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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*Źródło

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