Opis choroby *

Definicja

Craniofacial conodysplasia is characterised by craniofacial dysplasia, cone-shaped physes of the hands and feet, and neurological manifestations resembling cerebral palsy. It has been described in one family. The syndrome appeared to be transmitted as a dominant trait.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
85168

Kod OMIM
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Kod ICD10
Q87.5

Kod ICD11
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*Źródło

orphanet