## Opis choroby \*

Definicja

A rare autosomal recessive microcephalic primordial dwarfism characterized by congenital microcephaly and craniofacial features associated with a spectrum of limb abnormalities ranging from mild to severe. Short stature is frequently observed and often is severe.

Dane

## Klasyfikacja

Zespół wad wrodzonych

**Kod ORPHA** 572761

**Kod OMIM** 

**Kod ICD10** Q87.1

Kod ICD11

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## \*Źródło

orphanet