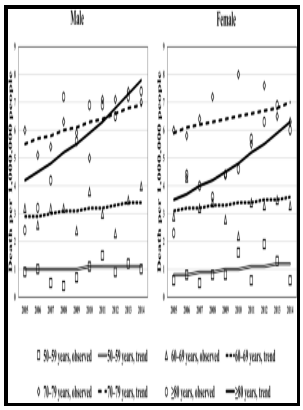
Clinicopathological aspects of Creutzfeldt-Jakob disease

Elsevier - Iatrogenic Creutzfeldt



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-Clinicopathological aspects of Creutzfeldt-Jakob disease
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Heightened incidence of sporadic Creutzfeldt

Department of Health and Human Services, the Public Health Service, the Centers for Disease Control and Prevention, or the authors' affiliated institutions.

Heightened incidence of sporadic Creutzfeldt

In contrast to other inherited human prion diseases, GSS syndrome shows unique neuropathological features consisting of widespread, multicentric PrP plaques.

Heightened incidence of sporadic Creutzfeldt

Population data were obtained from the 2005 and 2010 censuses. All of the more than 40 different known PRNP alleles carrying prion disease-causing mutations have been found to date lie within a short segment of the PRNP gene that encodes the amino acid sequence of the prion protein PrP.

Involvement of hippocampus in Creutzfeldt

Belay, in , 2017 Familial CJD Familial CJD patients generally have clinicopathologic phenotype similar to nongenetic forms of CJD. Strongly Disagree Strongly Agree 1 2 3 4 5 4.

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