

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:	01 Mar 2023 - 15:33

URL:	https://laboboeken.nexuzhealth.com/pboek/internet/GHB%20CME/13390
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RELATED CONTENT

Related Diseases

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- [Centrum Menselijke Erfelijkheid - KUL](#)

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- [BRCA1 associated protein 1](#)
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- [cyclin dependent kinase inhibitor 2A](#)
- [melanocyte inducing transcription factor](#)
- [protection of telomeres 1](#)
- [TERF2 interacting protein](#)
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Related Gene Panels

- [Melanoma and Familial Atypical Multiple Mole Melanoma Syndrome \(8 genes\) - KUL](#)
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