Opis choroby *

Definicja

Dislocation of the hip-dysmorphism syndrome is a rare multiple congenital anomalies syndrome characterized by bilateral congenital dislocation of the hip, characteristic facial features (flat midface, hypertelorism, epicanthus, puffiness around the eyes, broad nasal bridge, carp-shaped mouth), and joint hyperextensibility. Congenital heart defects, congenital dislocation of the knee, congenital inguinal hernia, and vesicoureteric reflux have also been reported. There have been no further descriptions in the literature since 1995.

Dane

Klasyfikacja Synonimy

Zespół wad wrodzonych Collins-Pope syndrome

Zespół Collinsa i Pope

 Kod ORPHA
 Kod OMIM
 Kod ICD10

 2412
 601450
 Q87.2

Kod ICD11

_

*Źródło

orphanet