

Test Catalog

Diagnostic. Prognostic. Predictive. Predisposition.



BCR-ABL1 Standard p210, p190

Alternative Name

Philadelphia chromosome, BCR-ABL1 Major, BCR-ABL1 Minor, BCR/ABL1 Standard

Methodology

Molecular

Test Description

Real-time RT-PCR for quantitative detection of t(9;22) BCR-ABL1 fusion transcripts that result in major p210 (e13a2 and/or e14a2) or minor p190 (e1a2) fusion proteins with option to add p230 detection (micro or atypical variant). Analytical sensitivity is 0.002% for p210 and 0.005% for p190, depending on quality and quantity of the isolated RNA and absence of interfering substances. Log reduction score and percent abnormal are reported, and longitudinal data will appear as a NeoTRACK Result on the report. Testing is New York approved for p210 and p190 only. p230 testing may be ordered as a reflex if p210 and p190 are negative, or as a stand-alone test, <u>BCR-ABL1 Non-Standard p230</u>. For p230, results are reported as percent abnormal.

Clinical Significance

Useful for diagnosis and monitoring of Philadelphia chromosome-positive cases of CML and ALL. Also useful for monitoring minimal residual disease (MRD) for ALL and AML.

Specimen Requirements

- Bone Marrow: 2 mL EDTA tube. Sodium heparin acceptable.
- Peripheral Blood: 5 mL EDTA tube. Sodium heparin acceptable.

Note: Test is RNA-based, NOT suitable for Freeze & Hold option.

Storage & Transportation

Use cold pack for transport, making sure cold pack is not in direct contact with specimen. Ship same day as drawn whenever possible; specimens <7 days old preferred.

CPT Code(s)*

81206, 81207

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.

^{*}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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