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

Synonimy
Hip dysplasia-enchondromata-ecchondroma syndrome
Dysplazja bioder - enchondroma - ecchondroma

Kod ORPHA
3408

Kod OMIM
191520

Kod ICD10
M91.8

Kod ICD11
LD24.2Y

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*Źródło

orphanet