

# Maximize CONSISTENCY

## KAPA HyperPure Beads: Attract what matters.

In the NGS workflow continuum, sample prep holds the key to unlocking the potential of every sample. Roche Sample Prep Solutions provide a **dynamic**, **reliable**, and **focused** portfolio of high-performance, high-quality library preparation reagents that enable you to process more samples successfully, obtain more information from every sample, and optimize sequencing resources.

This includes **KAPA HyperPure Beads** that offer a tunable and highly consistent solution for reaction purification and size selection in next-generation sequencing DNA library construction workflows.

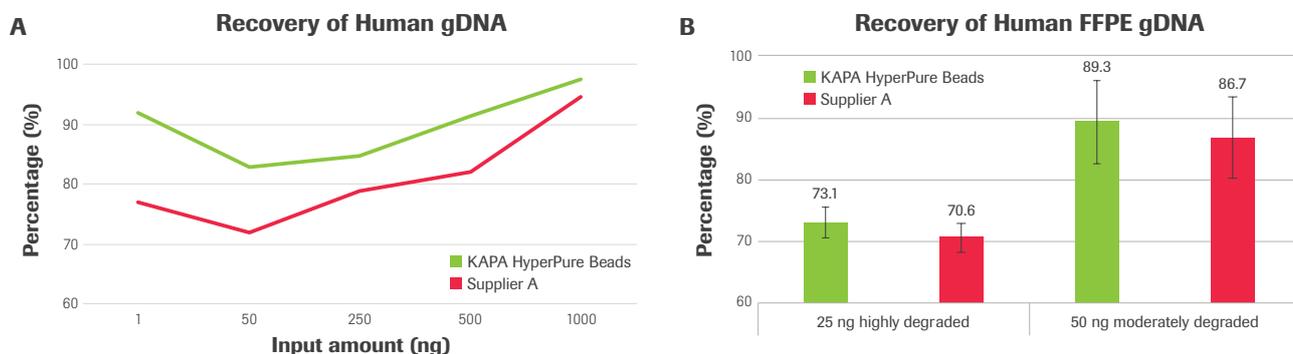
## Benefits

- KAPA HyperPure Beads offer best-in-class nucleic acid recovery for many KAPA NGS DNA workflows, providing tunable and consistent size selection
- Improved bead wash efficiency maximizes library diversity and reduces sequencing costs
- Fast, efficient reaction cleanups remove adapters, adapter-dimers, primers, primer-dimers, nucleotides, salts, and enzymes
- KAPA HyperPure Beads are ready-to-use and automation-friendly



## Industry-leading recovery of DNA

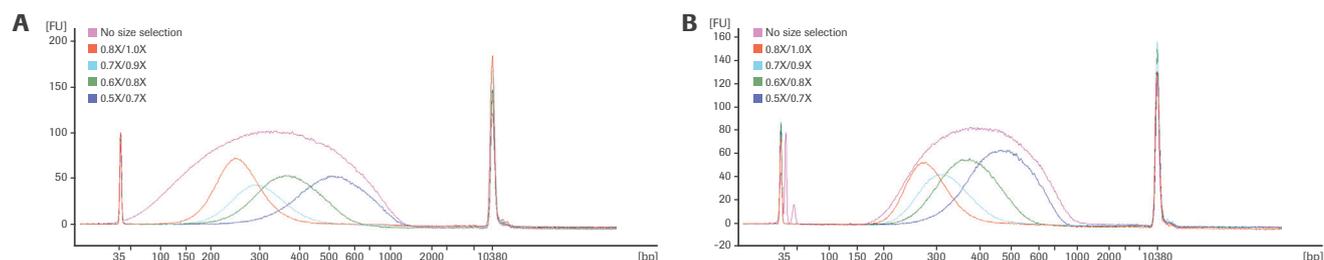
- KAPA HyperPure Beads achieve comparable or better DNA recovery than the market leader (Supplier A) across a broad spectrum of input amounts, sizes, and quality
- Improved general manipulation of DNA samples, e.g., buffer exchange to remove EDTA or other salts prior to enzymatic fragmentation and sample concentration



**Figure 1. KAPA HyperPure Beads provide superior recovery performance to Supplier A in DNA workflows. (A)** Recovery of unfragmented human gDNA. **(B)** Recovery of highly degraded and moderately degraded formalin-fixed paraffin-embedded (FFPE) human genomic DNA. Highly degraded DNA obtained from FFPE clinical research samples ( $n=12$ ); moderately degraded DNA obtained from Horizon Reference FFPE DNA ( $n=9$ ). For both **(A)** and **(B)**, KAPA HyperPure Beads and Supplier A beads were used at a 3X ratio to clean up DNA inputs of various amounts. Recovery was measured using the Qubit Fluorometer 3 dsDNA HS Assay Kit before and after cleanup.

## Achieve tunable and highly reproducible size selection

- Reproducible size selection of nucleic acids—capture of fragmented input DNA, adapter-ligated molecules or amplified libraries in NGS library preparation workflows
- Enables researchers to consistently select for fragment sizes of interest



**Figure 2. KAPA HyperPure Beads provide tunable size selection of double-stranded DNA.** At each step of the KAPA HyperPrep library preparation workflow, DNA was subjected to double-sided size selection using different bead-to-sample ratios (see legend) and then analyzed for size distribution with a Bioanalyzer® 2100 High Sensitivity Kit. Each panel shows the resulting fragment size ranges compared to same-stage DNA that was not size-selected. DNA concentrations were normalized prior to loading onto the Bioanalyzer. **(A)** Post-fragmentation size selection of input DNA. High-quality hgDNA was mechanically fragmented with a Covaris® E220 Focused Ultrasonicator using conditions optimized to yield a mode fragment length of 250 – 400 bp, then subjected to size selection. **(B)** Post-amplification size selection. Libraries were amplified and final libraries were then subjected to size selection.

### Ordering Information

Roche cat. no.	KAPA code	Description	Kit size
08963835001	KK8007	KAPA HyperPure Beads (5 mL)	5 mL
08963843001	KK8008	KAPA HyperPure Beads (30 mL)	30 mL
08963851001	KK8009	KAPA HyperPure Beads (60 mL)	60 mL
08963878001	KK8011	KAPA HyperPure Beads (4 x 60 mL)	4 x 60 mL
08963860001	KK8010	KAPA HyperPure Beads (450 mL)	450 mL

Published by:

**Roche Sequencing Solutions, Inc.**  
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Pleasanton, CA 94588

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06/22

