3p25.3 microdeletion syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. 3p25.3 microdeletion syndrome. ORPHA:435638

3p25.3 microdeletion syndrome is a rare chromosomal anomaly characterized by intellectual disability, epilepsy or EEG abnormalities, poor speech, ataxia, and stereotypic hand movements.