Opis choroby *

Definicja

A rare genetic disease characterized by generalized joint laxity leading to recurrent dislocation of major joints, such as the hip (often with congenital hip dislocation), shoulder, elbow, or patella. Patients often experience muscle and joint pain (sometimes with effusion) and may develop degenerative joint changes at a relatively early age. Skin abnormalities are absent.

Dane

Klasyfikacja
Choroba

Synonimy
Familial joint instability syndrome
EDS XI
Rodzinna luźność stawów
Zespół niestabilności stawów
Zespół rodzinnej niestabilności stawów
Familial joint laxity
Joint instability syndrome

Kod ORPHA
2295

Kod OMIM
147900

Kod ICD10
Q79.6

Kod ICD11
LD28.Y

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*Źródło

orphanet