Opis choroby *

Definicja

Familial hypercholanemia is a very rare genetic disorder characterized clinically by elevated serum bile acid concentrations, itching, and fat malabsorption reported in patients of Old Order Amish descent.

Dane

Klasyfikacja
Choroba

Synonimy
Hereditary hypercholanemia
Dziedziczna hipercholanemia

Kod ORPHA
238475

Kod OMIM
607748

Kod ICD10
E88.8

Kod ICD11
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*Źródło

orphanet