Belgian Genetic Tests database

ANALYTE:
BRAF

NAME:	B-Raf proto-oncogene, serine/threonine kinase
SYMBOL:	BRAF
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	BRAF1
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Cardiofaciocutaneous syndrome (5 genes)
- Cardiomyopathy, hereditary (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Congenital structural heart defects (gene panel)
- Dermatogenetic panel, severe, rare and hereditary genodermatoses (gene panel - 394 genes)
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epidermal nevus syndrome (gene panel)
- Epilepsy gene panel
- Heart / Cardio disorders / Cardiopathy (gene panel)
- Hereditary Spastic Paraplegia (94 genes)
- Hypermethylation promoter MLH1 and p.V600 of BRAF1
- Hypertrophic cardiomyopathy (gene panel)
- Intellectual Disability (gene panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Lynch syndrome - MLH1 promoter hypermethylation and BRAF V600E mutation
- Maffucci syndrome (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Overgrowth & vascular anomalies (gene panel)
- Pediatric oncopredisposition (gene panel)
- Primary ciliary dyskinesia (PCD) Heterotaxyies (gene panel)
- RASopathy (gene panel)
- Short Stature (gene panel)

- Short stature/ Growth retardation/ (gene panel)
- Skin disorders (gene panel)
- Sturge-Weber syndrome (gene panel)
- Vascular malformations (somatic)

Related Diseases

- Cardiofaciocutaneous syndrome
- Classic hairy cell leukemia
- Craniopharyngioma
- Cushing disease
- Differentiated thyroid carcinoma
- Hashimoto-Pritzker syndrome
- Langerhans cell histiocytosis
- NON RARE IN EUROPE: Melanoma
- Noonan syndrome
- Noonan syndrome with multiple lentigines
- Pilomyxoid astrocytoma
- Selection of therapeutic option in colorectal cancer
- Selection of therapeutic option in melanoma
- Selection of therapeutic option in non-small cell lung carcinoma
- Syringocystadenoma papilliferum

Related Gene Panels

- Cardiofaciocutaneous syndrome (5 genes))
- Cardiomyopathy, hereditary (208 genes) - VUB
- Congenital heart disease (29 genes) - VUB
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Congenital structural heart defects - UGent

- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Growth retardation/short stature (genepanel) - UZA
- Hereditary Spastic Paraplegia (94 genes) - KUL
- Heterotaxie PCD - UGent
- Hypertrophic cardiomyopathy (75 genes) - IPG
- Intellectual Disability (104 genes) - IPG
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Maffucci syndrome (65 genes) - KUL
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Overgrowth & vascular anomalies (65 genes) - KUL
- Pediatric oncopredisposition - UGent
- RASopathy - KUL
- Short Stature (46 genes) - IPG
- Skin disorders - UGent
- Sturge-Weber syndrome (65 genes) - KUL
- Vascular malformations (somatic) (19 genes) - UCL
- cardiopathy panel - UGent
- epidermal nevus syndrome (65 genes) - KUL
