

## 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