

Opis choroby *

Definicja

A type of primary congenital hypothyroidism, a permanent thyroid hormone deficiency that is present from birth due to thyroid resistance to TSH.

Dane

Klasyfikacja

Choroba

Kod ORPHA

90673

Kod OMIM

275200

Kod ICD10

E03.1

Kod ICD11

5A00.01

*Źródło

orphanet