Opis choroby *

Definicja

A rare familial partial lipodystrophy characterized by adult onset of distal lipoatrophy and severe insulin resistance in the liver and peripheral tissues, hyperinsulinemia, and diabetes mellitus. Acanthosis nigricans and hypertension have been reported in association.

Dane

Klasyfikacja
Choroba

Synonimy
AKT2-related FPLD
FplD zależna od AKT2

Kod ORPHA
79085

Kod OMIM
-

Kod ICD10
E88.1

Kod ICD11
5A44

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*Źródło

orphanet