

# Essential LBx Panel

The **oncoReveal™ Essential LBx** panel is a robust NGS assay that interrogates 197 targets across 34 genes of interest\* from multiple solid tumor cancer types. This is a focused panel that can detect four types of variants from cell-free DNA (cfDNA) that has been extracted from plasma: single nucleotide variants (SNVs), small insertion/deletion (indel) variants, copy number amplification (CNA), and microsatellite instability (MSI). The panel uses proprietary Stem-Loop Inhibition-Mediated amplification (SLIMamp®) technology, a tiled amplicon-based library prep chemistry for efficient single-tube target enrichment.

## oncoReveal™ Essential LBx Panel (34 genes)

AKT1	CTNNB1	FGFR2	KIT	PPP2R1A	SF3B1
ALK	EGFR	FGFR3	KRAS	PTCH1	SMAD4
AR	▲ ERBB2	GNA11	▲ MET	PTEN	TERT (promoter)
ATM	ERBB3	GNAQ	NRAS	RAC1	<b>TP53</b>
BRAF	ESR1	GNAS	PDGFRA	RET	
CDK4	▲ FGFR1	HRAS	PIK3CA	RNF43	

Copy Number Amplifications (CNAs) can also be detected in genes indicated by ▲ Genes marked in green indicate full CDS coverage

### Simple NGS library prep workflow

Maintain control of samples and results with single-tube, tiled amplification that can be performed in-house by any NGS lab

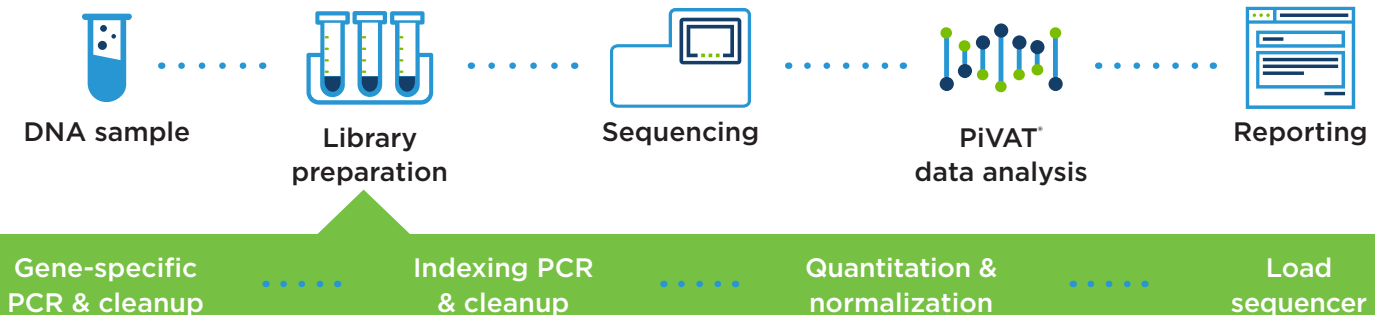
### Sensitive and robust chemistry

Achieve variant detection as low as 0.1% VAF<sup>†</sup> even with limited DNA input or poor sample quality

### Reduced fully-loaded lab costs

Improve lab efficiency and reduce “no calls”, repeat testing, and difficult interpretation decisions

### Simple, one-day workflow



\* Content is based on data from ongoing clinical trials along with sources that include the College of American Pathologists (CAP), the Association for Molecular Pathology (AMP), the National Comprehensive Cancer Network (NCCN), and the Catalog of Somatic Mutations in Cancer (COSMIC) database.

<sup>†</sup>VAF, variant allele frequency  
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## Panel specifications\*

Enrichment chemistry	Multiplex PCR using tiled amplicons
Number of pools	1 pool
Number of genes/amplicons	34/197
Number of targets	Hotspots in 33 genes; full CDS coverage of TP53; 27 MSI sites; CNAs in 3 genes
Variant types	SNVs, indels, CNAs, and MSI
Average amplicon size	90bp
Recommended DNA input range	10ng to 30ng
Sample types	cfDNA from plasma
Mapping rate	≥90%
% on-target aligned reads	≥90%
Coverage uniformity (% targets with >0.2X mean coverage)	≥90%
Recommended Reads Per Sample	~10 million paired-end reads
Total assay time (from DNA to sequencer)	<8 hours

\* Mapping rate, percentage of on-target aligned reads, and coverage uniformity metrics are based on internal testing performed using reference standard materials

## Ordering information

Select the panel AND one of the index kit options listed below.

Panel	Part number
oncoReveal™ Essential LBx Panel (24 reactions)	HLA-HS-1006-24

Pillar Unique Dual Index Kit Options	Reactions	Part number
Pillar Biosciences LBx Indexing Kit A	24 Combinations, 96 reactions	IDX-PI-1013-96
Pillar Biosciences LBx Indexing Kit B	24 Combinations, 96 reactions	IDX-PI-1014-96

TO ORDER OR LEARN MORE:  
[pillarbiosci.com](http://pillarbiosci.com)

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