Opis choroby *

Definicja

A rare genetic disease characterized by the association of osteosarcoma with limb anomalies (such as bilateral radioulnar synostosis and clinodactyly, as well as other abnormalities of the hands and feet) and erythroid macrocytosis without anemia. There have been no further descriptions in the literature since 1977.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
Osteosarcoma-limb anomalies-erythroid macrocytosis syndrome
Kostniakomięsak - wady kończyn - makrocytoza krwinek czerwonych

Kod ORPHA
2760

Kod OMIM
165660

Kod ICD10
C41.9

Kod ICD11
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*Źródło

orphanet