by the same evaluator, trained in the use of all the tests, in order to standardize the evaluation. Basic sociodemographic information was also collected. The study was approved by the Local Scientific Ethics Committee (CLOBI) under number 03-2014.

Statistical analysis was performed using the statistical program Stata 13.0 SE (StataCorp. LP 2013). The presence of normal cognition was defined for scores ≥26 points on the MoCA and cognitive impairment as score <26, as determined for the general population. The comparison between the groups with and without cognitive impairment was performed using the Mann-Whitney U-Test.

RESULTS

Among the total cases, females predominated (55%), while average age and formal education were 76.6±6.6 and 7.4±5.3 years, respectively. The mean duration of the disease was 5.5±3.2 years and 85% used Levodopa as part of their baseline therapy (Table 1).

In terms of level of functioning, 65% were at initial functional stages according to the Hoehn and Yahr scale, exhibiting slight repercussions on the BADL and the IADL scales, and had low risk of depression (Table 2).

Regarding cognitive aspects, the average score obtained on the MMSE was 26.7 and on the Clock Drawing Test was 6.8 points. The mean value on the MoCA was 20.7 points, proving abnormal (score <26) in 80% of the cases (Tables 2 and 3). For the different MoCA sub areas, statistically significant differences in visuospatial, attention and delayed recall functions were evident between the normal and abnormal MoCA groups (Table 3).

DISCUSSION

The main finding was that cognitive impairment in PD, defined as an MoCA score lower than 26 according to the recommended score, is a common finding in PD patients without cognitive impairment complaints. Despite normal MMSE scores in all cases, 80% of the patients had abnormal performance on the MoCA according to overall score. This finding indicated that

the majority of patients with PD had some degree of cognitive deterioration on more extensive and sensitive neuropsychological tests such as the MoCA. On the other hand, the comparison of the different cognitive areas showed that visuospatial, attention and delayed recall functions were the most relevant, a result in line with other investigations. 16,17 These findings suggest that initial cognitive impairment in PD patients occurs in a range of cognitive domains and highlights the importance of a thorough cognitive assessment using a simple but sensitive neuropsychological test. It is noteworthy that average scores on the Clock Drawing Test were lower than normal, correlating with alterations on the visuospatial/executive section of the MoCA; data to be considered as a way of compensating for the deficiencies of the MMSE in the evaluation of patients with PD.¹⁷⁻¹⁹ The present study corroborates the facts documented by many other investigations, showing that the MoCA can detect the presence of cognitive deterioration in patients with PD. This is of paramount importance because the presence of MCI represents a risk factor for

Table 1. Clinical and epidemiological characteristics of patients with PD.

| Variable | | Total (n=40) | Percentage |
|-------------------|----------------------|-----------------|------------|
| Gender | Female | 22 | 55 % |
| | Male | 18 | 45 % |
| Age | | 76.6±6.6 | |
| Marital status | Married | 20 | 50 % |
| | Widowed | 9 | 23 % |
| Education | | 7.4±5.3 | |
| Duration of disea | Duration of disease | | |
| Anti-Parkinson's | Levodopa | 34 | 85 % |
| treatment | Dopaminergic agonist | 5 | 12.5 % |
| | Amantadine | 1 | 2.5 % |
| Hoehn and Yahr | Stage 1 | 15 | 37.5 % |
| | Stage 2 | 11 | 27.5 % |
| | Stage 3 | 8 | 20 % |
| | Stages 4 and 5 | 0 | 0% |

Table 2. Cognitive and functional scale scores of PD patients.

| Variable | Minimum | Maximum | Mean | Standard deviation |
|--|---------|---------|------|--------------------|
| Basic activities of daily living (Barthel) | 30 | 100 | 89.6 | 15.9 |
| Instrumental activities of daily living (Lawton) | 0 | 8 | 4.8 | 2.3 |
| Geriatric depression scale (Yesavage) | 0 | 11 | 2.3 | 2.6 |
| MMSE | 24 | 30 | 26.7 | 2.9 |
| Clock Drawing Test | 1 | 10 | 6.8 | 2.9 |
| MoCA | 11 | 30 | 20.7 | 4.8 |

MMSE: mini-mental state examination; MoCA: Montreal Cognitive Assessment.

Table 3. MoCA Results for PD patients.

| Cognitive subarea | MoCA <26 points Mean and SD | MoCA ≥26 points Mean and SD | Z ^a score | P-value |
|------------------------|--------------------------------|--------------------------------|----------------------|---------|
| Total | 32 (80 %) | 8 (20 %) | NA | NA |
| Visuospatial/Executive | 1.5±1.3 | 3.9±1.6 | -3.3 | 0.001* |
| Naming | 2.5±0.9 | 3.0±0.0 | -1.8 | 0.075 |
| Attention | 3.3±1.3 | 5.5±0.8 | -3.7 | 0.000* |
| Language | 2.1±0.9 | 2.8±0.5 | -1.9 | 0.057 |
| Abstraction | 1.8±0.5 | 2.0±0.0 | -1.3 | 0.190 |
| Delayed recall | 1.3±1.3 | 3.4±1.4 | -3.2 | 0.001* |
| Orientation | 5.5±1.0 | 6.0±0.0 | -1.7 | 0.074 |

MoCA: Montreal Cognitive Assessment. aMann-Whitney U-Test *Significance 0.001.

subsequent development of dementia and instruments such as the MMSE lack the sensitivity and specificity to detect the condition.²⁰

In conclusion, patients with PD can present cognitive impairment even when not perceived by the patients. The use of cognitive tests with greater sensitivity and specificity is required to detect these deficits and the MoCA represents a good alternative for this purpose.

