

Opis choroby *

Definicja

Mucopolysaccharidosis type 1 (MPS 1) is a rare lysosomal storage disease belonging to the group of mucopolysaccharidoses. There are three variants, differing widely in their severity, with Hurler syndrome being the most severe, Scheie syndrome the mildest and Hurler-Scheie syndrome giving an intermediate phenotype.

Dane

Klasyfikacja

Choroba

Synonimy

Alpha-L-iduronidase deficiency

MPS1

MPSI

Mukopolisacharydoza typu I

Niedobór alfa-L-iduronidazy

MPS1

MPSI

Mucopolysaccharidosis type I

Kod ORPHA

579

Kod OMIM

607016

Kod ICD10

E76.0

Kod ICD11

5C56.30

[*Źródło](#)

orphanet