Adult-onset dystonia-parkinsonism

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Adult-onset dystonia-parkinsonism. ORPHA:199351

Adult-onset dystonia-parkinsonism is a rare neurodegenerative disease usually presenting before the age of 30 and which is characterized by dystonia, L-dopa-responsive parkinsonism, pyramidal signs and rapid cognitive decline.