

Opis choroby *

Definicja

Coxoauricular syndrome is an extremely rare primary bone defect, described only in a mother and her three daughters to date, characterized by short stature, hip dislocation, minor vertebral and pelvic changes, and microtia with hearing loss. There have been no further descriptions in the literature since 1981.

Dane

Klasyfikacja

Zespół wad wrodzonych

Kod ORPHA

1508

Kod OMIM

122780

Kod ICD10

Q87.1

Kod ICD11

LD24.E

*Źródło

orphanet