Mannosidosis

National Cancer Institute

Source

National Cancer Institute. Mannosidosis. NCI Thesaurus. Code C61275.

A rare autosomal recessive lysosomal storage disease characterized by a deficient activity of the enzymes alpha-D-mannosidase or beta-mannosidase. Clinical signs and symptoms include hepatomegaly, splenomegaly, hearing loss, mental retardation, skeletal abnormalities, and recurrent respiratory infections.