

## Opis choroby \*

### Definicja

A rare X-linked malformation syndrome characterized by craniofacial abnormalities, grooved nails, intellectual disability and various skeletal and soft tissue abnormalities.

### Dane

|                       |                                 |
|-----------------------|---------------------------------|
| <b>Klasyfikacja</b>   | <b>Synonimy</b>                 |
| Zespół wad wrodzonych | CFND                            |
|                       | CFND                            |
|                       | CFNS                            |
|                       | Zespół czaszkowo-czołowo-nosowy |
|                       | CFNS                            |
|                       | Craniofrontonasal syndrome      |

|                  |                 |                  |
|------------------|-----------------|------------------|
| <b>Kod ORPHA</b> | <b>Kod OMIM</b> | <b>Kod ICD10</b> |
| 1520             | 304110          | Q87.1            |

**Kod ICD11**  
LD25.3

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### \*Źródło

orphanet