## 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**
-

**Kod ICD10**
Q27.1

**Kod ICD11**
LA90.40

---

*Źródło

orphanet