Opis choroby *

Definicja

Familial thyroid dyshormonogenesis is a type of primary congenital hypothyroidism (see this term), a permanent thyroid hormone deficiency that is present from birth, which results from inborn errors of thyroid hormone synthesis.

Dane

Klasyfikacja
Choroba

Synonimy
Thyroid dyshormonogenesis
Rodzinna dyshormonogeneza tarczycy

Kod ORPHA
95716

Kod OMIM
607200

Kod ICD10
E03.1

Kod ICD11
5A00.00

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*Źródło

orphanet