

## Opis choroby \*

### Definicja

A severe form of otopalatodigital syndrome spectrum disorder, and is characterized by dysmorphic facies, severe skeletal dysplasia affecting the axial and appendicular skeleton, extraskelatal anomalies (including malformations of the brain, heart, genitourinary system, and intestine) and poor survival.

### Dane

Klasyfikacja	Synonimy
Zespół wad wrodzonych	OPD II syndrome OPD syndrome 2 OPD II syndrome OPD syndrome 2

Kod ORPHA	Kod OMIM	Kod ICD10
90652	304120	Q87.0

**Kod ICD11**  
LD25.1

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### \*Źródło

orphanet