Rodzinny guz neuroendokrynowy żołądka typu 1 Kod Orpha: 464756 Kod OMIM:

Opis choroby *

Definicja
A rare neoplastic disease characterized by occurrence of atypical and aggressive gastric type 1 neuroendocrine tumors (NET) in early adulthood. The tumors often show nodal infiltration requiring total gastrectomy. Synchronous gastric adenocarcinoma has also been reported. Patients present high serum gastrin concentrations and iron-deficiency anemia (rather than megaloblastic anemia, which is a typical feature in patients with sporadic gastric type 1 NET, where the tumor usually arises on the background of autoimmune atrophic gastritis).
Dane

Klasyfikacja
Choroba

Kod ORPHA
464756

Kod OMIM
-

Kod ICD10
C16.9

Kod ICD11
-

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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.