PET evaluation of the relationship between D₂ receptor binding and glucose metabolism in patients with parkinsonism

Makoto Nakagawa,* Yasuo Kuwabara,** Takayuki Taniwaki,*** Masayuki Sasaki,***

Hirofumi Koga,** Koichiro Kaneko,** Kazutaka Hayashi,**

Jun-ichi Kira*** and Hiroshi Honda**

*Department of Radiology, Fukuoka Red Cross Hospital

**Department of Clinical Radiology, Graduate School of Medical Sciences, Kyushu University

***Department of Neurology, Graduate School of Medical Sciences, Kyushu University

****Department of Radiological Sciences, School of Health Sciences, Faculty of Medicine, Kyushu University

Objective: To clarify the relationship between D₂ receptor binding and the cerebral metabolic rate for glucose (CMRGlu) in patients with parkinsonism, we simultaneously measured both of these factors, and then compared the results. Methods: The subjects consisted of 24 patients: 9 with Parkinson's disease (PD), 3 with Juvenile Parkinson's disease (JPD), 9 with multiple system atrophy (MSA), and 3 with progressive supranuclear palsy (PSP). The striatal D₂ receptor binding was measured by the C-11 raclopride transient equilibrium method. CMRGlu was investigated by the F-18 fluorodeoxyglucose autoradiographic method. Results: The D₂ receptor binding in both the caudate nucleus and putamen showed a positive correlation with the CMRGlu in the PD-JPD group, but the two parameters demonstrated no correlation in the MSA-PSP group. The left/right (L/R) ratio of D₂ receptor binding in the putamen showed a positive correlation with that of CMRGlu in the MSA-PSP group, while the two demonsrated no correlation in the PD-JPD group. Conclusion: Our PET study showed striatal D₂ receptor binding and the CMRGlu to be closely related in patients with parkinsonism, even though the results obtained using the L/R ratios tended to differ substantially from those obtained using absolute values. The reason for this difference is not clear, but this finding may reflect the pathophysiology of these disease entities.

Key words: D₂ receptor binding, CMRGlu, ¹¹C-raclopride, ¹⁸FDG, Parkinson's disease