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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 ?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)adaptersincolumns1–12oftheTruSeqIndexPlate Fixture. 2 [Plate]ArrangeIndex2(i5)adaptersinrowsA–HoftheTruSeqIndexPlateFixture. 3 [Plate]PlacetheplateontheTruSeqIndexPlateFixture. 4 Add5?lofeachIndex1(i7)adapterasfollows.

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