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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 μ l 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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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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