

Opis choroby *

Definicja

A rare genetic hyperlipidemia characterized by excessive increase in plasma triglyceride levels due to the accumulation of chylomicrons. Clinical manifestations include recurrent episodes of severe acute pancreatitis, abdominal pain, nausea, fatigue, diarrhea, constipation, hepatosplenomegaly, eruptive xanthomas, and failure to thrive. Children may often be asymptomatic. The condition is not associated with severe atherosclerosis.

Dane

Klasyfikacja

Choroba

Kod ORPHA

444490

Kod OMIM

615947

Kod ICD10

E78.3

Kod ICD11

-

*Źródło

orphanet