

## Opis choroby \*

### Definicja

A rare, progressive metabolic liver disease due to marked to complete lysosomal acid lipase deficiency and characterized by dyslipidemia and massive lipid accumulation leading to hepatomegaly and liver dysfunction, splenomegaly, accelerated atherosclerosis.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

LAL deficiency

Niedobór LAL

#### Kod ORPHA

275761

#### Kod OMIM

278000

#### Kod ICD10

E75.5

#### Kod ICD11

5C56.0Y

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#### [\\*Źródło](#)

orphanet