

Trans10 Chemically Competent Cell

Cat. No. CD101

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

Description

Trans10 Chemically Competent Cell is specifically designed for chemical transformation of DNA. It permits a transformation efficiency of over 10^8 cfu/ μ g DNA (tested by pUC19 plasmid DNA). The competent cell is resistant to streptomycin sulfate (Str^R).

Genotype

F⁻ *mcrA* Δ (*mrr-hsdRMS-mcrBC*) ϕ 80 *lacZ* Δ M15 Δ *lacX74* *recA1* *ara* Δ 139 Δ (*ara-leu*)7697 *galU* *galK* *rpsL* (Str^R) *endA1* *nupG*

Features

- High transformation efficiency: $>10^8$ cfu/ μ g (pUC19 DNA).
- Str^R.
- Blue/white selection.
- Toxic gene cloning and stable replication of plasmid DNA.

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 μ l of Trans10 Chemically Competent Cell on ice, aliquot 50 μ l of the cells into a prechilled 1.5 ml tube, add target DNA (1 to 5 μ l) into the tube. Do not mix by pipetting up and down. Incubate the cells on ice for 30 minutes.
- Heat-shock the cells for 30 seconds at 42°C without shaking. Immediately transfer the tube to ice. Incubate on ice for 2 minutes without shaking.
- Add 500 μ l 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 μ l 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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