GENETIC TEST:
Gorlin syndrome (PTCH1; SUFU genes)

FULL NAME:	Gorlin syndrome (PTCH1; SUFU genes)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal Diagnosis, Predictive and Pre-symptomatic diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA
METHOD CATEGORY:	Sequence analysis: entire coding region Deletion/duplication analysis
METHOD TECHNIQUE:	Next Generation Sequencing (NGS)
RIZIV CODE:	565530-565541
ACCREDITATION (ISO 15189):	2021-07-08 / 2026-02-02
TURNAROUND TIME (MAXIMUM):	4 - 6 months
CREATED:	19 Jul 2019 - 11:46
CHANGED:	01 Mar 2023 - 15:02

URL:	https://laboboeken.nexuzhealth.com/pboek/internet/GHB%20CME/13402

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