

## Opis choroby \*

### Definicja

A rare group of neurodegenerative disorders with a prenatal onset characterized by hypoplasia and/or atrophy of the cerebellum and pons. Involvement of supratentorial structures is variable. Multiple forms have been described based on severity, age of onset and clinical presentation.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Grupa fenomenów	PCH Atrofia mostowo-mózdkowa Hipoplazja mostowo-mózdkowa PCH Pontoneocerebellar atrophy Pontoneocerebellar hypoplasia

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
98523	-	Q04.3

**Kod ICD11**  
LD20.01

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### \*Źródło

orphanet