

## Opis choroby \*

### Definicja

A rare, genetic pituitary hormone deficiency characterized by gonadotropin (Gn) deficiency with low sex steroid levels associated with low levels of follicle stimulating hormone (FSH) and luteinizing hormone (LH). This disorder may be associated with a normal (normosmic) or impaired sense of smell (Kallmann syndrome).

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

Gonadotropic deficiency  
Isolated congenital gonadotropin deficiency  
Isolated gonadotropin-releasing hormone deficiency  
Gonadotropic deficiency  
Isolated congenital gonadotropin deficiency  
Isolated gonadotropin-releasing hormone deficiency

#### Kod ORPHA

238666

#### Kod OMIM

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#### Kod ICD10

E23.0

#### Kod ICD11

5A61.0

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#### [\\*Źródło](#)

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