## 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 Synonimy

Choroba 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 Kod OMIM
506090

- Kod OMIM Kod ICD10 - C25.9

**Kod ICD11** 

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## <u>\*Źródło</u>

orphanet