

pMC1neo and pMC1neo Poly A Vectors

Instruction Manual

Catalog #213201

Revision C0

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pMC1neo and pMC1neo Poly A Vectors

MATERIALS PROVIDED

Material Provided	Quantity
pMC1neo	(25 μg)
pMC1neo poly A	(25 μg)
AG1 Strain: recA1, endA1, gyrA96, thi-1, hsdR17, (r _k -, m _k +), supE44, relA1, (uncharacterized mutation improves transformation efficiency)	0.5 ml

STORAGE CONDITIONS

Vectors: -20°C

AG1 Strain (Bacterial Glycerol Stock): -80°C

VECTOR SEQUENCES

The complete sequence and list of restriction sites for the pMC1neo and pMC1neo Poly A vectors are available at www.genomics.agilent.com.

PREPARATION OF HOST CELLS

The host strain has been sent as a glycerol stock. For the appropriate media and plates, please refer to the following table:

Bacterial strain	Plates for bacterial streak	Media for glycerol stock
AG-1	LB agar	LB agar

On arrival, prepare the following from the glycerol stock:

Note Do not allow the contents of the vial to thaw. The vials can be stored at -20 or -80° C, but most strains remain viable longer if stored at -80° C.

- 1. Revive the stored cells by scraping off splinters of solid ice with a sterile wire loop.
- 2. Streak the splinters onto an LB agar plate. Restreak the cells fresh each week.

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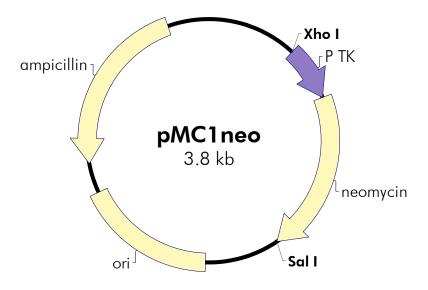
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Preparation of a -80°C Glycerol Stock

- 1. In a sterile 50-ml conical tube, inoculate 10 ml of the appropriate broth with one or two colonies from the plate. Grow the cells to late log phase.
- 2. Add 4.5 ml of a sterile glycerol–broth solution (5 ml of glycerol + 5 ml of broth) to the bacterial culture from step 1. Mix well.
- 3. Aliquot into sterile centrifuge tubes (1 ml/ tube).

This preparation may be stored at -20°C for 1-2 years or at -80°C for more than 2 years.

Vector Maps



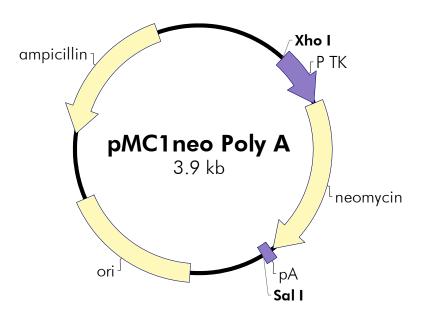


Figure 1 Map of the pMC1 neo vector (top) and pMC1 neo Poly A vector (bottom). The complete sequence and list of restriction sites for the vectors are available at www.genomics.agilent.com.

PREPARATION OF MEDIA AND REAGENTS

LB Broth (per Liter)

10 g of NaCl
10 g of tryptone
5 g of yeast extract
Add deionized H₂O to a final volume of
1 liter
Adjust to pH 7.0 with 5 N NaOH
Autoclaye

LB Agar (per Liter)

10 g of NaCl
10 g of tryptone
5 g of yeast extract
20 g of agar
Add deionized H₂O to a final volume of
1 liter
Adjust pH to 7.0 with 5 N NaOH
Autoclave
Pour into petri dishes (~25 ml/100-mm plate)

REFERENCE

1. Thomas, K. R. and Capecchi, M. R. (1987) Cell 51(3):503-12.

MSDS INFORMATION

Material Safety Data Sheets (MSDSs) are provided online at http://www.chem.agilent.com/en-US/search/library/Pages/MSDSSearch.aspx. MSDS documents are not included with product shipments.