

## Opis choroby \*

### Definicja

Pseudodiastrophic dysplasia is characterized by rhizomelic shortening of the limbs and severe clubfoot deformity, in association with elbow and proximal interphalangeal joint dislocations, platyspondyly, and scoliosis. It has been described in about 10 patients. An autosomal recessive inheritance has been suggested. Pseudodiastrophic dysplasia differs from diastrophic dysplasia (see this term) on the basis of clinical, radiographic, and histopathologic findings. Clubfoot can be treated by surgical therapy, and neonatal contractures and scoliosis can be relieved by physical therapy. Several of the reported patients died in the neonatal period or during infancy.

### Dane

### Klasyfikacja

Zespół wad wrodzonych

#### Kod ORPHA

85174

#### Kod OMIM

264180

#### Kod ICD10

Q78.8

#### Kod ICD11

LD24.E

---

### \*Źródło

orphanet