DISEASE:
Ataxia-telangiectasia-like disorder

NAME:	Ataxia-telangiectasia-like disorder
DESCRIPTION:	A rare genetic disease characterized by slowly progressive cerebellar degeneration resulting in ataxia, oculomotor apraxia, and other cerebellar symptoms. There is an increased frequency of spontaneous chromosomal aberrations, as well as hypersensitivity to ionizing radiation, while telangiectasia is absent.
ORPHACODE:	251347
SYNONYMS:	ATLD
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	MRE11
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RELATED CONTENT

Related Genetic Tests

- Ciliopathy (gene panel)
- Hereditary cancer (Breast, ovary, colon) (26 genes)
- Hereditary cancer (gene panel)

Related Laboratories

- Centre de Génétique Humaine - CHU Sart-Tilman
- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Gent

Related Analytes

- MRE11 homolog, double strand break repair nuclease

Related Gene Panels

- Cancer (Breast, ovary, colon,…) (26 genes) - ULG
- Ciliopathy (120 genes) - UGent
- Hereditary predisposition to cancer (47 genes) - IPG

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