Authors contributions. Krisly Arguedas Vásquez: planned, organized and supervised the project, Erick Miranda Valverde: wrote, edited and supervised the manuscript, Daniel Valerio Aguilar: carried out the study and wrote the manuscript, Henri-Jacques Hernández Gabarain: organized and carried out the study.

REFERENCES

- Janvin C, Aarsland D, Larsen JP, Hugdahl K. Neuropsychological profile of patients with Parkinson's disease without dementia. Dement Geriatr Cogn Disord. 2003;15(3):126-31.
- Emre M, Aarsland D, Brown R, Burn DJ, Duyckaerts C, Mizuno Y, et al. Clinical diagnostic criteria for dementia associated with Parkinson's disease. Mov Disord. 2007;22(12):689-707.
- Muslimovic D, Post B, Speelman JD, Schmand B. Cognitive profile of patients with newly diagnosed Parkinson disease. Neurology. 2005; 65(8):1239-45.
- Dalrymple-Alford JC, Livingston L, MacAskill MR, Graham C, Melzer TR, Porter RJ, et al. Characterizing Mild Cognitive Impairment in Parkinson's Disease. Mov Disord. 2011;26(4):629-36.
- Biundo R, Weis L, Pilleri M, Facchini S, Formento-Dojot P, Vallelunga A, Antonini A. Diagnostic and screening power of neuropsychological testing in detecting mild cognitive impairment in Parkinson's disease. J Neural Transm. (Vienna) 2013;120(4):627-33.
- Nasreddine ZS, Phillips NA, Bédirian V, Charbonneau S, Whitehead V, Collin I, et al. The Montreal Cognitive Assessment, MoCA: A Brief Screening Tool For Mild Cognitive Impairment. J Am Geriatr Soc. 2005; 53(4):695-9.
- Chou KL, Amick MM, Brandt J, Camicioli R, Frei K, Gitelman D, et al. A Recommended Scale for Cognitive Screening in Clinical Trials of Parkinson's Disease. Mov Disord. 2010;25(15):2501-7.
- Folstein M, Folstein S, Mchugh P. "Mini Mental State Examination" a practical method for grading the cognitive state of patient for clinicians. J Psychiat Res. 1975;12(3):189-98.
- Ridha B, Rossor M. The Mini Mental State Examination. Practical Neurology. 2005;5:298-303.
- Cacho J, García-García R, Arcaya J, Vicente JL, Lantada N. Una propuesta de aplicación y puntuación del test del reloj en la enfermedad de Alzheimer. Rev Neurol. 1999;28(7):648-55.

- Juby A, Tench Sh, Baker V. The value of clock drawing in identifying executive cognitive dysfunction in people with a normal Mini-Mental State Examination score. CMAJ. 2012;167(8):859-64.
- 12. Trigás-Ferrin M, Ferreira-González L, Meijide-Míguez H. Scales for functional assessment of the elderly. Galicia Clin.. 2011;72:11-16.
- Yesavage JA, Brink TL, Rose TL. Development and validation of a geriatric depression screening scale: a preliminary report. J Psychiatr Res. 1983;17(1):37-49.
- Perlmutter J. Assessment of Parkinson Disease Manifestations. Curr Protoc Neurosci. Chapter 10: Unit10.1 · October 2009
- Poewe W. The natural history of Parkinson's disease. J Neurol. 2006; 253 suppl 7:VII2-6
- Nazem S, Siderowf AD, Duda JE, Have TT, Colcher A, Horn SS, et al. Montreal Cognitive Assessment Performance in Patients with Parkinson's Disease with "Normal" Global Cognition According to Mini-Mental State Examination Score. J Am Geriatr Soc. 2009;57(2):304-8.
- Camargo CHF, Tolentino ES, Bronzini A, Ladeira MA, Lima R, Schultz-Pereira GL, Young-Blood MR. Comparison of the use of screening tools for evaluating cognitive impairment in patients with Parkinson's disease. Dement Neuropsychol. 2016;10(4):344-50.
- Han-Yeong Jeong, Jee-Young Lee, Hee Kyung Park, et al. Clock drawing test to screen for dementia in parkinsonian patients with low educational backgrounds. Neurology Asia. 2016;21(4):357-65.
- Saka E, Elibol B. Enhanced cued recall and clock drawing test performances differ in Parkinson's and Alzheimer's disease-related cognitive dysfunction. Parkinsonism Relat Disord. 2009;15(9):688-91.
- Hoops S, Nazem S, Siderowf AD, Duda JE, Xie SX, Stern MB, Weintraub D. Validity of the MoCA and MMSE in the detection of MCI and dementia in Parkinson disease. Neurology. 2009;73(21):1738-45.