Mucopolysaccharidosis Type IIIC

National Cancer Institute

Source

National Cancer Institute. Mucopolysaccharidosis Type IIIC. NCI Thesaurus. Code C84899.

A rare autosomal recessive lysosomal storage disease caused by deficiency of the enzyme acetyl-CoA:alpha-glucosaminide acetyltransferase. It is characterized by behavioral changes, sleep disturbances, and mental developmental delays.