ANALYTE:
ANOS1

NAME:	anosmin 1
SYMBOL:	ANOS1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	Adhesion molecule-like, X-linked KALIG-1 Kallmann syndrome interval gene 1 WAP four-disulfide core domain 19 WFDC19 anosmin-1
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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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)
- Hypogonadotropic hypogonadism (33 genes)
- Inherited Kidney Diseases (Gene Panel)
- Kallmann syndrome (ANOS1 gene)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Renal or urinary tract malformation (CAKUT) (gene panel)

Related Diseases

- Kallmann syndrome
- Normosmic congenital hypogonadotropic hypogonadism

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
- Hypogonadotropic Hypogonadism/Kallmann (61 genes) - ULG
- Hypogonadotropic hypogonadism (33 genes) - VUB
- Nephropathies, hereditary (219 genes) - KUL

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