

OncoTrack CGP (Liquid Biopsy)

Comprehensive Genomic Profiling of Solid Tumors from Blood

High precision multibiomarker test

SNVs & InDels - 523 genes | CNVs - 59 genes | Fusion - 23 genes | MSI | TMB



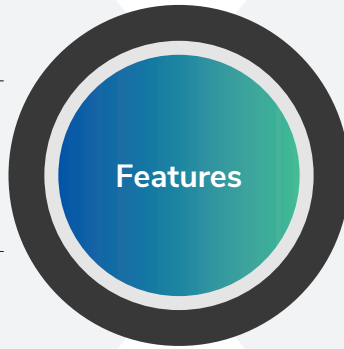
Provides comprehensive genomic profiling from blood sample for all solid tumors



Achieve low limits of detection with UMI-based hybrid-capture library preparation and deep sequencing.



Sophisticated variant calling algorithms and high depth of sequencing enable detection of key biomarkers in cfDNA with 0.5% limit of detection (LOD)



Leverage minimally invasive blood samples as a complement to tissue biopsy or as an alternative when tissue is not readily available



Detects low-frequency somatic variants across 523 genes with high sensitivity and specificity



Well - validated as per CAP guidelines. High throughput Illumina's sophisticated NGS platforms



Use in Clinical Practice

- Advanced solid tumors
- Before first-line therapy or at progression

Biomarkers

Single Nucleotide Variants (SNVs)

Small Insertions (<20bp)

Small Deletions (<20bp)

Multinucleotide Variants (MNVs)

Gene Amplifications

Gene Deletions

Gene Rearrangements (Fusions)

Tumor Mutational Burden (TMB)

Microsatellite Instability (MSI)

