1p35.2 microdeletion syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. 1p35.2 microdeletion syndrome. ORPHA:456298

A very rare, chromosomal anomaly characterized by an intrauterine and postanatal growth retardation, short stature, developmental delay, learning difficulties, hearing loss, hypermetropia,and a recognisable facial dysmorphism including prominenet forehead, long, myopathic facies, fine eyebrows, small mouth and micrognathia.