Opis choroby *

Definicja

A rare, genetic lipodystrophy characterized by abnormal subcutaneous fat distribution, resulting in preservation of visceral, neck and axilliary 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

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*Źródło

orphanet