

Opis choroby *

Definicja

A rare inherited cancer-predisposing syndrome characterized by occurrence of multiple synchronous primary carcinoids of the small intestine. Clinical presentation is otherwise indistinguishable from sporadic carcinoids and includes abdominal pain, flushing, and diarrhea, often becoming manifest only after a long asymptomatic period. Most patients present with low grade tumors. Occurrence of pulmonary carcinoids has also been reported.

Dane

Klasyfikacja

Choroba

Synonimy

Hereditary neuroendocrine tumor of small bowel

Hereditary neuroendocrine tumor of small bowel

Kod ORPHA

456333

Kod OMIM

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Kod ICD10

C17.9

Kod ICD11

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*Źródło

orphanet