Opis choroby *

Definicja

A form of renal agenesis characterized by complete absence of kidney development, absent ureters and subsequent absence of fetal renal function resulting in Potter sequence with pulmonary hypoplasia related to oligohydramnios, which is fatal shortly after birth.

Dane

Klasyfikacja
Podtyp kliniczny

Kod ORPHA
1848

Kod OMIM
615721

Kod ICD10
Q60.1

Kod ICD11
LB30.00

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*Źródło

orphanet