Synostoza promieniowo-łokciowa
Kod Orpha: 3269 Kod OMIM: 179300

Opis choroby *

Definicja
Congenital radioulnar synostosis is a rare bone disorder that may be isolated or associated with other disorders and that is characterized by failure of segmentation of the radius and ulna during embryological development, causing limited rotational movements of the forearm, which may lead to difficulties with some activities of daily living.
Dane

Klasyfikacja
Wada morfologiczna

Synonimy
Radioulnar fusion
Zrost promieniowo-łokciowy

Kod ORPHA
3269

Kod OMIM
179300

Kod ICD10
Q74.0

Kod ICD11
LB90.3

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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.