

## Opis choroby \*

### Definicja

A rare multiple congenital anomalies/dysmorphic syndrome characterized by anomalous ossification and skeletal patterning of the axial and appendicular skeleton, facial dysmorphism and conductive and sensorineural hearing loss.

### Dane

### Klasyfikacja

Choroba

#### Kod ORPHA

1826

#### Kod OMIM

617137

#### Kod ICD10

Q78.5

#### Kod ICD11

LD25.1

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### \*Źródło

orphanet