## Opis choroby *

Definicja

A rare, hereditary, familial primary hyperparathyroidism disease characterized by primary hyperparathyroidism due to single or multiple parathyroid tumors in at least two first-degree relatives in the absence of evidence of other endocrine disorders, tumors and/or systemic manifestations.

Dane

**Klasyfikacja**
Choroba

Synonimy
FIHPT
FIHPT

**Kod ORPHA**
99879

**Kod OMIM**
617343

**Kod ICD10**
E21.0

**Kod ICD11**
5A51.0

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*Źródło

orphanet