DISEASE:
Tetralogy of Fallot

NAME:	Tetralogy of Fallot
DESCRIPTION:	Tetralogy of Fallot is a congenital cardiac malformation that consists of an interventricular communication, also known as a ventricular septal defect, obstruction of the right ventricular outflow tract, override of the ventricular septum by the aortic root, and right ventricular hypertrophy.
ORPHACODE:	3303
XREF(S):	Orphanet OMIM MeSH MedDRA ICD-10 OMIM

ANALYTE(S):	KDR ZFPM2 GATA4 JAG1 NKX2-5 GATA5 GJA5 NKX2-6 GDF1 CITED2 GATA6 FLT4 TBX1
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RELATED CONTENT

Related Genetic Tests

- Cardiopathies, hereditary (gene panel)
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Laboratories

- Centre de Génétique Médicale UCL
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- Cbp/p300 interacting transactivator with Glu/Asp rich carboxy-terminal domain 2
- fms related receptor tyrosine kinase 4
- GATA binding protein 4
- GATA binding protein 5
- GATA binding protein 6
- growth differentiation factor 1
- gap junction protein alpha 5
- jagged canonical Notch ligand 1
- kinase insert domain receptor
- NK2 homeobox 5
- NK2 homeobox 6
- T-box transcription factor 1
- zinc finger protein, FOG family member 2

Related Gene Panels

- Cardiopathies, hereditary (102 genes) - KUL
- Cleft lip and palate / dysmorphic facial features / craniofacial anomalies (255 genes)) - UCL
- Congenital heart disease (29 genes) - VUB
- Hypogonadotropic Hypogonadism/Kallmann (61 genes) - ULG
