
DISEASE:
McLeod neuroacanthocytosis syndrome

NAME:	McLeod neuroacanthocytosis syndrome
DESCRIPTION:	McLeod neuroacanthocytosis syndrome (MLS) is a form of neuroacanthocytosis (see this term) and is characterized clinically by a Huntington's disease-like phenotype with an involuntary hyperkinetic movement disorder, psychiatric manifestations and cognitive alterations, and biochemically by absence of the Kx antigen and by weak expression of the Kell antigens.
ORPHACODE:	59306
SYNONYMS:	MLS X-linked McLeod syndrome
XREF(S):	<u>Orphanet</u> <u>ICD-10</u> <u>OMIM</u>
ANALYTE(S):	<u>XK</u>
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Source URL: <http://gentest.healthdata.be/disease/2830>

RELATED CONTENT

Related Analytes

- X-linked Kx blood group antigen, Kell and VPS13A binding protein

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