Opis choroby *

Definicja

A rare disorder characterized by the association of mullerian duct and distal limb anomalies. Females present with anomalies ranging from a vaginal septum to complete duplication of uterus and vagina, and males present with micropenis. The limb anomalies varied from postaxial polydactyly to severe upper limb hypoplasia with split hand.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
2491

Kod OMIM
146160

Kod ICD10
Q87.8

Kod ICD11
LD2F.1Y

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*Źródło

orphanet