

## Opis choroby \*

### Definicja

Congenital hypopituitarism is characterized by multiple pituitary hormone deficiency, including somatotroph, thyrotroph, lactotroph, corticotroph or gonadotroph deficiencies, due to mutations of pituitary transcription factors involved in pituitary ontogenesis. Congenital hypopituitarism is rare compared with the high incidence of hypopituitarism induced by pituitary adenomas, transsphenoidal surgery or radiotherapy.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

Familial congenital hypopituitarism

Rodzinna wrodzona niedoczynność przysadki

Złożone niedobory hormonów przysadki, formy genetyczne

Multiple pituitary hormone deficiencies, genetic forms

#### Kod ORPHA

95494

#### Kod OMIM

613986

#### Kod ICD10

E23.0

#### Kod ICD11

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#### \*Źródło

orphanet