

EHA101 Chemically Competent Cell



Cat. No. ACC-108

Lot. No. (See product label)

Product Name

EHA101 Chemically Competent Cell

Product Overview

The genotype of EHA101 Chemically Competent Cell is C58 (rif^R) Ti pEHA101 (pTiBo542 D T-DNA) (kan^R) Nopaline. Background of EHA101 strain is C58. It contains rifampicin resistant gene (rif) in nuclear genes as screening label. EHA101 carries amber basic Ti plasmid pEHA101 (pTiBo542DT-DNA) to facilitate transformation. Ti plasmid pEHA101 (pTiBo542DT-DNA) contains vir gene, which is essential for insertion of T-DNA into plant genome. Ti plasmid pEHA101 (pTiBo542DT-DNA) is disabled to transfer its own T-DNA but enabled to transfer foreign binary vector T-DNA. pEHA101 (pTiBo542DT-DNA) Ti plasmid contains Kanamycin resistance gene.

Applications

EHA101 Chemically Competent Cell is suitable for transgenic operations of corn, rice, tobacco and other plants.

Notes

1. Volume of DNA from ligation mix should not exceed 1/10 of the cell mixture; DNA for transformation should be purified and free of organic substances such as ethanol.
2. Do not pipette or vortex cells.
3. Plating volume can be adjusted accordingly.
4. Please avoid excessive use of rifampicin. Maximum concentration of rifampicin in selection is 25 µg/mL.

Kit Components

EHA101: 100 µL/tube * 10 tube/50 tube/100 tube.

Assay Protocol

1. Thaw Agrobacterium competent cells at room temperature or in the palm, and place in ice bath.
2. Add 0.01-1 µg of plasmid DNA to 100 µL of competent cells. Carefully flick the tube to mix cells and DNA. Do not pipette or vortex.
3. Place the tube on ice for 5 minutes, in liquid nitrogen for 5 minutes, in 28°C or 37°C water bath for 5 minutes, and in ice bath for 5 minutes.
4. Add 700 µL of antibiotic-free LB or YEB medium to the mixture and shake for 2 to 3 hours at 28°C.
5. Centrifuge culture at 6000 rpm for 1 minute and dispose ~700 µL supernatant. Resuspend cell pellet in the rest 100 µL medium.
6. Spread 50-100 µL cell suspension to LB or YEB plate containing proper antibiotics and incubate at 28°C for 2 - 4 days.

Transformation efficiency

Transformation efficiency of EHA101 Chemically Competent Cell using pK7WGF2 plasmid (spectinomycin resistance) with 50 µg/mL spectinomycin is >10⁴ cfu/µg DNA. Transformation efficiency is reduced to half when the plate contains 50 µg/mL spectinomycin and 20 µg/mL rif.

Storage

Store at - 80 °C for 12 months.

FOR RESEARCH OR FURTHER MANUFACTURING USE ONLY