Opis choroby *

Definicja

A rare genetic disease characterized by synchronous or metachronous occurrence of primary hyperparathyroidism and ossifying fibroma of the maxilla and/or mandible, associated with an increased risk of parathyroid carcinoma. Occurrence of renal cysts or tumors, multiple uterine polyps, and thyroid tumors has also been reported.

Dane

Klasyfikacja
Choroba

Synonimy
HPT-JT
HPT-JT

Kod ORPHA
99880

Kod OMIM
145001

Kod ICD10
E21.0

Kod ICD11
5A51.0

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*Źródło

orphanet