

## Opis choroby \*

### Definicja

A rare primary bone dysplasia characterized by severe intrauterine and postnatal growth retardation and short stature in association with craniofacial dysmorphism (such as large forehead, triangular face, low-set ears, and micro-retrognathism) and osteochondrodysplastic lesions. Radiographic findings include epiphyseal maturation delay, abnormal metaphyses, a narrow thorax, small pelvis, and short and broad metacarpal bones and phalanges. There have been no further descriptions in the literature since 1996.

### Dane

#### Klasyfikacja

Zespół wad wrodzonych Mievis-Verellen-Dumoulin syndrome  
Zespół Mievisa, Verellena i Dumoulina

#### Synonimy

#### Kod ORPHA

2867

#### Kod OMIM

601350

#### Kod ICD10

Q87.1

#### Kod ICD11

LD24.2Y

---

#### \*Źródło

orphanet