

Opis choroby *

Definicja

A rare familial partial lipodystrophy characterized by adult onset of distal lipoatrophy with gluteofemoral fat loss, as well as increased fat accumulation in the face and trunk and visceral adiposity. Additional manifestations include diabetes mellitus, atherogenic dyslipidemia, eyelid xanthelasmas, arterial hypertension, cardiovascular disease, hepatic steatosis, acanthosis nigricans on axillae and neck, hirsutism, and muscular hypertrophy of the lower limbs.

Dane

Klasyfikacja	Synonimy
Choroba	FPLD3 FpID zależna od PPARG FpID3 Rodzinna częściowa lipodystrofia typu 3 Familial partial lipodystrophy type 3 PPARG-related FPLD

Kod ORPHA	Kod OMIM	Kod ICD10
79083	604367	E88.1

Kod ICD11
5A44

*Źródło

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