Familial partial lipodystrophy

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Familial partial lipodystrophy. ORPHA:98306

Familial partial lipodystrophy (FPLD) is a group of rare genetic lipodystrophic syndromes characterized, in most cases, by fat loss from the limbs and buttocks, from childhood or early adulthood, and often associated with acanthosis nigricans, insulin resistance, diabetes, hypertriglyceridemia and liver steatosis.