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

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*Źródło

orphanet