46,XX ovarian dysgenesis-short stature syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. 46,XX ovarian dysgenesis-short stature syndrome. ORPHA:444048

A rare, genetic disorder of sex development characterized by primary amenorrhea, short stature, delayed bone age, decreased levels of estradiol, elevated levels of follicle-stimulating hormone and luteinizing hormone, absent or underdeveloped uterus and ovaries, delayed development of pubic and axillary hair, and normal 46,XX karyotype.