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

---

*Źródło

orphanet