

Opis choroby *

Definicja

Hypothyroidism due to mutations in transcription factors involved in pituitary development or function is a type of central congenital hypothyroidism (see this term), a permanent thyroid deficiency that is present from birth, characterized by low levels of thyroid hormones caused by disorders in the development or function of the pituitary.

Dane

Klasyfikacja

Choroba

Kod ORPHA

226307

Kod OMIM

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Kod ICD10

E03.1

Kod ICD11

5A00.01

*Źródło

orphanet