Familial multiple lipomatosis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Familial multiple lipomatosis. ORPHA:199276

Familial multiple lipomatosis is a rare, benign, genetic skin disease characterized by numerous, painless, encapsulated lipomas located in the subcutaneous adipose tissue of the trunk and extremities, with relative sparing of the neck and shoulders. Association with gastroduodenal lipomatosis, brain anomalies or lipomatosis, and refractory epilepsy has been reported.