Wrodzona uogólniona hipertrichoza typu Ambrasa Kod Orpha: 1023 Kod OMIM: 145701

Opis choroby *

Definicja
Congenital generalized hypertrichosis, Ambras type is an extremely rare type of hypertrichosis lanuginosa congenita, a congenital skin disease, that is characterized by the presence of vellus-type hair on the entire body, especially on the face, ears and shoulders, with the exception of palms, soles, and mucous membranes. Facial and dental anomalies can also be observed, such as triangular, coarse face, bulbous nasal tip, long palpebral fissures, delayed tooth eruption and absence of teeth.

Dane

Klasyfikacja
Podtyp kliniczny

Synonimy
Ambras syndrome
Zespół Ambrasa

Kod ORPHA
1023

Kod OMIM
145701

Kod ICD10
Q84.2

Kod ICD11
LD27.0Y

*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.