## Opis choroby *

Definicja

A rare, genetic lipodystrophy characterized by a loss of subcutaneous adipose tissue from the trunk, buttocks and limbs; fat accumulation in the neck, face, axillary and pelvic regions; muscular hypertrophy; and usually associated with metabolic complications such as insulin resistance, diabetes mellitus, dyslipidemia and liver steatosis.

Dane

**Klasyfikacja**
Choroba

Synonimy
Dunnigan syndrome
Rodzinna lipodystrofia częściowa typu 2
Zespół Dunnigana
FPLD2
Familial partial lipodystrophy type 2

**Kod ORPHA**
2348

**Kod OMIM**
151660

**Kod ICD10**
E88.1

**Kod ICD11**
5A44

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*Źródło

orphanet