

## Opis choroby \*

### Definicja

A rare neuroendocrine tumor of pancreas characterized by a well-differentiated epithelial pancreatic neuroendocrine neoplasm measuring at least 0.5 cm, without distinct hormonal syndrome. Tumors <0.5 cm are called microadenomas. Microadenomatosis is the multifocal occurrence of microadenomas. Histopathologic examination shows an organoid growth pattern and expression of synaptophysin and chromogranin A on immunohistochemistry. Tumors are often discovered incidentally, or patients may present with symptoms related to local or metastatic tumor spread. Microadenomas are considered benign, while larger tumors may behave in a malignant manner with extrapancreatic spread, metastasis, or recurrence.

### Dane

#### Klasyfikacja

#### Choroba

#### Synonimy

Non-functioning PNET

Rak trzustki nieczynny neuroendokrynnie

Non-functioning pancreatic NET

Non-functioning pancreatic neuroendocrine tumor

Non-functioning well-differentiated NEN of pancreas

Non-functioning well-differentiated neuroendocrine neoplasm of pancreas

Non-functioning well-differentiated pancreatic NEN

Non-functioning well-differentiated pancreatic neuroendocrine neoplasm

#### Kod ORPHA

506075

#### Kod OMIM

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#### Kod ICD10

C25.9

#### Kod ICD11

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