Opis choroby *

Definicja

A rare disorder of lipid metabolism characterized by childhood onset of steatorrhea due to isolated pancreatic colipase deficiency, while other exocrine pancreatic enzymes are normal. Early formation of gallstones, as well as vitamin B12 deficiency with megaloblastic anemia have also been reported. There have been no further descriptions in the literature since 1982.

Dane

Klasyfikacja
Choroba

Kod ORPHA
309108

Kod OMIM
614338

Kod ICD10
K90.3

Kod ICD11
5C62

---

*Źródło

orphanet