

Opis choroby *

Definicja

A rare bone development disorder characterized by abnormal bowing of the fibula with subsequent non-healing fractures and formation of a false joint (pseudoarthrosis), and instability and angulation at the pseudoarthrosis site. The defect is typically unilateral and often associated with pseudoarthrosis of the tibia and neurofibromatosis type 1.

Dane

Klasyfikacja	Synonimy
Podtyp kliniczny	Congenital pseudoarthrosis of the fibula Congenital pseudoarthrosis of the fibula

Kod ORPHA	Kod OMIM	Kod ICD10
295022	-	Q74.2

Kod ICD11
LB9Y

*Źródło

orphanet