GENETIC TEST:
Hepatorenal disorders (gene panel)

FULL NAME:	Hepatorenal disorders (gene panel)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Carrier diagnosis, Post-natal Diagnosis, Prenatal diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA, Amniotic fluid, Chorionic villi
METHOD CATEGORY:	Sequence analysis: entire coding region Deletion/duplication analysis
METHOD TECHNIQUE:	Next Generation Sequencing (NGS)
RIZIV CODE:	565493-565504
ACCREDITATION (ISO 15189):	2022-12-22 / 2027-12-21
TURNAROUND TIME (MAXIMUM):	3 months

CREATED:	06 Nov 2019 - 14:04
CHANGED:	24 Jan 2023 - 15:12

Source URL: http://gentest.healthdata.be/genetic_test/925

RELATED CONTENT

Related Diseases

- Autosomal recessive polycystic kidney disease
- Bardet-Biedl syndrome
- GRACILE syndrome
- HNF1B-related autosomal dominant tubulointerstitial kidney disease
- Infantile nephronophthisis
- Isolated complex III deficiency
- Isolated neonatal sclerosing cholangitis
- Joubert syndrome
- Joubert syndrome with hepatic defect
- Juvenile nephronophthisis
- Late-onset nephronophthisis
- Meckel syndrome
- Mitochondrial DNA-associated Leigh syndrome
- Primary Fanconi renotubular syndrome
- Senior-Loken syndrome

Related Laboratories

- Centre de Génétique Médicale UCL

Related Analytes

- BCS1 homolog, ubiquinol-cytochrome c reductase complex chaperone

- coiled-coil and C2 domain containing 2A
- doublecortin domain containing 2
- enoyl-CoA hydratase and 3-hydroxyacyl CoA dehydrogenase
- HNF1 homeobox B
- inversin
- MKS transition zone complex subunit 1
- nephrocystin 1
- nephrocystin 3
- nephrocystin 4
- PKHD1 ciliary IPT domain containing fibrocystin/polyductin
- DNA polymerase gamma, catalytic subunit
- transmembrane protein 216

Related Gene Panels

- Hepatorenal disorders (13 genes) - UCL
