

## Opis choroby \*

### Definicja

A rare gastroenterologic disease characterized by recurrent acute pancreatitis and/or chronic pancreatitis in at least 2 first-degree relatives, or 3 or more second-degree relatives in 2 or more generations, for which no predisposing factors are identified. This rare inherited form of pancreatitis leads to irreversible damage to both exocrine and endocrine components of the pancreas.

### Dane

### Klasyfikacja

Choroba

#### Kod ORPHA

676

#### Kod OMIM

167800

#### Kod ICD10

K86.1

#### Kod ICD11

DC32.2

---

### \*Źródło

orphanet