

Opis choroby *

Definicja

A rare renal disease characterized by congenital unilateral or bilateral narrowing of the renal artery leading to severe arterial hypertension and progressive renal failure in the neonate. Manifestations include hypertensive encephalopathy and/or neurological signs and symptoms due to hyponatremia, polyuria, renal electrolyte loss, proteinuria, and hematuria.

Dane

Klasyfikacja

Choroba

Synonimy

Congenital renovascular hypoplasia

Wrodzona hipoplazja nerkowo-naczyniowa

Kod ORPHA

97598

Kod OMIM

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Kod ICD10

Q27.1

Kod ICD11

LA90.40

*Źródło

orphanet