

## Opis choroby \*

### Definicja

A rare peroxisome biogenesis disorder (the most severe variant of Peroxisome biogenesis disorder spectrum) characterized by neuronal migration defects in the brain, dysmorphic craniofacial features, profound hypotonia, neonatal seizures, and liver dysfunction.

### Dane

Klasyfikacja	Synonimy	
Choroba	Cerebrohepatorenal syndrome Zespół mózgowo-wątrobowo-nerkowy ZS Severe PBD-ZSD Severe peroxisome biogenesis disorder- Zellweger spectrum disorder ZS	
Kod ORPHA	Kod OMIM	Kod ICD10
912	614887	Q87.8
Kod ICD11		
5C57.0		

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### \*Źródło

orphanet