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

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*Źródło

orphanet