Belgian Genetic Tests database

ANALYTE:
SIX1

NAME:	SIX homeobox 1
SYMBOL:	SIX1
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XREF(S):	Orphanet Genatlas HGNC OMIM SwissProt Reactome Ensembl
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Craniosynostosis (gene panel)
- Hearing loss (deafness), (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability (virtual gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Renal or urinary tract malformation (CAKUT) (gene panel)
- Skeletal dysplasia (gene panel)
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Diseases

- BOR syndrome
- Branchiootic syndrome
- Rare autosomal dominant non-syndromic sensorineural deafness type DFNA

Related Gene Panels

- Cakut (congenital anomalies of the kidney and urinary tract-1) (69 genes) - IPG
- Cleft lip and palate / dysmorphic facial features / craniofacial anomalies (255 genes)) - UCL
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB

- Craniosynostosis (32 genes) - KUL
- Hearing loss (deafness) (genepanel) - UZA
- Hearing loss (deafness) syndromic (59 genes) - UZA
- Intellectual disability (gene panel)
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Panel Nephro-ULG-V1
- Skeletal dysplasia (genepanel) - UZA
