ANALYTE:
NKX2-5

NAME:	NK2 homeobox 5
SYMBOL:	NKX2-5
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	CSX1 NKX2.5 NKX4-1 tinman (Drosophila) homolog tinman paralog (Drosophila)
XREF(S):	Orphanet Reactome SwissProt Ensembl Genatlas HGNC OMIM
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RELATED CONTENT

Related Genetic Tests

- Bicuspid aortic valve
- Cardiomyopathy, hereditary (gene panel)
- Cardiopathies, hereditary (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Congenital structural heart defects (gene panel)
- Dilated Cardiomyopathy (Gene panel)
- Endocrine Disorders - Hypothyroidism (gene panel - 42 genes)
- 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)
- 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)
- Primary immune deficiencies (gene panel)
- Thyroid disgenesis (38 genes)

Related Diseases

- Athyreosis
- Atrial septal defect, ostium secundum type
- Atrial septal defect-atrioventricular conduction defects syndrome
- Deletion 5q35

- Familial atrial fibrillation
- Familial bicuspid aortic valve
- Familial isolated congenital asplenia
- Familial progressive cardiac conduction defect
- Hypoplastic left heart syndrome
- NON RARE IN EUROPE: Ventricular septal defect
- Tetralogy of Fallot
- Thyroid ectopia

Related Gene Panels

- Bicuspid aortic valve - UGent
- Cardiomyopathy, hereditary (208 genes) - VUB
- Cardiopathies, hereditary (102 genes) - KUL
- Congenital heart disease (29 genes) - VUB
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Congenital structural heart defects - UGent
- Dilated Cardiomyopathy (79 genes) - IPG
- Endocrine Disorders - Hypothyroidism (42 genes) - ULB
- Heterotaxie PCD - UGent
- Intellectual disability (gene panel)
- Primary Electrical disorders/Brugada syndrome (genepanel) - UZA
- Primary cardiac arrhythmias (113 genes) - VUB
- Primary immune deficiencies - UGent
- Thyroid disgenesis (38 genes) - VUB
- cardiopathy panel - UGent
