

Opis choroby *

Definicja

A rare primary bone dysplasia characterized by Perthes-like pelvic anomalies (premature closure of the capital femoral epiphyses and widened femoral necks with flattened femoral heads), arthralgias of hips and knees, and occurrence of enchondromata and ecchondromata. There have been no further descriptions in the literature since 1971.

Dane

Klasyfikacja

Zespół wad wrodzonych Hip dysplasia-enchondromata-ecchondroma syndrome

Synonimy

Dysplazja bioder - enchondroma - ecchondroma

Kod ORPHA

3408

Kod OMIM

191520

Kod ICD10

M91.8

Kod ICD11

LD24.2Y

*Źródło

orphanet