Belgian Genetic Tests database

ANALYTE:
FOXC2

NAME:	forkhead box C2
SYMBOL:	FOXC2
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	MFH-1 mesenchyme forkhead 1
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM SwissProt Reactome
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Source URL: http://gentest.healthdata.be/analyte/3506

RELATED CONTENT

Related Genetic Tests

- Arteriovenous malformation (gene panel)
- Capillary malformation - arteriovenous malformation (2 genes)
- Cerebral cavernous malformation (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Congenital structural heart defects (gene panel)
- Intellectual disability (virtual gene panel)
- Nephropathies, hereditary (gene panel)
- Primary lymphedema / fetal hydrops (gene panel)
- Rendu-Osler-Weber disease (4 genes)
- Venous malformation (3 genes)
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Diseases

- Lymphedema-distichiasis syndrome

Related Gene Panels

- 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
- Intellectual disability (gene panel)

- Lymphedema / fetal hydrops (27 genes) - UCL
- Nephropathies, hereditary (219 genes) - KUL
- Vascular malformations (germline) (38 genes) - UCL
- test-test
