

Opis choroby *

Definicja

A rare functioning neuroendocrine tumor of pancreas characterized by a typically well-differentiated neoplasm composed of cells expressing serotonin. Patients may present with atypical carcinoid syndrome with abdominal pain, diarrhea, weight loss, and/or flushing. Carcinoid syndrome is usually present only when there are liver metastases. The tumors tend to be larger than non-functioning tumors and are associated with a poorer prognosis because they are almost always metastatic.

Dane

Klasyfikacja

Choroba

Synonimy

Serotonin-producing PNET

Serotonin-producing pancreatic NET

Serotonin-producing pancreatic neuroendocrine tumor

Serotonin-producing PNET

Serotonin-producing pancreatic NET

Serotonin-producing pancreatic neuroendocrine tumor

Kod ORPHA

506090

Kod OMIM

-

Kod ICD10

C25.9

Kod ICD11

-

*Źródło

orphanet