Zespół Holzgreve'a, Wagnera i Rehdera
Kod Orpha: 2167 Kod OMIM: 236110

Opis choroby *

Definicja
Holzgreve syndrome is an extremely rare, lethal, multiple congenital anomalies/dysmorphic syndrome characterized by renal agenesis with Potter sequence, cleft lip/palate, oral synechiae, cardiac defects, and skeletal abnormalities including postaxial polydactyly. Intestinal nonfixation and intrauterine growth restriction are also associated. There have been no further descriptions in the literature since 1988.
Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
Cleft palate-Potter sequence-congenital heart anomalies-mesoaxial polydactyly-multiple malformations syndrome
Holzgreve-Wagner-Rehder syndrome

Kod ORPHA
2167

Kod OMIM
236110

Kod ICD10
Q87.8

Kod ICD11
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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.