Opis choroby *

Definicja

A rare, adrenogenital syndrome characterized by generalized, partial tissue insensitivity to glucocorticoids leading to variable phenotype, including asymptomatic individuals with only biochemical alterations or patients with ambiguous genitalia at birth in females, hypertension, acne, hirsutism, precocious puberty, male-pattern hair loss, anxiety and depression in both sexes, menstrual irregularities in women, and oligospermia in men.

Dane

Klasyfikacja
Choroba

Kod ORPHA
786

Kod OMIM
615962

Kod ICD10
E25.8

Kod ICD11
5A71.00

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*Źródło

orphanet