

EPIPLEX RNA REAGENT KIT

Sample#	Reagents for 8 samples (includes 8 enrichment and 8 solution reactions)
Price	\$2,800; early access discount available
RNA modifications	m6A, inosine
Binding reagents	engineered small protein scaffolds
RNA species	poly-A(+) RNA, total RNA (rRNA depletion at cDNA step recommended)
RNA amount	>= 20 ng polyA+ RNA >= 250 ng total RNA
Library type	stranded RNA library
Quantification	fold-enrichment relative to spike-in controls
Sample definition	an enrichment reaction with a paired solution reaction
Sequencing depth	>= 20 - 50M reads per sample*, 200 cycles
Analysis	Alida Bio's analysis pipeline
Supported genomes	human, mouse – others per request
Turnaround time	Library prep using reagent kit: 7 hours

* Sequencing depth depends on sensitivity expectation. Prominent modification sites are accessible at lower coverage.

DATA ANALYSIS OUTPUT

PEAK LOCATIONS	BED FILES
<i>Transcript regions ("peaks") with RNA modifications for visualization in any genome viewer.</i>	
COVERAGE TRACKS	BIGWIG FILES
<i>Read coverage corrected for the non-enriched input coverage and normalized to spike-in controls for visualization in any genome viewer. One file per RNA modification.</i>	
GENOME ALIGNMENT	BAM FILES
<i>Aligned and deduplicated reads for visualization in any genome viewer. One file per RNA modification.</i>	
RAW SEQUENCING DATA	FASTQ
<i>Unprocessed sequencing reads, demultiplexed per sample for publication and data storage.</i>	
SUMMARY OUTPUTS	HTML, TSV, CSV, PNG
<i>RNA-seq TPM values Peaks table with peak location, annotated gene features, #reads per peak, fold-enrichment, q-values, and more Fold-enrichment correlation plots of user-defined conditions Modification distribution along the transcript DRACH motif enrichment (m6A) and A-to-G (inosine) mutations under peaks</i>	