Belgian Genetic Tests database

DISEASE:
Inherited Creutzfeldt-Jakob disease

NAME:	Inherited Creutzfeldt-Jakob disease
DESCRIPTION:	A rare form of genetic prion disease characterized by typical CJD features (rapidly progressive dementia, personality/behavioral changes, psychiatric disorders, myoclonus, and ataxia) with a genetic cause and sometimes a family history of dementia.
ORPHACODE:	282166
SYNONYMS:	Inherited CJD
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	PRNP
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Source URL: http://gentest.healthdata.be/disease/2023

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- Neurodegeneration (gene panel)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)

Related Analytes

- prion protein

Related Gene Panels

- Neurodegeneration (99 genes) - IPG