

## **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**

#### Synonimy

Zespół wad wrodzonych Ackerman fused molar roots syndrome

Pyramidal molar - jaskra - nieprawidłowa warga  
górną

#### **Kod ORPHA**

2561

#### **Kod OMIM**

200970

#### **Kod ICD10**

K00.2

#### **Kod ICD11**

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#### \*Źródło

orphanet