Opis choroby *

Definicja

A rare multiple congenital anomalies/dysmorphic syndrome characterized by a variable combination of dental, cutaneous, ocular, and bone abnormalities, including pyramidal and fused molar roots, taurodontism, an abnormal upper lip without a cupid's bow and thickened and wide philtrum, juvenile glaucoma, syndactyly, and clinodactyly. There have been no further descriptions in the literature since 1973.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
Ackerman fused molar roots syndrome
Pyramidal molar - jaskra - nieprawidłowa warga górna

Kod ORPHA
2561

Kod OMIM
200970

Kod ICD10
K00.2

Kod ICD11
-

---

*Źródło

orphanet