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
-

Kod ICD10
E03.1

Kod ICD11
5A00.01

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*Źródło

orphanet