

## 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 Ackerman fused molar roots syndrome  
Pyramidal molar - jaskra - nieprawidłowa warga  
górną

#### Synonimy

#### Kod ORPHA

2561

#### Kod OMIM

200970

#### Kod ICD10

K00.2

#### Kod ICD11

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#### \*Źródło

orphanet