DISEASE:
Congenital hydronephrosis

NAME:	Congenital hydronephrosis
DESCRIPTION:	Congenital hydronephrosis is a renal urinary disease characterized by distension and dilation of the renal pelvis and calyces secondary to various congenital obstructive malformations of the kidneys and urinary tract that can evolve to renal atrophy.
ORPHACODE:	2190
XREF(S):	Orphanet MedDRA ICD-10 OMIM
ANALYTE(S):	TBX18
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RELATED CONTENT

Related Genetic Tests

- Renal or urinary tract malformation (CAKUT) (gene panel)

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

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- T-box 18

Related Gene Panels

- Cakut (congenital anomalies of the kidney and urinary tract-1) (69 genes) - IPG