

Apr 13, 2020

Preparation of XL1-Blue competent cells using MgCl₂ and CaCl₂

DOI

dx.doi.org/10.17504/protocols.io.3trgnm6

Rashmi Karki¹, Monica Rieth¹

¹Southern Illinois University-Edwardsville

Labyrieth



Monica Rieth

Southern Illinois University-Edwardsville

OPEN  ACCESS



DOI: dx.doi.org/10.17504/protocols.io.3trgnm6

Protocol Citation: Rashmi Karki, Monica Rieth 2020. Preparation of XL1-Blue competent cells using MgCl₂ and CaCl₂. protocols.io <https://dx.doi.org/10.17504/protocols.io.3trgnm6>

License: This is an open access protocol distributed under the terms of the **Creative Commons Attribution License**, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited

Protocol status: Working

We use this protocol and it's working

Created: June 06, 2019

Last Modified: April 13, 2020

Protocol Integer ID: 24145

Keywords: XL1-Blue competent cells, Calcium-Chloride method



Abstract

Preparation of chemically competent XL1-Blue cells using MgCl_2 and CaCl_2

The purpose is to prepare batches of chemically competent bacteria for the purposes of subcloning and protein expression. Competent bacteria are able to readily and passively take up foreign DNA due their compromised cell walls as a result of exposure to divalent metal ions.

This protocol is based on current methods reported in:

Sambrook, Joseph et al. (2001). Molecular cloning : a laboratory manual. Cold Spring Harbor, N.Y. :Cold Spring Harbor Laboratory Press

Guidelines

This protocol must be carried out under sterile conditions. Wherever possible working benchtop and spaces should be wiped down with 70% EtOH solution. Sterilized, gloved hands are a must at all times.

Materials

MATERIALS

- CaCl₂
- MgCl₂
- DMSO
- Micropipettes and tips
- Eppendorf tubes (1.5 & 2.0 ml)
- Paper towels
- LB Broth
- XL1-Blue cells
- sterile culture tubes (17 × 100 mm)
- high-speed preparatory centrifuge

Safety warnings

- ⚠ Proper clothing is required as it needs to be done in cold room (4 °C)
Keep the DMSO at room temperature
Wear gloves at all times



Before start

Autoclave 100 ml LB- broth in a 1000 ml flask

Sterilize and pre-chill the 50-mL conical tubes

Prepare 30 ml of ice cold MgCl_2 - CaCl_2 solution (80mM MgCl_2 , 20 mM CaCl_2)

Prepare ice cold 0.1 CaCl_2

Pre-warm the LB plates



Day 1

- 1 Inoculate a 5 mL culture of liquid LB media from a glycerol stock (-80°C) of XL1-Blue competent cells
- 2 Incubate it for 16-20 hours at 37°C at 220-225 rpm

Day 2

- 3 Remove the culture tubes from incubator
- 4 Make four dilutions of the culture in liquid broth
 - 4.1 Dilution 1: 20 μL of the culture into 980 μL LB
 - 4.2 Dilution 2: 20 μL from dilution 1 into 980 μL LB
 - 4.3 Dilution 3: 20 μL from dilution 2 into 980 μL LB
 - 4.4 Dilution 4: 20 μL from dilution 3 into 980 μL LB
 - 4.5 Plate 200 μL of dilution 4 in LB plate and let it sit for some time until all the culture has been soaked
 - 4.6 Keep the plates inverted in incubator for 16-20 hours at 37°C .

Day 3

- 5 Remove the plates from the incubator.
- 6 Pick a single bacterial colony (2-3mm in diameter) from a plate (dilution 4) that has been incubated for 16-20 hours at 37°C.
- 7 Transfer into 100 ml sterilized LB broth or SOB medium in a 1-liter flask.
- 8 Incubate for 6 hr at 37°C with shaking at 220-250 rpm.
- 9 Measure OD₆₀₀ of culture every 15-20 minutes to ensure that the culture does not grow to a higher density. The transformation is efficient at OD₆₀₀~ 0.45.
- 10 Harvest the cell by arresting the growth when OD₆₀₀~0.45. Immediately put the flask in the ice.
- 11 Transfer the bacterial cells to pre sterilized and ice-cold 50 ml conical tube. Cool the cultures to 0°C by storing the tubes on the ice for 10 minutes.
- 12 Centrifuge at 2700g (4100 rpm in a Sorvall GSA rotor) or (RC-3B refrigerated centrifuge) for 10 minutes at 4 °C
- 13 Decant the medium from the cell pellets and keep the tubes in an inverted position over the paper towel for 1 minute to allow the traces of media to drain away
- 14 Resuspend the pellets by swirling or gentle vortexing in 30 ml of ice-cold MgCl₂-CaCl₂ solution (80mM MgCl₂, 20 mM CaCl₂) and mix until you get rid of all visible chunks
- 15 Centrifuge at 2700g (4100 rpm in a Sorvall GSA rotor) for 10 minutes at 4 °C
- 16 Decant the medium from the cell pellets and keep the tubes in the inverted position over the paper towel for 1 minute to allow the traces of media to drain away
- 17 Resuspend the pellets by swirling or gentle vortexing in 2 ml of ice-cold 0.1 CaCl₂ solutions for each 50 ml of the original culture and leave the tubes on ice for 15 minutes.



- 18 Add 140 μ l DMSO (cryo-protectant) per 4ml. Swirl gently to mix and let it sit for 15 minutes.
- 19 Add additional 140 μ l DMSO per 4 ml. Mix gently and return the tube to the icebox.
- 20 Transfer and store in aliquots of 100 μ l in sterile 1.5 mL Eppendorf tube.
- 21 Freeze at -80 °C.