

## 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 xanthelasma, arterial hypertension, cardiovascular disease, hepatic steatosis, acanthosis nigricans on axillae and neck, hirsutism, and muscular hypertrophy of the lower limbs.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

FPLD3

FplD zależna od PPARG

FplD3

Rodzinna częściowa lipodystrofia typu 3

Familial partial lipodystrophy type 3

PPARG-related FPLD

#### Kod ORPHA

79083

#### Kod OMIM

604367

#### Kod ICD10

E88.1

#### Kod ICD11

5A44

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#### \*Źródło

orphanet