GENETIC TEST:
Noonan syndrome (Screening PTPN11)

FULL NAME:	Noonan syndrome (Screening PTPN11)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal Diagnosis, Prenatal diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA, Amniotic fluid, Chorionic villi
METHOD CATEGORY:	Targeted variant analysis
METHOD TECHNIQUE:	Bi-directional Sanger Sequence analysis
RIZIV CODE:	565456-565460
ACCREDITATION (ISO 15189):	2021-07-08 / 2026-02-02
TURNAROUND TIME (MAXIMUM):	1o weeks
CREATED:	22 Jul 2019 - 10:47
CHANGED:	01 Mar 2023 - 15:58

URL:	https://laboboeken.nexuzhealth.com/pboek/internet/GHB/15589

Source URL: http://gentest.healthdata.be/genetic_test/126

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