

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	Synonimy
Grupa fenomenów	Secondary hypothyroidism Wtórna niedoczynność tarczycy

Kod ORPHA	Kod OMIM	Kod ICD10
226298	-	E03.1

Kod ICD11
5A61.41

*Źródło

orphanet