

## Opis choroby \*

### Definicja

A form of classic congenital adrenal hyperplasia due to 21-hydroxylase deficiency characterized by abnormal genital development with variable levels of virilization in females and normal genitalia in males in association with glucocorticoid insufficiency with salt-wasting due to aldosterone deficiency, accelerated growth velocity and bone maturation, premature adrenarche and precocious puberty leading to reduced adult height.

### Dane

#### Klasyfikacja

Podtyp kliniczny

#### Synonimy

Classic 21-OHD CAH, salt wasting form

Klasyczny 21-OHD CAH, postać z utratą soli

#### Kod ORPHA

315306

#### Kod OMIM

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#### Kod ICD10

E25.0

#### Kod ICD11

5A71.01

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#### \*Źródło

orphanet