Mucopolysaccharidosis Type IIID

National Cancer Institute

Source

National Cancer Institute. Mucopolysaccharidosis Type IIID. NCI Thesaurus. Code C84900.

A rare autosomal recessive lysosomal storage disease caused by deficiency of the enzyme N-acetylglucosamine-6-sulfatase. It is characterized by behavioral changes, sleep disturbances and mental developmental delays.