

resistance to EGFR inhibitor therapies in metastatic colorectal cancer, this test is pivotal for approximately 40% of patients. Rapid results from our assay are typically available

within 48-72 hours.

Key Benefits:

- Rapid Results: Delivering actionable insights within 48 hours, supporting swift clinical decisions.
- Minimal Sample Requirement: Leveraging quantitative methodology for efficient testing.
- FDA Approved: Validated rigorously to ensure reliability and accuracy.
- Aligned with NCCN Guidelines: Part of a comprehensive approach to oncology care.

Clinical Utility

FDA Approved

Aligned with NCCN Guidelines Rapid Results Minimal Sample Requirements

Personalized Treatment Precision Medicine Early Detection

Optimizing Patient Outcomes:

The KRAS precision medicine test is customized to an individual's genetic blueprint, holds immense promise for enhancing treatment effectiveness and improving patient outcomes. By harnessing genetic data, healthcare providers can finely tune interventions and therapies to match the distinct genetic traits of each patient, resulting in actionable insights delivered in a matter of days rather than weeks.

Jones RP, Sutton PA, et. al. 2017 Specific mutations in KRAS codon 12 are associated with worse overall survival in patients with advanced and recurrent colorectal cancer. Br J Cancer. Mar 28;116(7):923-929

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