

## Opis choroby \*

### Definicja

A lysosomal storage disease with multisystemic involvement leading to a massive accumulation of glycosaminoglycans and a wide variety of symptoms including distinctive coarse facial features, short stature, cardio-respiratory involvement and skeletal abnormalities. It manifests as a continuum varying from a severe form with neurodegeneration to an attenuated form without neuronal involvement.

### Dane

Klasifikacja	Synonimy
Choroba	Hunter syndrome
	MPS2
	MPSII
	Mukopolisacharydoza typu II
	Niedobór 2-sulfatazy idorunianu
	Zespół Huntera
	Iduronate 2-sulfatase deficiency
	MPS2
	MPSII
	Mucopolysaccharidosis type II

**Kod ORPHA**  
580

**Kod OMIM**  
309900

**Kod ICD10**  
E76.1

**Kod ICD11**  
5C56.31

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\*Źródło

orphanet