Belgian Genetic Tests database

ANALYTE:
TGFB3

NAME:	transforming growth factor beta 3
SYMBOL:	TGFB3
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	prepro-transforming growth factor beta-3
XREF(S):	Orphanet SwissProt Ensembl Genatlas HGNC OMIM Reactome
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RELATED CONTENT

Related Genetic Tests

- Aneurysm, Thoracic Aortic, familial (gene panel)
- Cardiomyopathy, hereditary (gene panel)
- Cardiomyopathy: hypertrophic cardiomyopathy, dilated cardiomyopathy, restrictive cardiomyopathy, left ventricular non-compaction cardiomyopathy, arrhythmogenic right ventricular cardiomyopathy (gene panel)
- Cardiopathies, 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)
- Familial Thoracic Aortic Aneurysm (gene panel)
- Heart / Cardio disorders / Cardiopathy (gene panel)
- Intellectual disability (virtual gene panel)
- Primary Electrical disorders / Brugada syndrome / Long QT syndrome (LQT) / Short QT syndrome (SQT) / Arrhythmogenic right ventricular cardiomyopathy (ARVC) / Catecholaminergic polymorphic ventricular tachycardia (CPVT) (gene panel)
- Skeletal dysplasia (gene panel)
- Stroke (gene panel)
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Diseases

- Familial isolated arrhythmogenic ventricular dysplasia, biventricular form
- Familial isolated arrhythmogenic ventricular dysplasia, left dominant form
- Familial isolated arrhythmogenic ventricular dysplasia, right dominant form
- Familial thoracic aortic aneurysm and aortic dissection

Related Gene Panels

- Cardiomyopathy (genepanel) - UZA
- Cardiomyopathy, hereditary (208 genes) - VUB
- Cardiopathies, hereditary (102 genes) - KUL
- Cleft lip and palate / dysmorphic facial features / craniofacial anomalies (255 genes)) - UCL
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Congenital structural heart defects - UGent
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Familial Thoracic Aortic Aneurysm (21 genes) - UGent
- Familial Thoracic Aortic Aneurysm (genepanel) - UZA
- Familiale thoracale aorta aneurysmata (19 genes) - UGent
- Intellectual disability (gene panel)
- Primary Electrical disorders/Brugada syndrome (genepanel) - UZA
- Skeletal dysplasia (genepanel) - UZA
- Stroke - UGent
- cardiopathy panel - UGent
- tests
