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Mucolipidosis Type III Gamma

National Cancer Institute

Source

National Cancer Institute. <u>Mucolipidosis Type III Gamma</u>. NCI Thesaurus. Code C129978.

An autosomal recessive condition caused by mutation(s) in the GNPTAG gene, encoding N-acetylglucosamine-1-phosphotransferase subunit gamma. It is characterized by a slowing of the growth rate in childhood, joint stiffness, mild cognitive impairment, and cardiorespiratory insufficiency.

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