Congenital adrenal hyperplasia due to 11-beta-hydroxylase deficiency

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Congenital adrenal hyperplasia due to 11-beta-hydroxylase deficiency. ORPHA:90795

Congenital adrenal hyperplasia due to 11 beta-hydroxylase (CYP11B1) deficiency is a rare form of congenital adrenal hyperplasia (CAH; see this term) characterized by glucocorticoid deficiency, hyperandrogenism, hypertension and virilization in females.