Opis choroby *

Definicja

Central or secondary congenital hypothyroidism is a type of permanent congenital hypothyroidism (see this term) characterized by permanent thyroid hormone deficiency that is present from birth and secondary to a disorder in the thyroid-stimulating hormone (TSH) - thyrotropin-releasing hormone (TRH) system.

Dane

Klasyfikacja
Grupa fenomenów

Synonimy
Secondary hypothyroidism
Wtórna niedoczynność tarczycy

Kod ORPHA
226298

Kod OMIM
-

Kod ICD10
E03.1

Kod ICD11
5A61.41

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*Źródło

orphanet