Congenital adrenal hyperplasia due to 17-alpha-hydroxylase deficiency

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Congenital adrenal hyperplasia due to 17-alpha-hydroxylase deficiency. ORPHA:90793

Congenital adrenal hyperplasia due to 17-alpha-hydroxylase deficiency is a very rare form of congenital adrenal hyperplasia (CAH; see this term) characterized by glucocorticoid deficiency, hypergonadotrophic hypogonadism and severe hypokalemic hypertension.