

Opis choroby *

Definicja

A rare bone development disorder characterized by localized, asymmetric osteochondral overgrowth affecting single or multiple epiphyses, most commonly the distal femur, proximal tibia, and talus. The lesions are typically restricted to one side of the epiphysis, with the medial side being affected twice as often as the lateral side. The condition is usually diagnosed in children, and three times more often in boys than in girls. Patients present with pain, limitation in range of motion, and deformity or swelling of the affected joint.

Dane

Klasyfikacja	Synonimy
Zespół wad wrodzonych	Trevor disease Choroba Trevera

Kod ORPHA	Kod OMIM	Kod ICD10
1822	127800	Q74.8

Kod ICD11
LD24.2Y

*Źródło

orphanet