Classic congenital adrenal hyperplasia due to 21-hydroxylase deficiency, simple virilizing form

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Classic congenital adrenal hyperplasia due to 21-hydroxylase deficiency, simple virilizing form. ORPHA:315311

The simple virilizing form of classical congenital adrenal hyperplasia due to 21-hydroxylase deficiency (classical 21 OHD CAH; see this term) is characterized by genital ambiguity and virilization of the external genitalia in females, hypocortisolism and precocious pseudopuberty without salt-wasting.