Belgian Genetic Tests database

ANALYTE:
MET

NAME:	MET proto-oncogene, receptor tyrosine kinase
SYMBOL:	MET
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	DFNB97 HGFR RCCP2 hepatocyte growth factor receptor
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/analyte/2583

RELATED CONTENT

Related Genetic Tests

- Ciliopathy / polycystic kidney and liver diseases / ADTKD/ nephronophtisis / Bardet-Biedl syndromes and kidney cancers (gene panel)
- Congenital structural heart defects (gene panel)
- Epidermal nevus syndrome (gene panel)
- Familial cancer predisposition (gene panel)
- Hearing loss (deafness), (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability (virtual 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)
- Nephrogenetics / Nephropathy (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Neuromuscular disorders (gene panel)
- Onco-endocrine pathologies (gene panel)
- Overgrowth & vascular anomalies (gene panel)
- Paraganglioma and pheochromocytoma (gene panel)
- Renal carcinoma (4 genes)
- Renal cell carcinoma (kidney cancer) (gene panel)
- Skeletal dysplasia (gene panel)
- Sturge-Weber syndrome (gene panel)

Related Diseases

- Hereditary papillary renal cell carcinoma
- NON RARE IN EUROPE: Autism
- Osteofibrous dysplasia
- Papillary renal cell carcinoma
- Pediatric hepatocellular carcinoma
- Rare autosomal recessive non-syndromic sensorineural deafness type DFNB
- Selection of therapeutic option in non-small cell lung carcinoma

Related Gene Panels

- Ciliopathy, polycystic kidney and liver diseases, ADTKD, nephronophtisis, Bardet-Biedl syndromes and kidney cancers (146 genes) - IPG
- Congenital structural heart defects - UGent
- Hearing loss (deafness) (genepanel) - UZA
- Hereditary cancer predisposition - UGent
- Intellectual disability (gene panel)
- Kidney cancer (Renal Cell Carcinoma (RCC)) (14 genes) - KUL
- Kidney cancer (Transitional Cell Carcinoma (TCC)) (14 genes) - KUL
- Maffucci syndrome (65 genes) - KUL
- Nephropathy panel - UGent
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Neuromuscular disorders - UGent
- 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
- Renal carcinoma (4 genes) - UCL
- Renal cell carcinoma - UGent
- Skeletal dysplasia (genepanel) - UZA
- Sturge-Weber syndrome (65 genes) - KUL
- epidermal nevus syndrome (65 genes) - KUL
