

Opis choroby *

Definicja

A rare pancreatic disease characterized by chronic non-alcoholic pancreatitis that presents with abdominal pain, steatorrhea, obstructive jaundice and responds well to steroid therapy and is seen in two subforms: type 1 which affects elderly males, involves other organs and has increased immunoglobulin G4 (IgG4) levels and type 2 which affects both sexes equally but presents at a younger age and has no other organ involvement or increased IgG4 levels.

Dane

Klasyfikacja	Synonimy
Grupa fenomenów	AIP AIP

Kod ORPHA	Kod OMIM	Kod ICD10
103919	-	K86.1

Kod ICD11
DC33

*Źródło

orphanet