Maroteaux-Lamy Syndrome

National Cancer Institute

Source

National Cancer Institute. Maroteaux-Lamy Syndrome. NCI Thesaurus. Code C61264.

A rare autosomal recessive lysosomal storage disease caused by deficiency of the enzyme N-acetylgalactosamine-4-sulfatase. It is characterized by organomegaly, short stature, joint stiffness, otitis media, and respiratory illnesses.