

Opis choroby *

Definicja

Ulbright-Hodes syndrome is characterised by renal dysplasia, growth retardation, phocomelia or mesomelia, radiohumeral fusion, rib abnormalities, anomalies of the external genitalia and a potter-like facies. The syndrome has been described in three infants (one pair of sibs and an unrelated case), all of whom died shortly after birth from respiratory distress resulting from pulmonary hypoplasia and oligohydramnios caused by renal dysplasia. The mode of transmission appears to be autosomal recessive.

Dane

Klasyfikacja

Zespół wad wrodzonych

Synonimy

Renal dysplasia-limb defects syndrome

Dysplazja nerek - mezomelia - zrost kości
promieniowej i ramiennej

Dysplazja nerek - wady kończyn

Renal dysplasia-mesomelia-radiohumeral fusion
syndrome

Kod ORPHA

3404

Kod OMIM

266910

Kod ICD10

Q87.8

Kod ICD11

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*Źródło

orphanet