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