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
-

Kod ICD10
E23.0

Kod ICD11
5A61.0

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*Źródło

orphanet