## 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

---

*Źródło

orphanet