## 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