GENETIC TEST:
Caroli Disease

FULL NAME:	Caroli Disease
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:	06 Nov 2019 - 10:17
CHANGED:	24 Jan 2023 - 14:58

RELATED CONTENT

Related Diseases

- Caroli disease

Related Laboratories

- Centre de Génétique Médicale UCL

Related Analytes

- PKHD1 ciliary IPT domain containing fibrocystin/polyductin