DISEASE:
Renal hypoplasia, bilateral

NAME:	Renal hypoplasia, bilateral
DESCRIPTION:	A form of renal hypoplasia characterized by bilateral small kidneys with a deficit in the number of nephrons present. The condition is typically asymptomatic but may be associated with hypertension, and some excretory functional limitations, as well as eventual chronic renal failure.
ORPHACODE:	97362
XREF(S):	Orphanet ICD-10
ANALYTE(S):	PAX2 PBX1
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Nephrotic syndrome, Focal Segmental Glomerulosclerosis (FSGS) , Alport syndrome and podocytopathy (gene panel)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)

Related Analytes

- paired box 2
- PBX homeobox 1

Related Gene Panels

- Nephrotic syndrome, FSGS, Alport syndrome (76 genes) - IPG