

TransDB3.1 Chemically Competent Cell

Cat. No. CD531

Storage: at -70°C for six months. Do not store in liquid nitrogen.

Description

*Trans*DB3.1 Chemically Competent Cell is specifically designed for chemical transformation of DNA. This cell contains the gyrA462 gene which provides resistance to the toxic effects from the ccdB gene. *Trans*DB3.1 Chemically Competent Cell can be used for transformation and propagation of plasmid containing the ccdB gene. It permits a transformation efficiency of over 10⁸ cfu/μg DNA (tested by pUC19 plasmid DNA).

Genotype

 $F \ gyrA462 \ end A1 \ \Delta(sr1-recA) \ mcrB \ mrr \ hsd S20(r_n, m_n) \ sup E44 \ ara-14 \ gal K2 \ lac Y1 \ pro A2 \ rps L20(Sm^R) \ xyl-5 \ \lambda-leu \ mtl 1 \ lac Y1 \ pro A2 \ rps L20(Sm^R) \ xyl-5 \ \lambda-leu \ mtl 1 \ lac Y1 \ pro A2 \ rps L20(Sm^R) \ xyl-5 \ \lambda-leu \ mtl 1 \ lac Y1 \ pro A2 \ rps L20(Sm^R) \ xyl-5 \ \lambda-leu \ mtl 1 \ lac Y1 \ pro A2 \ rps L20(Sm^R) \ xyl-5 \ \lambda-leu \ mtl 1 \ lac Y1 \ pro A2 \ rps L20(Sm^R) \ xyl-5 \ \lambda-leu \ mtl 1 \ lac Y1 \ pro A2 \ rps L20(Sm^R) \ xyl-5 \ \lambda-leu \ mtl 1 \ lac Y1 \ pro A2 \ rps L20(Sm^R) \ xyl-5 \ \lambda-leu \ mtl 1 \ lac Y1 \ pro A2 \ rps L20(Sm^R) \ xyl-5 \ \lambda-leu \ mtl 1 \ lac Y1 \ pro A2 \ rps L20(Sm^R) \ xyl-5 \ \lambda-leu \ mtl 1 \ lac Y1 \ pro A2 \ rps L20(Sm^R) \ xyl-5 \ had \ xy$

Features

- High transformation efficiency: >108 cfu/μg (pUC19 DNA).
- Transformation and propagation of plasmids containing the ccdB gene.
- · StrR.

Procedures

- Equilibrate a water bath to 42°C.
- Warm a vial of SOC medium or LB medium to room temperature. Warm selective plates at 37°C for 30 minutes.
- Thaw a vial of 100 μl of *Trans*DB3.1 Chemically Competent Cell on ice, aliquot 50 μl of the cells into a prechilled 1.5 ml tube, add target DNA (1 to 5 μl) into the tube. Do not mix by pipetting up and down. Incubate the cells on ice for 30 minutes.
- Heat-shock the cells for 45 seconds at 42°C without shaking. Immediately transfer the tube to ice. Incubate on ice for 2 minutes without shaking.
- Add 500 µl of prewarmed SOC medium or LB medium (without antibiotic) into the tube, mix well and shake at 37°C for 1 hour at 200 rpm for cell recovery and for the expression of antibiotic resistance.
- Spread 20 to 200 µl from each transformation vial on a prewarmed selective plate. The remaining can be stored at 2-8°C and plated the next day if needed.
- Invert the plate and incubate at 37°C overnight.
- Select colonies and analyze by restriction enzyme digestion, PCR, or sequencing.

Notes

- · Higher efficiency transformation can be achieved by transforming cells immediately following thawing.
- · Avoid repeated thawing.
- Gentle handling is required for the entire procedure.

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