

Opis choroby *

Definicja

A rare, genetic lipodystrophy characterized by abnormal subcutaneous fat distribution, resulting in preservation of visceral, neck and axillary fat and absence of lower limb and femorogluteal subcutaneous fat. Additional clinical features are acanthosis nigricans, insulin-resistant type II diabetes mellitus, dyslipidemia, and hypertension, leading to pancreatitis, hepatomegaly and hepatic steatosis.

Dane

Klasyfikacja

Choroba

Synonimy

CIDEC-related FPLD

FplD5

FplD zależna od CIDEC

FPLD5

Kod ORPHA

435651

Kod OMIM

615238

Kod ICD10

E88.1

Kod ICD11

5A44

[*Źródło](#)

orphanet