BRAF therascreen

BRAF V600 Mutation Analysis

The BRAF therascreen empowers oncologists by pinpointing the BRAF V600E mutation, prevalent in melanoma and colorectal cancer (CRC). Generally, BRAF V600 mutations are found in approximately 5-15% of CRC cases. This critical genetic insight facilitates personalized treatment plans, particularly in metastatic CRC, where it serves as a beacon for selecting patients for BRAFTOVI (encorafenib) in combination with cetuximab therapy. **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 Precision: Designed for metastatic colorectal cancer patients, enhancing treatment selection with BRAFTOVI® and Cetuximab (ERBITUX®) based on BRAF V600E mutation status.

Optimizing Patient Outcomes:

Clinical Utility

FDA Approved

Aligned with NCCN Guidelines Rapid Results Minimal Sample Requirement

Personalized Treatment Precision Medicine Early Detection

The BRAF 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.

For more information, please contact us at: **cs@emeritusdx.com**





