

Poster

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Changes of the corpus callosum in the parkinsonism

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ARTICLE INFO ABSTRACT * Correspondence to: The corpus callosum (CC) is the largest white-matter fibre tract in the brain that plays essential role in communicating sensory, motor and cognitive information between the Murat Golpinar two cerebral hemispheres. Corpus callosum area has been examined extensively in neu-Department of Anatomy, rodegenerative diseases as a marker for cortical pathology and for differential diagnosis. Faculty of Medicine, However, there have been very few studies that examined CC area in Parkinson's disease Ondokuz Mayıs University, (PD). In the present study, we aimed to determine whether there are alterations in the CC Samsun, Turkey of PD. The study included 20 (9 females and 11 males) adult controls and 20 (8 females e-mail:golpinarmurat@hotmail.com and 12 males) patients with PD. Structural MRI was done to both patients and controls. Dicom images were analysed using ImageJ software. Midsagittal surface area of the corpus callosum and intracranial cavity were measured using the planimetry technique. The projection area fraction (PAF) of the CC was measured in the midsagittal section of magnetic resonance images. The mean of midsagittal surface area of the corpus callosum and intracranial cavity in healthy subjects were 5.93±0.75 cm² and 147.20±9.11 cm², re-**Keywords:** spectively, and were 6.26±0.75 cm² and 151.10±13.68 cm² for patients. The mean of the Corpus callosum projection area fraction (PAF) for controls and patients were 4.04 cm² \pm 0.54 and 4.16 Magnetic resonance imaging cm²±0.53.There were no statistical differences between groups regarding the midsagittal surface area of the corpus callosum, intracranial cavity and PAF (p≤0.05). In the light Parkinsonism of the literature there are few studies for the assessment of the CC in PD, the obtained Planimetry data of those studies didn't show statistical significant differences between groups and Projection area fraction gender. We conclude that the patient with PD do not have CC atrophy in contrast to other

neurodegenerative diseases. J. Exp. Clin. Med., 2014; 31:143

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