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# McLeod neuroacanthocytosis syndrome

INSERM

## Source

INSERM. (1999). *Orphanet: an online rare disease and orphan drug data base*. [McLeod neuroacanthocytosis syndrome](#). ORPHA:59306

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.