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Balint's Syndrome Secondary to Cortico-Basal Degeneration

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Keywords

Balint's syndrome; Cortico-basal degeneration; Parieto-occipital junction

Clinical Image

A 70-year-old man with Cortico-Basal Degeneration (CBD) diagnosed 7 years ago presented to the neurology clinic complaining of 2 year worsening visual agnosia. Physical exam showed severe cognitive impairment (MMSE of 7/30), severe motor apraxia on rapid alternating movements, rigidity in both arms, and body bradykinesia. He walked slowly with small, shuffling steps. Neuroophthalmolgy exam revealed the classic triad of Balint's syndrome including simultanagnosia (inability of to perceive more than one object at a time), oculomotor apraxia (impaired visual scanning and difficulty in fixating the eyes despite normal eye movement), and optic ataxia (inability to move the hand to a specific object by using vision) [1,2]. PET scan showed hypometabolism in both parietal lobes which is commonly seen in CBD and brain MRI revealed atrophy in the parietooccipital junction (Figure 1). On the basis of these findings, patient was diagnosed with Balint's syndrome as the result of the progressive of CBD.

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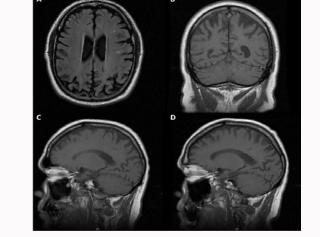


Figure 1: MRI shows atrophy in the parieto-occipital junction bilaterally: (A) Axial T2 FLAIR. (B) Coronal T1 FSE. (C) Sagittal T1 FSE (Left hemisphere). (D) Sagittal T1 FSE (Right hemisphere).

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