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, extraskeletal anomalies (including malformations of the brain, heart, genitourinary system, and intestine) and poor survival.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
OPD II syndrome
OPD syndrome 2
OPD II syndrome
OPD syndrome 2

Kod ORPHA
90652

Kod OMIM
304120

Kod ICD10
Q87.0

Kod ICD11
LD25.1

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*Źródło

orphanet