



pRY121 plasmid in *E. coli*

37658™

Description

This construct contains the *S. cerevisiae* GAL1 promoter region (from about 1 kb upstream of the GAL1 transcription start site to amino acid 33) fused to the bacterial lacZ gene.

Organism: *Saccharomyces cerevisiae* Meyen ex E.C. Hansen

Clone type: Clone

Host: *Escherichia coli* LE392

Shipping information: *Escherichia coli* GC10 containing the plasmid in glycerol stock

Storage Conditions

Product format: Frozen

Storage conditions: -80°C or colder

Intended Use

This product is intended for laboratory research use only. It is not intended for any animal or human therapeutic use, any human or animal consumption, or any diagnostic use.

BSL 1

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or national agencies.

Certificate of Analysis

For batch-specific test results, refer to the applicable certificate of analysis that can be found at www.atcc.org.

Insert Information

Insert size (kb): 0.6999999999999996

Insert information:

Gene: Galactose promoter region

Source: *Saccharomyces cerevisiae*

• **Insert end:** Sall/BamHI

Vector Information

Construct size (kb): 11.50

Vector name: pLG670Z

Type of vector: plasmid

Vector information: Excise insert: Sall/BamHI

Markers: lacZ+; ampR; URA3

Growth Conditions

Medium:

ATCC Medium 1227: LB Medium (ATCC medium 1065) with 50 mcg/ml ampicillin

Temperature: 37°C

Handling Procedures

Thaw contents of the vial in a 37°C water bath with gentle agitation. Transfer a loopful to a test tube containing 5 mL LB+50 µg/mL of ampicillin broth. A loopful of culture can also be streaked on an LB + amp agar plate. Incubate cultures at 37°C. Isolate DNA using standard plasmid preparation procedures.

Notes

Restriction digests of the clone gave the following sizes (in kb): EcoRI – 3.2 (doublet), 2.8, 2.2 ; HindIII – 6.2, 2.15, 1.45, 1.05 ; Sall – Sall – 11.5.

–ATCC Staff

Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner: pRY121 plasmid in *E. coli* (ATCC 37658)

References

References and other information relating to this material are available at www.atcc.org.

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