Belgian Genetic Tests database

ANALYTE:
PAX2

NAME:	paired box 2
SYMBOL:	PAX2
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XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM SwissProt
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RELATED CONTENT

Related Genetic Tests

- Ciliopathy / polycystic kidney and liver diseases / ADTKD/ nephronophtisis / Bardet-Biedl syndromes and kidney cancers (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- End-stage renal disease, ESRD (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability (virtual gene panel)
- Microphthalmia / Anophthalmia / Coloboma-Anterior Segment Dysgenesis (MAC-ASD) (gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Nephrotic syndrome, Focal Segmental Glomerulosclerosis (FSGS) , Alport syndrome and podocytopathy (gene panel)
- Renal or urinary tract malformation (CAKUT) (gene panel)
- Retinal dystrophy / RETNET (gene panel)

Related Diseases

- Familial idiopathic steroid-resistant nephrotic syndrome with focal segmental hyalinosis
- Genetic steroid-resistant nephrotic syndrome
- Renal coloboma syndrome
- Renal hypoplasia, bilateral

Related Gene Panels

- Cakut (congenital anomalies of the kidney and urinary tract-1) (69 genes) - IPG
- Ciliopathy, polycystic kidney and liver diseases, ADTKD, nephronophtisis, Bardet-Biedl syndromes and kidney cancers (146 genes) - IPG

- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- End-stage renal disease (106 genes) - IPG
- Intellectual disability (gene panel)
- Microphthalmia/Anophthalmia/Coloboma - Anterior Segment Dysgenesis - UGent
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Nephrotic syndrome, FSGS, Alport syndrome (76 genes) - IPG
- Panel Nephro-ULG-V1
- Retinal dystrophy - UGent
