

## 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