21-Hydroxylase Deficiency

National Cancer Institute

Source

National Cancer Institute. 21-Hydroxylase Deficiency. NCI Thesaurus. Code C131087.

A classic form of congenital adrenal hyperplasia that is characterized by severe 21-hydroxylase deficiency, resulting in glucocorticoid and mineralocorticoid deficiency, without clinically significant salt wasting, and androgen excess, which causes virilization in female infants.