

ANALYTE:
SIX5

NAME:	SIX homeobox 5
SYMBOL:	SIX5
VERSION OF ORPHANET:	2023-06-22 14:14:43
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM SwissProt
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

Source URL: <http://gentest.healthdata.be/index.php/index.php/analyte/416>

RELATED CONTENT

Related Genetic Tests

- [Congenital malformation \(gene panel - 1721 genes\)](#)
- [Congenital malformation 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 \(CAKUT\) \(gene panel\)](#)
- [cleft lip with/without cleft palate \(virtual gene panel\)](#)

Related Diseases

- [BOR syndrome](#)

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](#)
- [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](#)

Source URL: <http://gentest.healthdata.be/index.php/index.php/analyte/416>