Male hypergonadotropic hypogonadism-intellectual disability-skeletal anomalies syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Male hypergonadotropic hypogonadism-intellectual disability-skeletal anomalies syndrome. ORPHA:2234

This syndrome is characterized by hypergonadotropic hypogonadism, intellectual deficit, congenital skeletal anomalies involving the cervical spine and superior ribs, and diabetes mellitus.