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Huntington disease-like 2

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base.

Huntington disease-like 2. ORPHA:98934

Huntington disease-like 2 (HDL2) is a severe neurodegenerative disorder considered part of the neuroacanthocytosis syndromes (see this term) characterized by a triad of movement, psychiatric, and cognitive abnormalities.