Opis choroby *

Definicja

A rare multiple congenital anomalies syndrome characterized by congenital microgastria and a uni- or bilateral limb reduction defect, that can include absent or hypoplastic thumbs, radius, ulna and/or amelia. Association with other variable abnormalities, including intestinal malrotation, asplenia, dysplastic kidneys, hypoplastic lungs, dysplastic corpus collosum, and abnormal genitalia, has been reported.

Dane

Klasyfikacja

Zespół wad wrodzonych

Kod ORPHA

2538

Kod OMIM 156810

Kod ICD10 Q87.8

Kod ICD11 LD2F.1Y

<u>*Źródło</u>

orphanet