Primary dystonia, DYT17 type

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Primary dystonia, DYT17 type. ORPHA:370103

Primary dystonia, DYT17 type is a rare, genetic, isolated dystonia initially presenting as torticollis, and later progressing to segmental or generalized dystonia. Dysphonia and dysarthria also occur later in the disease course.