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Progressive supranuclear palsy (PSP) is a neurodegenerative disease that occurs in adults over 65 years of age and is characterized by the formation of intraneuronal aggregates of the protein Tau in the brain. Proximal muscles appear to be involved from the initial stages of the disease and it is probable that an anterograde dystonic-dyskinetic motor syndrome may predate cognitive deterioration by several years. No method is currently available to early stage diagnosis of PSP. Amyloid-B Protein Precursor (Ab) is a neuronal protein that is deposited in the brain in Alzheimer's disease (AD). Ab has been shown to aggregate in progressive forms of amyloidosis and specifically in brain regions associated with the pathogenesis of PSP. The objectives of this study are to characterize molecular and immunocytochemical features of Ab in PSP and to determine if Ab is a unique or nonspecific marker of a group of diseases of the neuro-psychiatric system that include PSP, parkinsonism-dementia complex of Guam (PDC), non-Alzheimer's dementia, Alzheimer's disease and Down's syndrome. Furthermore, Ab may be a useful marker of the antecedent anterograde dystonic-dyskinetic motor syndrome that predates the development of cognitive deterioration in PSPS. Therefore, we propose to quantitatively determine if Ab in PSP is abnormal in immunocytochemical, biochemical, and immunoblotting assays.

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