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# Autosomal recessive ataxia due to ubiquinone deficiency

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Autosomal recessive ataxia due to ubiquinone deficiency. ORPHA:139485*

This syndrome is characterised by childhood-onset progressive ataxia and cerebellar atrophy.