

## Opis choroby \*

### Definicja

A rare genetic neurological disorder characterized by multiple lateral meningoceles, distinctive facial dysmorphism (including hypertelorism, downslanting palpebral fissures, posteriorly rotated ears, micrognathia, and high, narrow palate, among others), and skeletal abnormalities (e. g. vertebral anomalies, wormian bones, short stature, and scoliosis). Multiple additional features may present, such as conductive hearing impairment, hypotonia, and connective tissue and urogenital abnormalities. Cognition is usually normal.

### Dane

#### Klasyfikacja

Zespół wad wrodzonych  
Lehman syndrome  
Zespół Lehmana

#### Synonimy

#### Kod ORPHA

2789

#### Kod OMIM

130720

#### Kod ICD10

Q87.5

#### Kod ICD11

LA07.Y

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#### \*Źródło

orphanet