

[Open Peer Review on Qeios](#)

LIPE-related familial partial lipodystrophy

INSERM

Source

INSERM. (1999). *Orphanet: an online rare disease and orphan drug data base. LIPE-related familial partial lipodystrophy. ORPHA:435660*

A rare, genetic lipodystrophy characterized by abnormal subcutaneous fat distribution, resulting in excess accumulation of fat in the face, neck, shoulders, axillae, trunk and pubic region, and loss of subcutaneous fat from the lower extremities. Variable common additional features are progressive adult onset myopathy, insulin resistance, diabetes, hypertriglyceridemia, hepatic steatosis, and vitiligo.