

Client: Example Client ABC123
123 Test Drive
Salt Lake City, UT 84108
UNITED STATES

Physician: Doctor, Example

Patient: Patient, Example

DOB 9/30/1960
Gender: Male
Patient Identifiers: 01234567890ABCD, 012345
Visit Number (FIN): 01234567890ABCD
Collection Date: 00/00/0000 00:00

1p/19q Deletion by FISH

ARUP test code 3001309

1p Result Not Deleted

19q Result Not Deleted

This result has been reviewed and approved by Deepika Sirohi,
M.D. Controls performed as expected.

H=High, L=Low, *=Abnormal, C=Critical

Unless otherwise indicated, testing performed at:

ARUP LABORATORIES | 800-522-2787 | aruplab.com
500 Chipeta Way, Salt Lake City, UT 84108-1221
Tracy I. George, MD, Laboratory Director

Patient: Patient, Example
ARUP Accession: 20-230-112409
Patient Identifiers: 01234567890ABCD, 012345
Visit Number (FIN): 01234567890ABCD
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INTERPRETIVE INFORMATION: 1p/19q, FISH

Fluorescence in situ hybridization (FISH) analysis was performed on a section from a paraffin-embedded tissue block using differentially labeled fluorescent probes targeting 1p36/1q25 and 19p13/19q13 (Abbott Molecular). Cells were evaluated from regions of tumor identified on histopathologic review of a matching hematoxylin- and eosin-stained section. Controls performed appropriately.

This assay evaluates the average ratios of 1p to 1q and 19q to 19p, as well as the percentage of cells with a signal pattern consistent with a deletion (individual cell 1p/1q and 19q/19p ratios of 0.5 or lower). Based on the validation of this assay, 1p deletion is defined as a 1p/1q ratio below 0.80 combined with a deleted pattern in 24 percent or more of the scored cells, and 19q deletion is defined as a 19q/19p ratio below 0.80 combined with a deleted pattern in 26 percent or more of the scored cells.

Codeletion of 1p and 19q as the result of an unbalanced translocation is characteristic of oligodendrogliomas and a diagnostic feature according to the WHO Classification of Tumours of the Central Nervous System, Revised 4th Edition (2016). Codeletion is also predictive of a favorable response to combination chemotherapy. Isolated deletions of 1p or 19q are neither diagnostic nor predictive in a similar fashion. Polysomy, defined in this context as three or more signals for 1q and/or 19p in 30 percent or more of the tumor cells, suggests a less-favorable outcome in oligodendrogliomas. Based on the assay performance during test validation, the test is expected to detect 96 percent of 1p and 19q deletions in patients with oligodendrogliomas. Assay range and limit of detection were generated using normal and known positive cases respectively. Correlation with other laboratory data, especially histopathologic findings, is recommended for optimal risk stratification.

References:

- Jenkins RB et al. A t(1;19)(q10;p10) Mediates the Combined Deletions of 1p and 19q and Predicts a Better Prognosis of Patients with Oligodendroglioma. *Cancer Res* 66 (20): 9852-9861, 2006.
- Snuderl M et al. Polysomy for chromosomes 1 and 19 predicts earlier recurrence in anaplastic oligodendrogliomas with concurrent 1p/19q loss. *Clin Cancer Res* 15(20):6430-6437, 2009.
- Wiens et al. Polysomy of chromosomes 1 and/or 19 is common and associated with less favorable clinical outcome in oligodendrogliomas: fluorescent in situ hybridization analysis of 84 consecutive cases. *J Neuropathol Exp Neurol* 71(7):618-624, 2012.
- Clark K et al. How molecular testing can help (and hurt) in the workup of gliomas. *Am J Clin Pathol* 139(3):275-288, 2013.
- Senetta R et al. A "weighted" fluorescence in situ hybridization strengthens the favorable prognostic value of 1p/19q codeletion in pure and mixed oligodendroglial tumors. *J Neuropathol Exp Neurol* 72(5):432-41, 2013.
- Eckel-Passow JE et al. Glioma Groups Based on 1p/19q, IDH, and TERT Promoter Mutations in Tumors. *N Engl J Med* 25;372(26):2499-508, 2015.
- Louis DN, Ohgaki H, Wiestler OD, Cavenee WK, Ellison DW, Figarella-Branger D, Perry A, Reifenberger G, von Deimling A, Eds. WHO Classification of Tumours of the Central Nervous System, Revised 4th Edition. Lyon, France: International Agency for Research on Cancer, 2016.

Test developed and characteristics determined by ARUP Laboratories. See Compliance Statement A: aruplab.com/CS

1p/1q Ratio

1.00

H=High, L=Low, *=Abnormal, C=Critical

Chromosome 1 Polysomy	Not Detected
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19q/19p Ratio	1.00
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Chromosome 19 Polysomy	Not Detected
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1p19q FISH Reference Number	1234
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1p19q FISH Source	Brain
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19Q Total Cell Count	50
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1P Total Cell Count	50
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Scoring Method	Manual
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H=High, L=Low, *=Abnormal, C=Critical

VERIFIED/REPORTED DATES

Procedure	Accession	Collected	Received	Verified/Reported
1p Result	20-230-112409	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
19q Result	20-230-112409	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
1p/1q Ratio	20-230-112409	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Chromosome 1 Polysomy	20-230-112409	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
19q/19p Ratio	20-230-112409	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Chromosome 19 Polysomy	20-230-112409	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
1p19q FISH Reference Number	20-230-112409	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
1p19q FISH Source	20-230-112409	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
19Q Total Cell Count	20-230-112409	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
1P Total Cell Count	20-230-112409	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
Scoring Method	20-230-112409	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00

END OF CHART

H=High, L=Low, *=Abnormal, C=Critical

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