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Full name:	Hepatorenal disorders (13 genes) - UCL
Abbreviation:	Panel GP
Type of panel:	Custom panel
Laboratory:	Centre de Génétique Médicale UCL
Created:	06 Nov 2019 - 14:17
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Related Diseases

- Autosomal recessive polycystic kidney disease
- Infantile nephronophthisis
- Joubert syndrome
- Joubert syndrome with hepatic defect
- Juvenile nephronophthisis
- Late-onset nephronophthisis
- Meckel syndrome

Related Analytes

GENE	% OF CODING SEQUENCE SUFFICIENTLY COVERED TO DETECT HETEROZYGOUS MUTATIONS	COPY NUMBER VARIATION	COMMENTS
BCS1L	100.00	1	Part of Gastro-pneumo panel
CC2D2A	100.00	1	Part of Gastro-pneumo panel
DCDC2	100.00	1	Part of Gastro-pneumo panel
EHHADH	100.00	1	Part of Gastro-pneumo panel
HNF1B	100.00	1	Part of Gastro-pneumo panel
INVS	100.00	1	Part of Gastro-pneumo panel

GENE	% OF CODING SEQUENCE SUFFICIENTLY COVERED TO DETECT HETEROZYGOUS MUTATIONS	COPY NUMBER VARIATION	COMMENTS
MKS1	100.00	1	Part of Gastro-pneumo panel
NPHP1	100.00	1	Part of Gastro-pneumo panel
NPHP3	100.00	1	Part of Gastro-pneumo panel
NPHP4	100.00	1	Part of Gastro-pneumo panel
PKHD1	100.00	1	Part of Gastro-pneumo panel
POLG	100.00	1	Part of Gastro-pneumo panel
TMEM216	100.00	1	Part of Gastro-pneumo panel