Dziedziczny neuroendokrynny guz jelita cienkiego Kod Orpha: 456333 Kod OMIM:

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
-

Kod ICD10
C17.9

Kod ICD11
-

*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.