Open Peer Review on Qeios

Beta-mannosidosis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Beta-mannosidosis. ORPHA:118

Beta-mannosidosis is a very rare lysosomal storage disease characterized by developmental delay of varying severity and hearing loss, but that can manifest a wide phenotypic heterogeneity.