# **TOP10** Chemically Competent Cell

# **Product Specification**

Catalog Number	Specification
CS02010	10×100µL
CS02020	20×100µL

#### **Product introduction**

The product are the competent cells obtained from special treatment of Escherichia coli TOP10 strain, and can be used for DNA thermal shock conversion. TOP10 is a kind of strain which can be commonly used in plasmid cloning, its 80lacZ M15 product can be complementary to the amino terminus of the beta galactosidase that encoded by the vector, and also can be used for blue-white spot screening.

Using pUC19 plasmid to detect, the conversion efficiency can up to 10<sup>8</sup>, which is suitable for efficient cloning of plasmid DNA and can ensure stable replication of high copy plasmid.

## Protocol

1. Take the TOP10 chemically competent cells from - 80 °C, inserted rapidly in the ice, waiting for 5 minutes for the fungus block to melt, add the target DNA (plasmid or connection product) and then gently blowing it with pipette place in the ice for 5 minutes.

2. Heat shock 90 seconds in 42  $^\circ\!\mathrm{C}$  water bath , quickly put back on ice and put it aside for 5 minutes.

3. Add 500  $\mu$ L of antibiotic-free sterile medium (SOC or LB medium) to the centrifuge tube and mix for 37 minutes at 200 ° C for 60 minutes.

4. 3000 RPM instantaneous centrifugal harvesting bacteria, with a total of 100 mu L left and to gently blow the resuspension bacteria and smear it to the LB medium containing the selected plasmid with screening antibiotics

5. Upside down the plate and place it in a incubator at 37°C over night.

## Notes

1. It is better to slowly melt competent cells in the ice, not to place cells in the ice too long, long storage will reduce the conversion efficiency.

2. Gentle operation should be performed when mixing the plasmid.

3. Conversion of high concentration of plasmids can reduce the amount of bacteria that eventually to be used for plating.