The relationship between fractional anisotorpy in cerebral white matter by high-field (3 Tesla) MRI, and regional cerebral blood flow and oxygen metabolism with PET study in Alzheimer disease.

Hisashi Yonezawa¹⁾, Satoshi Takahashi¹⁾, Masako Kudo¹⁾, Satoko Obara¹⁾, Toshihide Shibata¹⁾, Makiko Okamoto¹⁾, Hideo Tohgi¹⁾, Takashi, Inoue²⁾, Toshiaki Sasaki³⁾ and Kazunori Terasaki³⁾

- 1 Department of Neurology, Iwate Medical University 19-1Uchimaru, Morioka 020-8505, Japan
- 2) Department of Neurosurgery , Iwate Medical University 19-1Uchimaru, Morioka 020-8505, Japan
- 3 Cyclotoron Research center, Iwate Medical University 358-58Tomegamori, Takizawa, 020-0173, Japan

Abstract

[Purpose] The correlation of fractional anisotropy (FA) of the cerebral white matter using high field MRI(3T) and regional cerebral blood flow (rCBF) and oxygen metabolism (regional cerebral oxygen metabolic rate (rCMRO₂) and regional oxygen extraction fraction (rOEF)) using PET is examined in Alzheimer disease (AD). [Subjects and Methods] Subjects were 17 patients with probable Alzheimer disease (AD) (4 males and 13 females; 70.4 4.8 years old, mean SD). The diagnosis of AD was made according to the criteria of NINCDS-ADRDA MRI scans were performed using a Signa VH/i 3.0 T MR imaging system. A spin echo type echo planar imaging (EPI) sequence with diffusion gradients applied in six directions was used for DT imaging (TR 3000 ms, TE 84 ms. To optimize the measurement of diffusion, only two b factors were used (b1=0, b2=1200 s /mm²). The scalar in variants of the fractional anisotropy (FA) was derived for every pixel. The ROI(round areas of 25 mm²) were placed on the subcortical white matter fiber tracts in the middle frontal gyrus, occipital apex, angular gyrus, and the anterior and posterior portions of the corpus callosum and the cingulate bundle in the coronal slices . rCBF, rCMRO2 and rOEF in all cerebral cortex were determined with the steady-state technique using oxygen-15-labeled tracers (15O₂, C15O₂ and C15O) by PET study. [Results] Unexpectedly, there was no correlation between rCBF, rOEF, and rCMRO2 in each cerebral cortex and FA value of right under of the cortex. There were the significant positive correlation in FA value of the left posterior portions of the cingulate bundle and rCBF of the left frontal cortex (r=0.50 p 0.05), of the left parietal cortex(r=0.50 p 0.05), and of the left temporal cortex(r=0.48 p 0.05), and the significant positive correlation in FA value of the right posterior portions of the cingulate bundle and rCBF of the right frontal cortex (r=0.48 p 0.05), and of the right temporal cortex(r=0.47 p 0.05), In addition, the significant positive correlation was recognized in FA value of the left posterior portions of the cingulate bundle and rCMRO₂ of the left frontal cortex (r=0.49 p 0.05), the left parietal cortex(r=0.63 p 0.01), and of the left temporal cortex(r=0.59 p 0.05), and in FA value of the right posterior portions of the cingulate bundle and rCMRO₂ of the right frontal cortex (r=0.69 p 0.01), the right parietal cortex(r=0.69 p 0.01), and of the right temporal cortex(r=0.72 p 0.005). [Conclusion] In the Alzheimer disease, there is the correlation for dysfunction of only posterior portions of the cingulate bundle and dicrease of rCBF and rCMRO₂ in frontal parietal and temporal cortex .