DISEASE:
Fraser syndrome

NAME:	Fraser syndrome
DESCRIPTION:	A rare congenital malformation mainly characterized by unilateral or bilateral cryptophthalmos, syndactyly and urogenital anomalies.
ORPHACODE:	2052
SYNONYMS:	Cryptophthalmos-syndactyly syndrome
XREF(S):	Orphanet OMIM OMIM ICD-10 MeSH OMIM
ANALYTE(S):	FRAS1 FREM2 GRIP1
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

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RELATED CONTENT

Related Genetic Tests

- Epidermolysis bullosa (gene panel)
- Renal or urinary tract malformation (CAKUT) (gene panel)
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Laboratories

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

Related Analytes

- Fraser extracellular matrix complex subunit 1
- FRAS1 related extracellular matrix 2
- glutamate receptor interacting protein 1

Related Gene Panels

- Cakut (congenital anomalies of the kidney and urinary tract-1) (69 genes) - IPG
- Cleft lip and palate / dysmorphic facial features / craniofacial anomalies (255 genes)) - UCL
- Epidermolysis bullosa and bladder diseases (60 genes) - KUL

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