Hurler Syndrome

National Cancer Institute

Source

National Cancer Institute. Hurler Syndrome. NCI Thesaurus. Code C61261.

An autosomal recessive inherited disorder of mucopolysaccharide metabolism. It is the most severe form of mucopolysaccharidosis type I. It is characterized by deficiency of the enzyme alpha-L-iduronidase resulting in the accumulation of mucopolysaccharides in the tissues.