Open Peer Review on Qeios

JMP syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. JMP syndrome. ORPHA:324999

Joint contractures, muscle atrophy, microcytic anemia and panniculitis-induced lipodystrophy (JMP) syndrome is a rare autoinflammatory disorder belonging to the proteasome disability syndrome (see this term) group and characterized by sclerodermic skin with the presence of erythematous lesions, joint contractures, generalized or partial lipodystrophy, muscle atrophy and short stature.