

# Test Catalog

Diagnostic. Prognostic. Predictive. Predisposition.





# **FLT3 Mutation Analysis**

#### **Alternative Name**

FLT3, FLT3 TKD, FLT3 ITD

# Methodology

Molecular

# **Test Description**

Detection of internal tandem duplication and exon 20 tyrosine kinase domain (TKD) mutations using fragment-length analysis and PCR. Positive results identify presence of TKD mutations or report ITD results quantitatively as allelic ratio.

# **Clinical Significance**

Testing for FLT3 and other gene mutations in AML patients with intermediate-risk cytogenetic abnormalities can improve risk stratification. The presence of an FLT3 mutation in a patient with AML implies aggressive disease.

#### **Specimen Requirements**

- Peripheral blood: 5 mL in EDTA tube.
- Bone marrow: 2 mL in EDTA tube.
- FFPE tissue: Paraffin block is preferred. Alternatively, send 1 H&E slide plus 5-10 unstained slides cut at 5 or more microns. Please use positively-charged slides and 10% NBF fixative. Do not use zinc fixatives.

Note: Test is suitable for Freeze & Hold option.

#### **Storage & Transportation**

Use cold pack for transport, making sure cold pack is not in direct contact with specimen.

#### CPT Code(s)\*

81245, 81246

#### Medicare MoIDX CPT Code(s)\*

81479

# **New York Approved**

Yes

#### **Level of Service**

Global

#### **Turnaround Time**

5 days

Please direct any questions regarding coding to the payor being billed.

<sup>\*</sup>The CPT codes provided with our test descriptions are based on AMA guidelines and are for informational purposes only. Correct CPT coding is the sole responsibility of the billing party.

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Committed to research as the means to improve patient care, we provide Pharma Services for pharmaceutical companies, in vitro diagnostic manufacturers, and academic scientist-clinicians. We promote joint publications with our client physicians. NeoGenomics welcomes your inquiries for collaborations. Please contact us for more information.

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