

## NGS Ligation Enzyme

Catalog No.	40105S	40105L
Reactions	24 reactions	96 reactions

### Description

The **NGS Ligation Enzyme** was developed for preparation of high-quality libraries for next generation sequencing. The reagent was derived from the ligation step of our NGS library prep kits. The reagent contains Ligation Enzyme and Ligation Buffer with enhancers for high library conversion efficiency.

The proprietary components in the buffer improve ligation of NGS adaptors to dA-tailed DNA fragments dramatically. The ligation is performed at room temperature (20°C) for only 20 minutes with high efficiency. The enzyme is also optimized for use with the BioDynami **End Repair/dA-Tailing Master Mix** (Cat.# 40104S; 40104L).

### Features

- Fast: 20 minutes ligation at room temperature
- Simple reaction setup
- Higher library conversion efficiency
- Input DNA amount: From 1 ng to 1 ug

### Component

Catalog No.	40105S	40105L
Ligation Buffer	336 ul	1344 ul
Ligation Enzyme	48 ul	192 ul

### Storage Condition

- Store kit at -20°C, stable up to 12 months.

### Reagent & Equipment Needed (not provided in this kit)

- Magnetic particle concentrator
- PCR thermal cycler
- 96-well PCR plate
- 80% ethanol (prepare before use)
- Magnetic Beads (BioDynami Cat.# 40051) or equivalent

## Protocol

### Adaptor Ligation

- 1) Add the following to one well of a 96-well PCR plate. Slow pipetting of the viscous **Ligation Buffer** is needed for precise aliquot.

DNA sample	30 ul
Adaptor	4 ul
Ligation Buffer	14 ul
<u>Ligation Enzyme</u>	<u>2 ul</u>
Total	50 ul

- 2) Mix by pipetting ten times.
- 3) Incubate at 20°C for 20 min.

### Subsequent Magnetic Beads Purification (Recommended)

- 1) Resuspend **Magnetic Beads** and transfer 40 ul to the above reaction mixture, mix by pipetting and incubate for 3 min.
- 2) Load the plate on a magnet, incubate for 2 min, and discard the supernatant carefully.
- 3) Add 180 ul of 80% ethanol, incubate for 30 sec, and discard the supernatant carefully. Remove all residual ethanol without disturbing the beads.
- 4) Remove the plate from the magnet, resuspend the beads in 22 ul of water or Tris-HCl (10 mM).
- 5) Load the plate on the magnet, incubate for 1 min, and transfer 20 ul supernatant (containing library) to a new tube without disturbing the beads.

### Quality Control

Kit components passed stringent functional quality test.

### Product Use Limitation




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