

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

Kod ORPHA	Kod OMIM	Kod ICD10
2348	151660	E88.1

Kod ICD11
5A44

*Źródło

orphanet