Pitt-Hopkins-like syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Pitt-Hopkins-like syndrome. ORPHA:221150

Pitt-Hopkins-like syndrome is a rare, genetic, syndromic intellectual disability disorder characterized by severe intellectual disability, lack of speech with normal, or mildly delayed, motor development, episodic breathing abnormalities, early-onset seizures and facial dysmorphism which only includes a wide mouth. Abnormal sleep-wake cycles, autistic behavior and stereotypic movements are commonly associated.