DISEASE:
Senior-Loken syndrome

NAME:	Senior-Loken syndrome
DESCRIPTION:	A rare autosomal recessive oculo-renal ciliopathy characterized by the association of nephronophthisis (NPHP), a chronic kidney disease, with retinal dystrophy.
ORPHACODE:	3156
SYNONYMS:	Nephronophthisis with retinal dystrophy Renal dysplasia-retinal aplasia syndrome SLSN

XREF(S):	Orphanet MeSH ICD-10 OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM OMIM
ANALYTE(S):	CEP290 INVS IQCB1 NPHP1 NPHP3 NPHP4 SDCCAG8 WDR19 CEP164 TRAF3IP1
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RELATED CONTENT

Related Genetic Tests

- Cholestasis (gene panel)
- Ciliopathy (gene panel)
- Ciliopathy / polycystic kidney and liver diseases / ADTKD/ nephronophtisis / Bardet-Biedl syndromes and kidney cancers (gene panel)
- Early-onset severe obesity
- Hepatorenal disorders (gene panel)
- Leber Congenital Amaurosis - Retinal dystrophy, early onset (gene panel)

Related Laboratories

- Centre de Génétique Humaine - CHU Sart-Tilman
- Centre de Génétique Médicale UCL
- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Gent

Related Analytes

- centrosomal protein 164
- centrosomal protein 290
- inversin
- IQ motif containing B1
- nephrocystin 1
- nephrocystin 3
- nephrocystin 4

- SHH signaling and ciliogenesis regulator SDCCAG8
- TRAF3 interacting protein 1
- WD repeat domain 19

Related Gene Panels

- Cholestasis (40 genes) - UCL
- Ciliopathy (120 genes) - UGent
- Ciliopathy, polycystic kidney and liver diseases, ADTKD, nephronophtisis, Bardet-Biedl syndromes and kidney cancers (146 genes) - IPG
- Early-onset severe obesity (44 genes) - ULG
- Leber Congenital Amaurosis - UGent
