

Opis choroby *

Definicja

A rare congenital limb malformation characterized by true congenital dislocation of the shoulder, developing in utero. It can be unilateral or bilateral and is usually associated with other abnormalities of the shoulder girdle, such as in the glenoid, the humeral head, the joint capsule, and the scapula. In addition, it may be accompanied by other malformations, like developmental hip dysplasia or cardiac malformation.

Dane

Klasyfikacja

Wada morfologiczna

Kod ORPHA

295030

Kod OMIM

-

Kod ICD10

Q68.8

Kod ICD11

LB91

*Źródło

orphanet