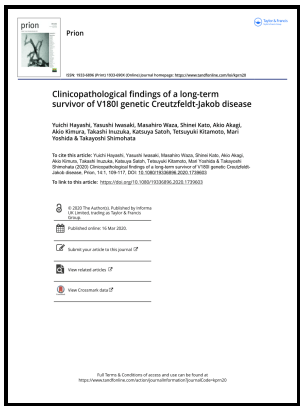
Clinicopathological aspects of Creutzfeldt-Jakob disease

Elsevier - Diagnostic Criteria



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Creutzfeldt-Jakob disease -- Japan.Clinicopathological aspects of Creutzfeldt-Jakob disease
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Neuropathology of prion diseases

By contrast, spongiform changes are rarely present in the brainstem and spinal cord, although PrP accumulation can be demonstrated at these sites.

A nationwide trend analysis in the incidence and mortality of Creutzfeldt

In the age groups of 60—69 and 70—79 years, AAPCs significantly increased across the entire study period. Consecutive EEGs revealed no periodic synchronous discharge PSD.

Clinicopathological features of the rare form of Creutzfeldt

Cite this article Stoeck, K.

A nationwide trend analysis in the incidence and mortality of Creutzfeldt

To calculate the age-adjusted rates of CJD-associated mortality and incidence, we used the direct age-standardisation method with the 2005 Japanese population split into 10-year age groups as the standard population. Clinical symptomatology is similar to sCJD, although about 10% will manifest vertical gaze palsy or neuropathy.

Involvement of hippocampus in Creutzfeldt

Penetrance of PRNP mutations is usually high, although the existence of healthy octogenarian carriers of certain mutations clearly argues in favor of the existence of non— PRNP-related disease modifiers.

Rodrigo Morales, PhD

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