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Autosomal dominant dopa-responsive dystonia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Autosomal dominant dopa-responsive dystonia. ORPHA:98808

Autosomal dominant dopa-responsive dystonia (DYT 5a) is a rare neurometabolic disorder characterized by childhood-onset dystonia that shows a dramatic and sustained response to low doses of levodopa (L-dopa) and that may be associated with parkinsonism at an older age.