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Prepare 5% extra reagent mix if you are preparing multiple samples. 2 Remove the PCR plate from magnetic stand 3 Add 30 µl of the elution pre mix to each well of the PCR plate using a single channel or multichannel pipette. 4 Gently pipette the entire volume of each well up and down 10–20 times to mix thoroughly. Ensure that the beads are fully resuspended. 5 Seal the PCR plate with a Microseal ` B ` adhesive seal.

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The TruSeq Exome Kit supports 12-plex pre-enrichment library pooling, enabling researchers to maximize sequencing throughput and variant identification by sequencing up to 12 libraries per flow cell lane. This enables sequencing of more exomes per run, so researchers can maximize their budgets.

TruSeq DNA Exome (formerly TruSeq Exome Kit)

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Illumina RNA Prep with Enrichment is designed to provide exceptional capture efficiency and coverage uniformity, minimizing required sequencing depth for accurate detection without bias. Only 2.5 µL probes are needed per enrichment reaction.

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