

Opis choroby *

Definicja

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

Synonimy

Kod ORPHA

2167

Kod OMIM

236110

Kod ICD10

Q87.8

Kod ICD11

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*Źródło

orphanet