## Opis choroby *

Definicja

A rare neuroendocrine neoplasm of pancreas characterized by morphologically recognizable neuroendocrine and non-neuroendocrine components, each constituting at least 30% of the tumor volume. Based on histopathology, mixed ductal- and mixed acinar-neuroendocrine carcinomas are distinguished. Patients usually present with unspecific symptoms related to tumor growth and/or metastasis, although occurrence of Zollinger-Ellison syndrome has been reported. Resectability of the tumor is the most important prognostic factor.

Dane

**Klasyfikacja**
Choroba

Synonimy
MiNEN of pancreas
Pancreatic MiNEN
Pancreatic mixed neuroendocrine-nonneuroendocrine neoplasm
MiNEN of pancreas
Pancreatic MiNEN
Pancreatic mixed neuroendocrine-nonneuroendocrine neoplasm

**Kod ORPHA**
506112

**Kod OMIM**
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**Kod ICD10**
C25.9

**Kod ICD11**
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*Źródło

orphanet