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