Congenital radioulnar synostosis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Congenital radioulnar synostosis. ORPHA:3269

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.