X-linked intellectual disability, Cabezas type

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. X-linked intellectual disability, Cabezas type. ORPHA:85293

An X-linked syndromic intellectual disability characterized by developmental delay, intellectual disability with significant speech impairment, and short stature in male patients. Variable additional clinical features have been associated, including macrocephaly, seizures, tremor, gait abnormalities, hypogonadism, truncal obesity, behavioral disturbances and unspecific facial dysmorphism.