DISEASE:
Von Hippel-Lindau disease

NAME:	Von Hippel-Lindau disease
DESCRIPTION:	Von Hippel-Lindau disease (VHL) is a familial cancer predisposition syndrome associated with a variety of malignant and benign neoplasms, most frequently retinal, cerebellar, and spinal hemangioblastoma, renal cell carcinoma (RCC), and pheochromocytoma.
ORPHACODE:	892
SYNONYMS:	Familial cerebelloretinal angiomatosis Lindau disease VHL Von Hippel-Lindau syndrome
XREF(S):	Orphanet OMIM MeSH MedDRA ICD-10
ANALYTE(S):	CCND1 VHL
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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)
- Hereditary cancer (gene panel)
- Neuroendocrine tumor (NET) (gene panel)
- Onco-endocrine pathologies (gene panel)
- Paraganglioma-pheochromocytoma (6 genes) - ULG
- Renal cell carcinoma (kidney cancer) (gene panel)
- Von Hippel Lindau
- Von Hippel Lindau disease
- Von Hippel Lindau syndrome

Related Laboratories

- Centre de Génétique Humaine - CHU Sart-Tilman
- Centre de Génétique Médicale UCL
- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Gent
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- cyclin D1
- von Hippel-Lindau tumor suppressor

Related Gene Panels

- Ciliopathy, polycystic kidney and liver diseases, ADTKD, nephronophtisis, Bardet-Biedl syndromes and kidney cancers (146 genes) - IPG
- Hereditary predisposition to cancer (47 genes) - IPG
- Neuroendocrine tumor (NET) (9 genes) - KUL
- Onco-endocine pathologies (50 genes) - UCL
- Paraganglioma-pheochromocytoma (6 genes) - ULG
- Renal cell carcinoma - UGent
