

## **Opis choroby \***

Definicja

A group of rare inherited endocrine disorders caused by a steroidogenic enzyme deficiency and characterized by adrenal insufficiency and variable degrees of hyper- or hypoandrogenism manifestations, depending on disease type and severity.

Dane

**Klasyfikacja**  
Grupa fenomenów

Synonimy  
CAH

**Kod ORPHA**  
418

**Kod OMIM**  
613571

**Kod ICD10**  
E25.0

**Kod ICD11**  
5A71.01

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\*[Źródło](#)

[orphanet](#)