Open Peer Review on Qeios

Multiminicore myopathy

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Multiminicore myopathy. ORPHA:598

Multi-minicore Disease (MmD) is a hereditary neuromuscular disorder characterized by multiple cores on muscle biopsy and clinical features of a congenital myopathy.