

Related products

| Product | Amount | Cat. no. |
|--|---------------|-----------|
| Anza™ T4 DNA Ligase Master Mix | 50 reactions | IVGN210-4 |
| Anza™ Alkaline Phosphatase Kit | 500 reactions | IVGN220-4 |
| Anza™ T4 PNK Kit | 50 reactions | IVGN230-4 |
| Anza™ DNA End Repair Kit | 20 reactions | IVGN250-4 |
| PureLink™ PCR Purification Kit | 50 preps | K3100-01 |
| One Shot™ TOP10 Chemically Competent <i>E. Coli</i> | 20 reactions | C4040-03 |
| One Shot™ INV110 Chemically Competent <i>E. Coli</i> | 20 reactions | C7171-03 |

To order additional Anza™ Restriction Enzymes and Anza™ Modifying Cloning Enzymes, go to thermofisher.com/Anza

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7 June 2016

ThermoFisher
S C I E N T I F I C

Anza™ DNA Blunt End Kit

| Cat. No. | Size | Lot no. | Exp. Date |
|-----------|---------------|---------|-----------|
| IVGN240-4 | 100 reactions | | |

Publication No. MAN0014312 Rev. B.0

Product description The Invitrogen™ Anza™ DNA Blunt End Kit is used to convert DNA with cohesive ends to blunt ended DNA for blunt end ligation.

The Anza™ DNA Blunting Enzyme Mix contains T4 DNA polymerase and Klenow Fragment.

The Anza™ 10X Blunting Buffer contains dNTPs to facilitate the synthesis of blunt ends.

| Components | Amount |
|---------------------------|--------|
| Anza™ DNA Blunting Enzyme | 100 µL |
| Anza™ 10X Blunting Buffer | 200 µL |

Storage Store at –20°C.

For research use only. Not for use in diagnostic procedures.

General guidelines

- DNA digested with Anza™ Restriction Enzymes can be used directly in the protocol following heat inactivation.

DNA blunting protocol

Use this protocol to convert sticky-ended DNA to blunt-ended DNA for insertion into a dephosphorylated blunt ended vector.

1. Prepare a reaction mix by adding the reagents listed in the following table to a clean microcentrifuge tube:

| Reagent | Volume |
|-------------------------------|--|
| Nuclease-free water | As required to reach final reaction volume |
| Anza™ 10X Blunting Buffer | 2 µL |
| DNA insert | 0.2–1 µg |
| Anza™ DNA Blunting Enzyme Mix | 1 µL |
| Final reaction volume | 20 µL |

2. Mix reagents by pipetting up and down.
3. Incubate at 20°C for 15 minutes.
4. Purify DNA insert from reaction mix using the PureLink™ PCR Purification Kit.
5. Ligate insert and vector (dephosphorylated and blunt ended) using the Anza™ T4 DNA Ligase Master Mix.
6. Use 1–5 µL of the ligation reaction mixture to transform competent cells.