

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

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.