

REPORT

Laboratory Director: George Nasioulas PhD.

Sample Information

 Name:
 XXXXXX XXXXXX
 Report No:
 24007794GR

 ID/Medical ID:
 TEST 3/11/2023
 Date Received:
 21/05/2024

 Date Of Birth:
 01/01/1956
 Date Of Report:
 21/05/2024

Patient Location: XX/XXX

Sample Details

Type of sample #1: PARAFFIN EMBEDDED TISSUE-BLOCK Code of sample #1: XXX

Barcode of sample #1: 24007794GR-1

MGMT, IDH1/IDH2, BRAF, 1p19q

DNA and RNA were extracted from the test sample (QIAmp DSP DNA Mini kit, Qiagen RNeasy FFPE Kit). The promoter methylation was detected using the MGMT Methylation Detection kit (Entrogen) and the QuantStudio 3 Real Time PCR System (ThermoFisher) platform. Mutations in exon 4 of the *IDH1* and *IDH2* genes and in exons 11 and 15 of the *BRAF* gene were detected using the targeted replication method (Ion AmpliSeq El Panel, Thermo Fisher Scientific) and the sequencing was performed on the Ion Gene Studio S5 Prime System (Thermo Fisher Scientific). Mutations were identified by DNA sequencing in the SeqStudio genetic analyzer (TermoFisher). 1p and/or 19q deletions were detected by FISH using the ZytoLight FISH Tissue Implementation Kit and the ZytoLight Glioma 1p/19q Probe Set. MGMT methylation, IDH and BRAF analysis were performed after sample's macrodissection.

Results

Material suitable for analysis.

The assayed sample does not carry a mutation in exon 4 of the genes IDH1 and IDH2.

Material suitable for analysis.

The assayed sample does not carry a mutation in exons 11 and 15 of the BRAF gene.

Material suitable for analysis.

The assayed sample was methylated at the promoter region of the MGMT gene.

GENEKOR | 52 Spaton Ave. Gerakas 15344 Athens | email: info@genekor.com, www.genekor.com | TEL. (+30) 210 6032138 FAX.(+30) 210 6032148

A company certified with ELOT EN ISO 9001:2015 (Cert. No 041150049) and accredited under the terms of ELOT EN ISO 15189:2012 (Cert. No. 822)

Material suitable for analysis.

In total 100 nuclei, were analyzed and the assayed sample does not carry deletion of the genetic regions 1p/19q

Scientific Director George Nasioulas, PhD Molecular Biologist

***Note: Each analysis has an internal error probability of 0,5-1%. This is due to rare events and factors involved in the production and analysis of specimens.