Cleveland Clinic

Cleveland Clinic Laboratories

Solid Tumor Gene Fusion Next-Generation Sequencing Panel

Background

Benign and malignant mesenchymal tumors (sarcomas and their mimics) are difficult to diagnose with many benign and malignant entities that differ in their clinical behavior and response to therapy. Many of these tumors harbor gene fusions that are crucial to establishing a definitive diagnosis. The Solid Tumor Gene Fusion Next-Generation Sequencing (NGS) Panel is a custom designed, 58-gene panel, high complexity laboratory developed test (LDT) designed for targeted sequencing of benign and malignant solid and soft tissue neoplasms. This assay identifies fusion transcripts in targeted regions of RNA from total nucleic acid (TNA) isolated from formalin-fixed, paraffin-embedded (FFPE) tissue specimens.

The test will identify the vast majority of known fusions in benign and malignant mesenchymal tumors, but also has the ability to identify a limitless number of as-of-yet undiscovered gene fusions. This technology only "primes" from one partner of the gene fusion, allowing for discovery of new gene fusion partners.

Clinical Indications

This test is intended for the diagnosis of benign or malignant mesenchymal tumors (sarcomas and their benign mimics) as well as other solid tumors.

Interpretation

The results of this test are to be interpreted in the context of the histological, immunohistochemical, and clinical features of the neoplasm.

Methodology

This test relies on Anchored Multiplex PCR (AMP™) technology to generate scalable, target-enriched libraries for NGS from formalin-fixed, paraffin embedded tissue sections. In AMP, unidirectional gene-specific primers (GSPs) are used to enrich libraries for known and unknown mutations. Adapters that contain both molecular barcodes and sample indices enable quantitative multiplex data analysis, read de-duplication, and accurate variant calling. Libraries are sequenced on the Illumina MiSeq instrument, which employs

Highlights of Solid Tumor Gene Fusion NGS Panel

- Comprehensive detection of gene fusions across 58 targeted genes aids in determining diagnosis, prognosis, and therapeutic options.
- FFPE tissue removes need to send fresh or frozen specimens.

The targeted genes included in the panel are:

Solid Tumor Gene Fusion NGS Panel (SRCNGS) – 58 Genes

ALK	*CSF1*	*FUS*	*NCOA1*	*NTRK1*
BCOR	*EPC1*	*GLI1*	*NCOA2*	*NTRK2*
BRAF	*ETV6*	*HMGA2*	*NCOA3*	*NTRK3*
CAMTA1	*EWSR1*	*JAZF1*	*NOTCH1*	*NUTM1*
CCNB3	*FOS*	*MEAF6*	*NOTCH2*	*PAX3*
CIC	*FOSB*	*MKL2*	*NOTCH3*	*PAX7*
CRTC1	*FOXO1*	*MYB*	*NR4A3*	*PDGFB*
PDGFD	*RAF1*	*STAT6*	*TRIM11*	
PGR	*RELA*	*TAF15*	*USP6*	
PHF1	*RET*	*TCF12*	*WWTR1*	
PLAG1	*ROS1*	*TFE3*	*YAP1*	
PRDM10	*SRF*	*TFEB*	*YWHAE*	
PRKD1	*SS18*	*TFG*		

Subpanel: Head & Neck Gene Fusion (HDNK) – 29 Genes

ALK	*FOS*	*MKL2*	*NTRK3*	*RET*
BRAF	*FOSB*	*MYB*	*NUTM1*	*SS18*
CAMTA1	*FOXO1*	*NCOA1*	*PAX3*	*STAT6*
CRTC1	*FUS*	*NR4A3*	*PAX7*	*TFE3*
ETV6	*GLI1*	*NTRK1*	*PLAG1*	*YAP1*
EWSR1	*HMGA2*	*NTRK2*	*PRKD1*	

Subpanel: NTRK Gene Fusion (NTRK) – 3 Genes

NTRK1 *NTRK2* *NTRK3*

"sequencing by synthesis;" a fluorescence, image-based, reversible-terminator technology to sequence targeted regions of the 58 genes included in the panel.

Sequencing data are analyzed for fusion variant detection using Archer® Analysis bioinformatics tools. Specimen quality control is monitored and recorded by in-house developed software (scripts). Raw sequencing data are de-multiplexed based on unique index sequence using the Illumina bcl2fastq program. The fastq.gz files are de-duplicated according to the unique molecular barcode present and aligned to the human reference genome hg19. Part of the fusion calling and annotation is performed utilizing the Archer® Quiver™ Fusion Database.

Limitations of the Assay

This test does not detect missense mutations, insertions, deletions, or copy number changes, and does not distinguish between variants that are inherited versus acquired.

References

1. Archer Dx, FusionPlex Anchored MultiPlex PCR (AMP) technology http://archerdx.com/fusionplex/ [Accessed: July 2018]
2. MiSeq System user Guide, Publication Number 15027617 Rev.0. *Illumina*, San Diego, CA. 9/2014.
3. NCCN Clinical Practice Guidelines in Oncology (NCCN Guidelines®), *Soft Tissue Sarcoma*, version 1.2019
4. Taylor BS, Barretina J, Maki RG, Antonescu CR, Singer S, Ladanyi M. Advances in sarcoma genomics and new therapeutic targets. *Nat. Rev. Cancer*. Jul 14 11(8), 541-57 (2011).

Test Overview

Test Name	Solid Tumor Gene Fusion NGS Panel
Ordering Mnemonic	SRCNGS
Methodology	Next-Generation Sequencing
Specimen Requirements	Formalin-fixed, paraffin-embedded (FFPE) tissue: • Ten (10) unstained, 4 μM sections of FFPE on charged, unbaked slides • One (1) H&E stained slide with best tumor area circled by a pathologist (minimum of 20% tumor content for best results)
Stability	Ambient: Transport and store slides at ambient temperature. Frozen: Unacceptable Refrigerated: Unacceptable
Days Performed	1 day per week
Days Reported	14 days
CPT Codes	81445, 88381

Technical Information Contact:

Kristen G. McDonnell, MB (ASCP)^{CM}CG^{CM}
216.314.1008
mcdonnk3@ccf.org

Scientific Information Contacts:

Brian Rubin, MD, PhD
rubinb2@ccf.org
216.445.5551

Elizabeth Azzato, MD, PhD
azzatoe@ccf.org
216.444.8116