

Opis choroby *

Definicja

Primary congenital hypothyroidism without thyroid developmental anomaly is a type of primary congenital hypothyroidism (see this term) in which the thyroid gland is anatomically normal.

Dane

Klasyfikacja

Kategoria

Kod ORPHA

95714

Kod OMIM

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Kod ICD10

E03.1

Kod ICD11

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*[Źródło](#)

[orphanet](#)