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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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TruSeq DNA Exome (formerly TruSeq Exome Kit)
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Procedure. 1 [Plate]ArrangeIndex1(i7)adaptersincolumns1-

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1 2 of the TruSeq Index Plate Fixture. 2 [Plate] Arrange Index 2 (i5) adapters in rows A–H of the TruSeq Index Plate Fixture. 3 [Plate] Place the plate on the TruSeq Index Plate Fixture. 4 Add 5 μ l of each Index 1 (i7) adapter as follows.

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