

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

*Źródło

orphanet