GENETIC TEST:
Hemophilia B

FULL NAME:	Hemophilia B
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Carrier diagnosis, Mutation confirmation, Post-natal Diagnosis, Prenatal diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA, Chorionic villi, Amniotic fluid, 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

ACCREDITATION (ISO 15189):	2022-12-22 / 2027-12-21
EQA:	• Genetics of heritable Bleeding Disorders, • Genetics of heritable Bleeding Disorders
TURNAROUND TIME (MAXIMUM):	3 months
CREATED:	19 Jul 2019 - 12:32
CHANGED:	14 Dec 2023 - 15:24

RELATED CONTENT

Related Diseases

- Bleeding disorder in hemophilia B carriers
- Hemophilia B
- Mild hemophilia B
- Moderate hemophilia B
- Severe hemophilia B

Related Laboratories

- Centre de Génétique Médicale UCL

Related Analytes

- coagulation factor IX