

# Mild Cognitive Impairment in Early Parkinson Disease is Associated with Posterior Cingulate Atrophy. A Voxel Based Morphometry Study

T. O. Dalaker<sup>1,2</sup>, R. Zivadinov<sup>1,3</sup>, J. P. Larsen<sup>2</sup>, M. Beyer<sup>4</sup>, J. Cox<sup>1,3</sup>, G. Alves<sup>2</sup>, K. Bronnick<sup>2</sup>, O-B. Tysnes<sup>5</sup>, R. Antulov<sup>1,6</sup>, and D. Aarsland<sup>2</sup>

<sup>1</sup>Buffalo Neuroimaging Analysis Center, State University of New York at Buffalo, Buffalo, NY, United States, <sup>2</sup>The Norwegian Centre for Movement Disorders, Stavanger University Hospital, Stavanger, Norway, <sup>3</sup>The Jacobs Neurological Institute, Department of Neurology, State University of New York at Buffalo, Buffalo, NY, United States, <sup>4</sup>Department of Radiology, Stavanger University Hospital, Stavanger, Norway, <sup>5</sup>Department of Neurology, Haukeland University Hospital, Bergen, Norway, <sup>6</sup>Department of Radiology, Clinical Hospital Centre Rijeka, Rijeka, Croatia

**Aim.** We investigated whether mild cognitive impairment (MCI) in early Parkinson Disease (PD) is predicted by extent of regional gray matter (GM) atrophy.

**Background.** Cognitive impairment is common in PD(1). Clinical studies have also revealed MCI in non-demented patients, especially in the domains of visuospatial functions and working memory. It has been shown that such MCI is associated with a higher risk of developing dementia(2).

**Methods.** Our study sample consisted of forty-three (43) non-demented, newly diagnosed PD patients and thirty-one (31) cognitively unimpaired sex-matched normal controls (NC) with mean age  $62.5 \pm 9.3$  years. In the PD sample, 21 patients (age  $68.6 \pm 9.1$  years, disease duration  $26.2 \pm 16.5$  months) were classified as MCI and 22 patients were not (age  $60.6 \pm 8.3$ , disease duration  $36.1 \pm 26.4$  months). MCI classification was based upon performance on an extensive and well recognized test battery consisting of seven neuropsychological tests in domains known to be affected in PD: verbal memory, psychomotor speed, visuospatial functioning and executive functioning. Optimized voxel-based morphometry(3) (VBM) was applied to high resolution 3D-T1 weighted MRI scans. Analyses were corrected for white matter hyperintensities using FLAIR lesion mask in order to avoid misclassification of white matter lesions as GM (figure 1). Statistical analysis was performed on modulated GM images and included covariates for appropriate differences in demographic and clinical characteristics between the groups.

**Results.** MCI patients showed a strong trend towards reduced GM matter volume in the posterior cingulate region of the right limbic lobe (figure 2) when compared to cognitively intact patients ( $p = 0.054$  corrected for family wise errors, FWE). There were no other significant areas of regional atrophy when comparing the PD patients with NC.

**Conclusion.** This study shows that the MCI in early PD is associated with region-specific GM atrophy. The cingulate is known to be a part of the anatomical circuits that are involved in cognitive performance such as visuospatial functions(4). Our study shows that cingulate atrophy is associated with early cognitive deficit in PD, and that this region might serve as a possible biomarker for increased risk of developing dementia in PD.

## References:

- 1.Aarsland D, Andersen K, Larsen JP, Lolk A, Kragh-Sorensen P. Prevalence and characteristics of dementia in Parkinson disease: an 8-year prospective study. Arch Neurol. 2003 Mar;60(3):387-92.
- 2.Janvin CC, Larsen JP, Aarsland D, Hugdahl K. Subtypes of mild cognitive impairment in Parkinson's disease: progression to dementia. Mov Disord. 2006 Sep;21(9):1343-9.
- 3.Good CD, Johnsrude IS, Ashburner J, Henson RN, Friston KJ, Frackowiak RS. A voxel-based morphometric study of ageing in 465 normal adult human brains. Neuroimage. 2001 Jul;14(1 Pt 1):21-36.
- 4.Vogt BA, Vogt L, Laureys S. Cytology and functionally correlated circuits of human posterior cingulate areas. Neuroimage. 2006 Jan 15;29(2):452-66.

