ANALYTE:
WT1

NAME:	WT1 transcription factor
SYMBOL:	WT1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	AWT1 NPHS4 WAGR WIT-2
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM SwissProt Reactome
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Congenital structural heart defects (gene panel)
- Disorders of sex development - Primary Ovarian insufficiency - Hypogonadotropic Hypogonadism (gene panel)
- End-stage renal disease, ESRD (gene panel)
- Familial cancer predisposition (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability (virtual gene panel)
- Leukodystrophy (gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Nephrotic syndrome, Focal Segmental Glomerulosclerosis (FSGS) , Alport syndrome and podocytopathy (gene panel)
- Pediatric oncopredisposition (gene panel)
- WAGR Syndrome
- Wilms tumor (DICER1; WT1 genes)

Related Diseases

- 46,XY complete gonadal dysgenesis
- 46,XY partial gonadal dysgenesis
- Denys-Drash syndrome
- Desmoplastic small round cell tumor
- Familial idiopathic steroid-resistant nephrotic syndrome with diffuse mesangial sclerosis
- Familial idiopathic steroid-resistant nephrotic syndrome with focal segmental hyalinosis

- Frasier syndrome
- Genetic steroid-resistant nephrotic syndrome
- Meacham syndrome
- Nephroblastoma
- Sporadic idiopathic steroid-resistant nephrotic syndrome with diffuse mesangial sclerosis
- WAGR syndrome

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Congenital structural heart defects - UGent
- Disorders of Sex Development - Primary Ovarian Insufficiency - Hypogonadotropic Hypogonadism - UGent
- End-stage renal disease (106 genes) - IPG
- Hereditary cancer predisposition - UGent
- Intellectual disability (gene panel)
- Leukodystrophy - UGent
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Nephrotic syndrome, FSGS, Alport syndrome (76 genes) - IPG
- Panel Nephro-ULG-V1
- Pediatric oncopredisposition - UGent
- Wilms' tumor (2 genes) - KUL

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