Opis choroby *

Definicja

Primary congenital hypothyroidism is a type of permanent congenital hypothyroidism (see this term), a permanent thyroid hormone deficiency that is present from birth.

Dane

Klasyfikacja
Grupa fenomenów

Kod ORPHA
226295

Kod OMIM
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Kod ICD10
E03.1

Kod ICD11
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*Źródło

orphanet