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Adult-onset cervical dystonia, DYT23 type

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Adultonset cervical dystonia</u>, <u>DYT23 type</u>. ORPHA:420492

Adult-onset cervical dystonia, DYT 23 type is a rare, genetic, isolated dystonia characterized by adult-onset, non-progressive, focal cervical dystonia typically manifesting with torticollis and occasionally accompanied by mild head tremor and essential-type limb tremor.

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