

## Opis choroby \*

### Definicja

A rare developmental defect during embryogenesis characterized by moderate to severe prenatal and postnatal growth retardation, microcephaly, a distinctive facial appearance, profound psychomotor delay, hip and knee contractures and rockerbottom feet.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych	Bowen syndrome, Hutterite type
	Zespół Bowena, typ Hutterite

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
1270	211180	Q87.8

**Kod ICD11**  
LD20.2

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### \*Źródło

orphanet