Opis choroby *

Definicja

A rare form of thyroid dysgenesis characterized by complete absence of thyroid tissue that results in primary congenital hypothyroidism, a permanent thyroid deficiency that is present from birth.

Dane

Klasyfikacja
Wada morfologiczna

Kod ORPHA
95713

Kod OMIM
225250

Kod ICD10
E03.1

Kod ICD11
5A00.01

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*Źródło

orphanet