AMPINEXT™ High-Sensitivity DNA Library Preparation Kit (Illumina)

The AMPINEXT'S etigh Sensitivity 2014 Library Preparation Kit (Illumina) is a complete set of optimized reagents to prepare a DNA library from very prepare a DNA library from very small smount of samples for use in next-generation sequencing applications. This kin is satisfied to preparing a DNA library using subgenerations using an Illumina sequencer. These applications include genomic DNA-seq, ChIP-seq, MeDIP/hMeDIP-seq, traditional bisulfite-seq, and targeted re-sequencing. The optimized protocol and components of the kit allow both non-barcoded (singleplexed) and barcoded (multiplexed) DNA libraries to be constructed quickly with reduced bias.

Ordering Information

Order Online »

ENZ-GEN505-0012	12Reactions
ENZ-GEN505-0024	24Reactions

Manuals, SDS & CofA

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- High sensitivity and flexibility -Use with non-barcoded (singleplexed) and barcoded (multiplexed) DNA library preparation.
- Fast and streamlined procedure
 - No clean-up is required between each step and all reactions take place in the same tube.
- Highly convenient The kit contains all required components for each step of DNA library preparation,
- Minimized bias Ultra HiFi amplification and optional PCRfree step.

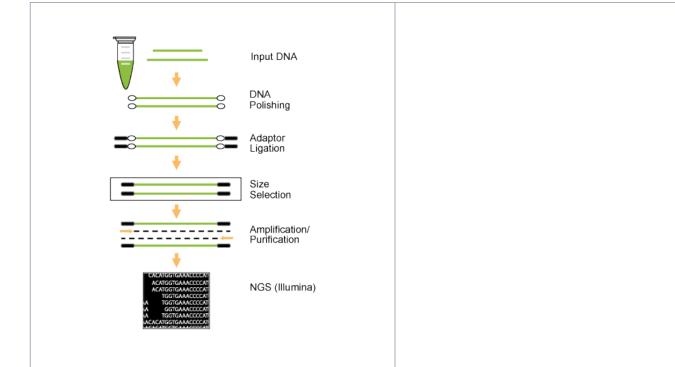


Figure 1. Schematic Procedure for using the AMPINEXT High-Sensitivity DNA Library Preparation Kit

Handling & Storage

Use/StabilityUpon receipt: Store the following components at -20°C immediately: 10X End Polishing

Buffer, End Polishing Enzyme Mix, End Polishing Enhancer, 2X Ligation Buffer, T4 DNA Ligase, Adaptors, 2X HiFi PCR Master Mix, Primer U, Primer I, and Elution Buffer. Store the following components at 4°C: MQ Binding Beads. Store all other components at

room temperature.

Shipping Blue Ice

Regulatory Status RUO - Research Use Only

Product Details

Application Notes A complete set of optimized reagents to prepare a DNA

library from very small amount of samples for use in next-

generation sequencing applications.

Assay Time 1 hour 30 minutes

Contents 10X End Polishing Buffer

End Polishing Enzyme Mix

End Polishing Enhancer

2X Ligation Buffer

T4 DNA Ligase

Adaptors (50 µM)

MQ Binding Beads

2X HiFi PCR Master Mix

Primer U (10 µM)

Primer I (10 µM)

Elution Buffer

Technical Info / Product Notes

Starting materials can include fragmented dsDNA isolated from various tissue or cell samples, dsDNA enriched from a ChIP reaction, MeDIP/hMeDIP reaction, or exon capture. DNA should be relatively free of RNA because large fractions of RNA will impair end repair and dA-tailing, resulting in reduced ligation capabilities. The input amount of DNA can be from 0.2 ng to 100 ng. For optimal preparation, the input amount should be 10 ng to 50 ng.

