46,XY complete gonadal dysgenesis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. 46,XY complete gonadal dysgenesis. ORPHA:242

46,XY complete gonadal dysgenesis (46,XY CGD) is a disorder of sex development (DSD) associated with anomalies in gonadal development that result in the presence of female external and internal genitalia despite the 46,XY karyotype.