Wolf-Hirschhorn syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Wolf-Hirschhorn syndrome. ORPHA:280

Wolf-Hirschhorn syndrome (WHS) is a developmental disorder characterized by typical craniofacial features, prenatal and postnatal growth impairment, intellectual disability, severe delayed psychomotor development, seizures, and hypotonia.