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 Synonimy

Wada morfologiczna Radioulnar fusion

Zrost promieniowo-łokciowy

Kod ORPHA Kod OMIM Kod ICD10

3269 179300 Q74.0

Kod ICD11 LB90.3

*Źródło

orphanet