

## 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

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#### Kod ICD10

C25.9

#### Kod ICD11

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#### \*Źródło

orphanet