GENETIC TEST:
Crigler Najjar Syndrome

FULL NAME:	Crigler Najjar Syndrome
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:	565471-565482
ACCREDITATION (ISO 15189):	2022-12-22 / 2027-12-21
TURNAROUND TIME (MAXIMUM):	3 months

CREATED:	24 Jul 2019 - 16:31
CHANGED:	24 Jan 2023 - 15:03

RELATED CONTENT

Related Diseases

- Crigler-Najjar syndrome type 1
- Crigler-Najjar syndrome type 2

Related Laboratories

- Centre de Génétique Médicale UCL

Related Analytes

- UDP glucuronosyltransferase family 1 member A1