

Open Peer Review on Qeios

Niemann-Pick disease type B

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Niemann-Pick disease type B.</u> ORPHA:77293

Niemann-Pick disease type B is a mild subtype of Niemann-Pick disease, an autosomal recessive lysosomal disease, and is characterized clinically by onset in childhood with hepatosplenomegaly, growth retardation, and lung disorders such as infections and dyspnea

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