Unverricht-Lundborg disease

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Unverricht-Lundborg disease. ORPHA:308

Unverricht-Lundborg disease (ULD) is a rare progressive myoclonic epilepsy disorder characterized by action- and stimulus-sensitive myoclonus, and tonic-clonic seizures with ataxia, but with only a mild cognitive decline over time.