Congenital hypogonadotropic hypogonadism

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Congenital hypogonadotropic hypogonadism. ORPHA:174590

Congenital hypogonadotropic hypogonadism (CHH) is a rare disorder of sexual maturation characterized by gonadotropin (Gn) deficiency with low sex steroid levels associated with low levels of follicle stimulating hormone (FSH) and luteinizing hormone (LH).