Belgian Genetic Tests database

ANALYTE:
GATA4

NAME:	GATA binding protein 4
SYMBOL:	GATA4
VERSION OF ORPHANET:	2023-06-22 14:14:43
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Cardiomyopathy, hereditary (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Congenital structural heart defects (gene panel)
- Disorders of sex development - Primary Ovarian insufficiency - Hypogonadotropic Hypogonadism (gene panel)
- Heart / Cardio disorders / Cardiopathy (gene panel)
- Intellectual disability (virtual gene panel)
- Primary cardiac arrhythmias (Atrial fibrillation / Brugada syndome / Catech. polymorphic ventricular tachycardia / Early repolaristion syndrome / Ideopathic ventricular fibrillation / Long QT syndrome / Sick sinus syndrome / Short QT syndrome) (gene pane)
- Primary ciliary dyskinesia (PCD) Heterotaxyies (gene panel)

Related Diseases

- 46,XY partial gonadal dysgenesis
- 8p23.1 microdeletion syndrome
- Atrial septal defect, ostium secundum type
- Complete atrioventricular canal-left heart obstruction syndrome
- Complete atrioventricular septal defect with ventricular hypoplasia
- Complete atrioventricular septal defect-tetralogy of Fallot
- Familial atrial fibrillation
- NON RARE IN EUROPE: Ventricular septal defect
- Partial atrioventricular septal defect
- Partial atrioventricular septal defect with ventricular hypoplasia
- Tetralogy of Fallot

Related Gene Panels

- 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
- Disorders of Sex Development - Primary Ovarian Insufficiency - Hypogonadotropic Hypogonadism - UGent
- Heterotaxie PCD - UGent
- Intellectual disability (gene panel)
- Primary cardiac arrhythmias (113 genes) - VUB
- cardiopathy panel - UGent
