Belgian Genetic Tests database

ANALYTE:
VHL

NAME:	von Hippel-Lindau tumor suppressor
SYMBOL:	VHL
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	VHL1
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Ciliopathy (gene panel)
- Ciliopathy / polycystic kidney and liver diseases / ADTKD/ nephronophtisis / Bardet-Biedl syndromes and kidney cancers (gene panel)
- Erythrocytoses, polycythémies, thrombocytoses et neutropénies congénitales (gene panel)
- Familial cancer predisposition (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Kidney cancer (Renal cell carcinoma and transitional cell carcinoma (TCC) renal pelvis) (gene panel)
- Kidney cancer (renal cell carcinoma) (gene panel)
- Maffucci syndrome (gene panel)
- Myeloid/lymphoid neoplasms with germline predisposition (gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Neuroendocrine tumor (NET) (gene panel)
- Onco-endocrine pathologies (gene panel)
- Overgrowth & vascular anomalies (gene panel)
- Paraganglioma and pheochromocytoma (gene panel)
- Paraganglioma-pheochromocytoma (6 genes) - ULG
- Paraganglioma-pheochromocytoma (gene panel)
- Pediatric oncopredisposition (gene panel)
- Pheochromocytoma - paraganglioma syndrome (gene panel)
- Renal carcinoma (4 genes)
- Renal cell carcinoma (kidney cancer) (gene panel)
- Stroke (gene panel)
- Sturge-Weber syndrome (gene panel)
- Von Hippel Lindau
- Von Hippel Lindau disease
- Von Hippel Lindau syndrome

- « Inherited bone marrow failures syndromes » with or without organ dysfunction

Related Diseases

- Chuvash erythrocytosis
- Hereditary pheochromocytoma-paraganglioma
- Sporadic pheochromocytoma
- Sporadic pheochromocytoma/secreting paraganglioma
- Von Hippel-Lindau disease

Related Gene Panels

- Ciliopathy - UGent
- Ciliopathy, polycystic kidney and liver diseases, ADTKD, nephronophtisis, Bardet-Biedl syndromes and kidney cancers (148genes) - IPG
- Congenital or familial erythrocytosis (5 genes) - ULG
- Erythocyoses, polycythémies, thrombocytoses congénitales (gene panel) - ULG
- Hematologic Familiar Forms - ULG
- Hereditary cancer predisposition - UGent
- Kidney cancer (Renal Cell Carcinoma (RCC)) (14 genes) - KUL
- Kidney cancer (Transitional Cell Carcinoma (TCC)) (14 genes) - KUL
- Maffucci syndrome (65 genes) - KUL
- Nephropathies, hereditary (222 genes) - KUL
- Nephropathy panel - UGent
- Neuroendocrine tumor (NET) (9 genes) - KUL
- Onco-endocine pathologies (50 genes) - UCL
- Overgrowth & vascular anomalies (65 genes) - KUL
- Panel Nephro-ULG-V1
- Paraganglioma and pheochromocytoma (29 genes) - UCL
- Paraganglioma-pheochromocytoma (10 genes) - UGent
- Paraganglioma-pheochromocytoma (6 genes) - ULG
- Paraganglioma-pheochromocytoma (8 genes) - KUL

- Pediatric oncopredisposition - UGent
- Pheochromocytoma - paraganglioma syndrome - UGent
- Renal carcinoma (4 genes) - UCL
- Renal cell carcinoma - UGent
- Stroke - UGent
- Sturge-Weber syndrome (65 genes) - KUL
