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References Category:Auto CADA novel variant of the Alzheimer's disease associated presenilin 1 gene in a family with autosomal dominant Alzheimer's disease. Alzheimer's disease (AD) is the most common cause of dementia, although many other dementing conditions, including Lewy body dementia, hereditary spinocerebellar ataxia and frontotemporal dementia are also associated with AD. One of the genes responsible for early-onset AD is the presenilin 1 gene (PSEN1). Two major mutations, E280G and I143V, account for approximately 1/3 of early-onset AD. Recently, G209A and G209V mutations have been found to be responsible for a significant proportion of early-onset AD in Japanese populations. Here we report a new mutation, F148I, in the presenilin 1 gene (PSEN1) in a Japanese family, where the mother, son and son's eldest daughter all had AD. The clinical and pathological characteristics of the two families are similar. The E280G, I143V and G209A mutations can be predicted to cause an increase in the risk of AD. Therefore, identification of new variants of the PSEN1 gene and evaluation of the pathogenicity of the mutations may lead to further understanding of the pathogenesis of AD and to the development of new therapies. Protein kinase C-dependent phosphorylation of a novel cytosolic protein (pp71) in normal human placenta. Protein kinase C (PKC) has been shown to be involved in the stimulation of prostaglandin (PG) biosynthesis in human endometrial cells and may regulate

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