

Opis choroby *

Definicja

A rare subtype of neuroendocrine neoplasm, arising from enterochromaffin-like cells in the stomach, with a variable clinical presentation, disease course and prognosis, depending on the disease type and histological grade. Most patients are asymptomatic, with diagnosis usually occurring incidentally during gastroscopy, however, symptoms of dyspepsia, anemia, pain, weight loss and gastrointestinal bleeding can be observed. Association with Zollinger-Ellison syndrome and multiple endocrine neoplasia type I has been reported.

Dane

Klasyfikacja

Choroba

Synonimy

GNET

GNET

Gastric NET

Gastric neuroendocrine tumor

NET of stomach

Kod ORPHA

100075

Kod OMIM

-

Kod ICD10

C16.9

Kod ICD11

2B72.1

*Źródło

orphanet