

Dysplazja ektodermalna - ślepotą

Kod Orpha: 1806 Kod OMIM: 268320

Opis choroby *

Definicja

A rare multiple congenital anomalies/dysmorphic syndrome characterized by intellectual disability, severe visual impairment due to ocular malformations (microphthalmos and microcornea with sclerocornea), short stature, hypotrichosis, dental anomalies, and dysmorphic facial features (such as a narrow nasal bridge with marked distal flaring and low-set, protruding ears). There have been no further descriptions in the literature since 1992.

Dane

Klasyfikacja

Zespół wad wrodzonych

Kod ORPHA

1806

Kod OMIM

268320

Kod ICD10

Q87.8

Kod ICD11

LD27.0Y

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.