GENETIC TEST:
Coagulopathies (2 genes)

FULL NAME:	Coagulopathies (2 genes)
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, DNA
METHOD CATEGORY:	Sequence analysis: entire coding region Deletion/duplication analysis
METHOD TECHNIQUE:	Next Generation Sequencing (NGS) MLPA based techniques Bi-directional Sanger Sequence analysis
RIZIV CODE:	565471-565482
TURNAROUND TIME (MAXIMUM):	3 months

CREATED:	24 Jul 2019 - 16:22
CHANGED:	14 Dec 2023 - 15:12

RELATED CONTENT

Related Diseases

- Autosomal dominant macrothrombocytopenia
- Fetal and neonatal alloimmune thrombocytopenia
- Glanzmann thrombasthenia

Related Laboratories

- Centre de Génétique Médicale UCL

Related Analytes

- integrin subunit alpha 2b
- integrin subunit beta 3