GENETIC TEST:
Alagille syndrome (2 genes)

FULL NAME:	Alagille syndrome (2 genes)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Carrier diagnosis, Post-natal Diagnosis, Prenatal diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA, Chorionic villi, Amniotic fluid
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 - 11:01
CHANGED:	24 Jan 2023 - 14:34

RELATED CONTENT

Related Diseases

- Alagille syndrome due to 20p12 microdeletion
- Alagille syndrome due to a JAG1 point mutation
- Alagille syndrome due to a NOTCH2 point mutation

Related Laboratories

- Centre de Génétique Médicale UCL

Related Analytes

- jagged canonical Notch ligand 1
- notch receptor 2

Related Gene Panels

- Alagille syndrome (2 genes) - UCL