GENETIC TEST:
Hemophilia B

FULL NAME:	Hemophilia B
DESCRIPTION:	Screening of all coding exons of the F9 gene.
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) MLPA based techniques
RIZIV CODE:	565471-565482

TURNAROUND TIME (MAXIMUM):	3 - 4 months
CREATED:	19 Jul 2019 - 12:32
CHANGED:	14 Dec 2021 - 11:16
URL:	https://laboboeken.nexuzhealth.com/pboek/internet/GHB%20CME/14756

RELATED CONTENT

Related Diseases

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

Related Laboratories

- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- coagulation factor IX