



## **KAPA SARS-CoV-2 Target Enrichment Panel**

Better by Design

Monitor the presence of the SARS-CoV-2 virus and the emergence of new strains with the **KAPA SARS-CoV-2 Target Enrichment Panel**. Prepare libraries from input RNA using the robust **KAPA RNA HyperPrep Kit** and then enrich for viral sequences; this panel targets 100% of the reference SARS-CoV-2 genome (NC\_045512) and >99.7% of another 183 publicly available SARS-CoV-2 genomes (GenBank, March 2020).

- Identify multiple variants of SARS-CoV-2 in a single reaction\*
- Achieve 1X coverage of >97% of the SARS-CoV-2 genome down to 1000 viral copies and obtain genomic sequence from as few as 10 viral copies (in a background of 20 ng or 100 ng RNA with 1 million 2x75 bp reads)
- Save valuable time with hybridization as short as 1 hour



### Identify multiple variants of SARS-CoV-2 in a single reaction

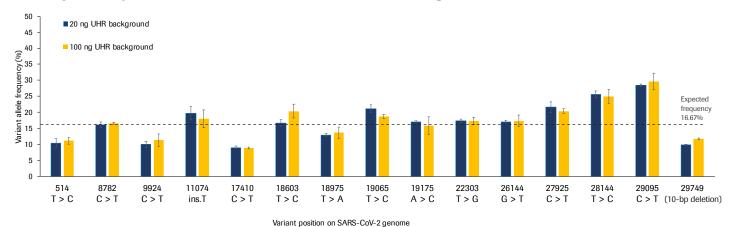


Figure 1. Variants from six strains of SARS-CoV-2 are identified from within a single sample. A total of 1,000,000 SARS-CoV-2 genome copies from 6 different strains was combined with either 20 ng (dark blue) or 100 ng (yellow) of human RNA and processed in triplicate using KAPA RNA HyperPrep and the KAPA SARS-CoV-2 Target Enrichment Panel. Alternate allele frequency analysis shows that all expected variants from the six SARS-CoV-2 strains were identified at close to the expected frequency. Datasets downsampled to 1 million reads prior to analysis.

# Cover more of the SARS-CoV-2 genome and identify viral sequence from low viral copy numbers

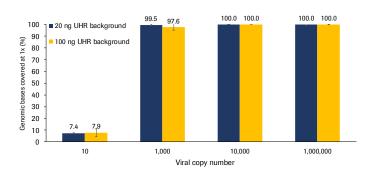
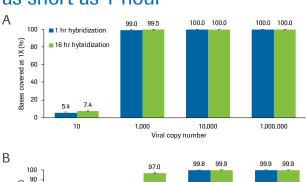


Figure 2. The KAPA SARS-CoV-2 Target Enrichment Panel achieves 1X coverage of >97% of the SARS-CoV-2 genome down to 1000 viral copies and genomic sequence from as few as 10 viral copies. Samples containing the indicated number of viral copies in a background of either 20 ng or 100 ng of human RNA were processed in triplicate using KAPA RNA HyperPrep and the KAPA SARS-CoV-2 Target Enrichment Panel. Datasets were downsampled to 1 million reads prior to analysis.



# Save valuable time with hybridization as short as 1 hour



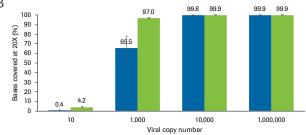


Figure 3. Shortening the hybridization time to 1 hour yields equivalent 1X SARS-CoV-2 genome coverage compared to 16-hour hybridization. Libraries generated from samples containing the indicated number of viral copies in a background of 20 ng of human RNA were hybridized to the SARS-CoV-2 TE panel for 1 hour or 16 hours and processed through the HyperCap v3 workflow in triplicate. A. The 1-hour hybridization time and 16-hour hybridization time yield similar SARS-CoV-2 genome coverage at 1X for all viral copy levels. B. Samples with higher viral loads also yield similar genome coverage at 20X for the 1-hour and 16-hour hybridization times, with somewhat reduced coverage for lower viral loads. 1 million Illumina NextSeq reads (2x75 bp).



#### Learn more at:

https://go.roche.com/SARS-CoV2-Panel

Request a no-charge evaluation of the SARS-CoV-2 Enrichment Panel at: https://go.roche.com/SARS-CoV2-Eval