



LABORATORY REPORT

Example Client, XYZ123
1234 Warde Road
Ann Arbor MI 48108

EXAMPLE, REPORT W
WX0000003827 M 07/08/1978 45 Y

Referral Testing

Collected: 09/14/2023 11:29 Received: 09/14/2023 11:29

Table with 6 columns: Test Name, Result, Flag, Ref-Ranges, Units, Site. Row 1: MPL Mutation Analysis, Clinical Indication: NA, QCRL. Row 2: Specimen Source: NA, QCRL. Row 3: Block/Specimen ID: NA, QCRL. Row 4: MPL Exon 10 Mutation: NOT DETECTED, QCRL.

Reference Range:
NOT DETECTED

Table with 6 columns: Test Name, Result, Flag, Ref-Ranges, Units, Site. Row 1: Gene: ., QCRL. Row 2: Amino Acid: ., QCRL. Row 3: Mutation Frequency: ., QCRL. Row 4: Mutation Type: ., QCRL. Row 5: Exon: ., QCRL. Row 6: Nucleotide Change: ., QCRL. Row 7: Reference: ., QCRL. Row 8: Interpretation: SEE NOTE, QCRL.

No mutation is detected in exon 10 of MPL, encompassing codons 505 and 515.

This data was reviewed and interpreted by Charles Ma, PhD. HCLD(ABB)

Table with 6 columns: Test Name, Result, Flag, Ref-Ranges, Units, Site. Row 1: Assay Details: SEE NOTE, QCRL.

This PCR-based advanced sequencing assay interrogates DNA from leukocytes for the presence of mutations in exon 10 of the thrombopoietin receptor (MPL), including codons 505 and 515. The sensitivity of mutation detection is 5% but may vary depending on the particular mutation type. Insertions up to 30bp and deletions up to 52bp have been successfully detected by the assay. Alterations outside of the tested areas of this gene will not be detected. Synonymous or known non-synonymous polymorphic changes (SNPs) are not reported. Mutations at these sites in MPL are associated with myeloproliferative neoplasms (MPNs), particularly essential thrombocythemia (ET) and primary myelofibrosis (PMF). Results of this assay should be correlated with morphology and other laboratory testing for final diagnosis and classification. If this test is negative, additional testing that may be useful for workup of MPNs, depending on presenting hematologic features, includes BCR-ABL1 rearrangement (test code 91065 or 12070X) or mutational analysis of JAK2 V617F (polycythemia vera (PV)/ET/PMF, 92473), CALR (ET/PMF, 92475), JAK2 exon 12 (PV, 92474) or CSF3R (chronic neutrophilic leukemia, 92477). Residual material from

LAB: L - LOW, H - HIGH, AB - ABNORMAL, C - CRITICAL, . - NOT TESTED



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Test Name Result Flag Ref-Ranges Units Site

this sample may be used except for BCR-ABL1 testing; call lab to add.

DNA was aligned to GRCh37(hg19) for analysis and transcript ID
ENST00000372470 was used as reference for MPL sequence.

For additional information, please refer to
(This link is being provided for informational/educational purposes
only.)

This test was developed and its analytical performance characteristics
have been determined by Quest Diagnostics Nichols Institute San Juan
Capistrano. It has not been cleared or approved by FDA. This assay has
been validated pursuant to the CLIA regulations and is used for
clinical purposes.

Test Performed at:
Quest Diagnostics Nichols Institute
33608 Ortega Highway
San Juan Capistrano, CA 92675-2042 I Maramica MD, PhD, MBA

QCRL: QUEST DIAGNOSTICS REFERENCE LAB CAPISTRANO 33608 Ortega Highway San Juan Capistrano CA 92675

Reported Date: 2023.09.14 11:30 MPLQL

LAB: L - LOW, H - HIGH, AB - ABNORMAL, C - CRITICAL, . - NOT TESTED

F314000005 Ordered By: KAJAL SITWALA, MD, PhD
WX0000003827 WX00000000002365
Printed D&T: 09/14/23 11:30

Kajal V. Sitwala, MD, PhD - Medical Director
Form: MM RL1
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