

## Opis choroby \*

### Definicja

GM1 gangliosidosis is a rare lysosomal storage disorder characterized biochemically by deficient beta-galactosidase activity and clinically by a wide range of variable neurovisceral, ophthalmological and dysmorphic features.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Choroba	Beta-galactosidase-1 deficiency Choroba Landinga Niedobór beta-galaktozydazy-1 Niedobór GLB1 GLB1 deficiency Landing disease

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
354	230650	E75.1

**Kod ICD11**  
5C56.00

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### \*Źródło

orphanet