

Opis choroby *

Definicja

Familial thyroid dysmorphogenesis 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 dysmorphogenesis

Rodzinna dysmorphogeneza tarczycy

Kod ORPHA

95716

Kod OMIM

607200

Kod ICD10

E03.1

Kod ICD11

5A00.00

[*Źródło](#)

orphanet