Belgian Genetic Tests database

DISEASE:
Frasier syndrome

NAME:	Frasier syndrome
DESCRIPTION:	A rare genetic, syndromic glomerular disorder characterized by the association of progressive glomerular nephropathy and 46,XY complete gonadal dysgenesis with a high risk of developing gonadoblastoma.
ORPHACODE:	347
XREF(S):	Orphanet OMIM MeSH ICD-10
ANALYTE(S):	WT1
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Source URL: http://gentest.healthdata.be/disease/1567

RELATED CONTENT

Related Genetic Tests

- Nephrotic syndrome, Focal Segmental Glomerulosclerosis (FSGS) , Alport syndrome and podocytopathy (gene panel)
- Wilms tumor (DICER1; WT1 genes)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- WT1 transcription factor

Related Gene Panels

- Nephrotic syndrome, FSGS, Alport syndrome (76 genes) - IPG
- Wilms' tumor (2 genes) - KUL