DISEASE:
Hereditary clear cell renal cell carcinoma

NAME:	Hereditary clear cell renal cell carcinoma
DESCRIPTION:	Hereditary clear cell renal cell carcinoma (ccRCC) is a hereditary renal cancer syndrome defined as development of ccRCC in two or more family members without evidence of constitutional chromosome 3 translocation, von Hippel-Lindau disease or other tumor predisposing syndromes associated with ccRCC, such as tuberous sclerosis or Birt-Hogg-Dubbé syndrome.
ORPHACODE:	422526
SYNONYMS:	Hereditary clear cell renal cell adenocarcinoma
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	OGG1 SLC49A4 FLCN RNF139 FHIT HSPBAP1 DIRC3
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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)
- Kidney cancer (Renal cell carcinoma and transitional cell carcinoma (TCC) renal pelvis) (gene panel)
- Kidney cancer (renal cell carcinoma) (gene panel)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- disrupted in renal carcinoma 3
- fragile histidine triad diadenosine triphosphatase
- folliculin
- HSPB1 associated protein 1
- 8-oxoguanine DNA glycosylase
- ring finger protein 139
- solute carrier family 49 member 4

Related Gene Panels

- Ciliopathy, polycystic kidney and liver diseases, ADTKD, nephronophtisis, Bardet-Biedl syndromes and kidney cancers (146 genes) - IPG

- Kidney cancer (Renal Cell Carcinoma (RCC)) (14 genes) - KUL
- Kidney cancer (Transitional Cell Carcinoma (TCC)) (14 genes) - KUL
