GENETIC TEST:
Melanoma / Familial Atypical Multiple Mole Melanoma Syndrome (gene panel)

FULL NAME:	Melanoma / Familial Atypical Multiple Mole Melanoma Syndrome (gene panel)
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:	565552-565563
ACCREDITATION (ISO 15189):	2021-07-08 / 2026-02-02
TURNAROUND TIME (MAXIMUM):	4 - 6 months
CREATED:	22 Jul 2019 - 09:37
CHANGED:	21 May 2024 - 15:01
URL:	https://laboboeken.nexuzhealth.com/pboek/internet/GHB%20CME/13390

RELATED CONTENT

Related Diseases

- Familial atypical multiple mole melanoma syndrome
- Familial melanoma

Related Laboratories

- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- ACD shelterin complex subunit and telomerase recruitment factor
- BRCA1 associated protein 1
- cyclin dependent kinase 4
- cyclin dependent kinase inhibitor 2A
- melanocyte inducing transcription factor
- protection of telomeres 1
- TERF2 interacting protein
- telomerase reverse transcriptase

Related Gene Panels

- Melanoma and Familial Atypical Multiple Mole Melanoma Syndrome (8 genes) - KUL