

## Opis choroby \*

### Definicja

8p23.1 deletion involves a partial deletion of the short arm of chromosome 8 characterized by low birth weight, postnatal growth deficiency, mild intellectual deficit, hyperactivity, craniofacial abnormalities, and congenital heart defects.

### Dane

|                       |   |
|-----------------------|---|
| <b>Klasyfikacja</b>   | <b>Synonimy</b>   |
| Zespół wad wrodzonych | Del(8)(p23.1)<br>Del(8)(p23.1)<br>Monosomia 8p23.1<br>Monosomy 8p23.1 |

|                  |                 |                  |
|------------------|-----------------|------------------|
| <b>Kod ORPHA</b> | <b>Kod OMIM</b> | <b>Kod ICD10</b> |
| 251071           | -               | Q93.5            |

**Kod ICD11**  
LD44.81

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### \*Źródło

orphanet