ANALYTE:
WNT4

NAME:	Wnt family member 4
SYMBOL:	WNT4
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	WNT-4
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
CHANGED:	26 Oct 2023 - 23:49

Source URL: http://gentest.healthdata.be/analyte/1554

RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Disorders of sex development - Primary Ovarian insufficiency - Hypogonadotropic Hypogonadism (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)
- Rokitansky syndrome

Related Diseases

- Mayer-Rokitansky-Küster-Hauser syndrome type 2
- Müllerian aplasia and hyperandrogenism
- SERKAL syndrome

Related Gene Panels

- Cakut (congenital anomalies of the kidney and urinary tract-1) (69 genes) - IPG
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Disorders of Sex Development - Primary Ovarian Insufficiency - Hypogonadotropic Hypogonadism - UGent
- Intellectual disability (gene panel)
- Nephropathies, hereditary (219 genes) - KUL

- Nephropathy panel - UGent
- Panel Nephro-ULG-V1
