DISEASE:
Familial juvenile hyperuricemic nephropathy type 1

NAME:	Familial juvenile hyperuricemic nephropathy type 1
DESCRIPTION:	Familial juvenile hyperuricemic nephropathy type 1 (FJHN1) is a rare kidney disorder characterized by hyperuricemia, progressive nephropathy, and gout occurring at an early age.
ORPHACODE:	209886
SYNONYMS:	FJHN type 1 Familial juvenile gouty nephropathy Familial nephropathy with gout UMOD-associated FJHN UMOD-associated familial juvenile hyperuricemic nephropathy
XREF(S):	http://www.orpha.net/consor/cgi-bin/OC_Exp.php?lng=en&Expert=209886
ANALYTE(S):	UMOD
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RELATED CONTENT

Related Genetic Tests

- Ciliopathy / polycystic kidney and liver diseases / ADTKD/ nephronophtisis / Bardet-Biedl syndromes and kidney cancers (gene panel)
- Renal or urinary tract malformation (CAKUT) (gene panel)

Related Laboratories

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

Related Analytes

- uromodulin

Related Gene Panels

- Cakut (congenital anomalies of the kidney and urinary tract-1) (69 genes) - IPG
- Ciliopathy, polycystic kidney and liver diseases, ADTKD, nephronophtisis, Bardet-Biedl syndromes and kidney cancers (146 genes) - IPG