



p423 GALL

87335™

Description

Clone type: Vector

Host: *Escherichia coli* HB101 (ATCC 33694)

Storage Conditions

Product format: Frozen

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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Certificate of Analysis

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

Vector Information

Construct size (kb): 6.459000110626221

Intact vector size: 6.459

Vector name: p423 GALL (plasmid)

Type of vector: plasmid

Construction: pRS423, GALL promoter

Host range: *Saccharomyces cerevisiae*; *Candida robusta*; *Escherichia coli*

Cloning sites: SpeI; BamHI; SmaI; EcoRI; EcoRV; ClaI; Sall; XhoI

Markers: HIS3; ampR

MCS: XhoI...SpeI, -, 2337-2394

Polylinker sites: XbaI; SpeI; BamHI; SmaI; PstI; EcoRI; EcoRV; HindIII; ClaI; Sall; XhoI

Promoters: GALL, <-, 2401-2831

Replicon: 2 micron

Terminator: CYC1, ->, 2077-2337

Growth Conditions

Medium:

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

Temperature: 37°C

Notes

Restriction digests of the clone give the following sizes (kb): SacI/XbaI--3.2, 2.8, 0.45; EcoRI--6.5; KpnI--5.5, 1.0.

- ATCC staff

High copy number shuttle expression vector.

- Nucleic Acids Res. 22: 5767-5768, 1994

One of 32 yeast expression vectors (ATCC 87318-87349) differing in promoter, selection marker and replicon.

- Nucleic Acids Res. 22: 5767-5768, 1994

The galactokinase (GALL) promoter is a deletion variant of GAL1, which lacks one of the 3 UAS elements required for full induction of the promoter by galactose.

- Nucleic Acids Res. 22: 5767-5768, 1994

The promoter is tightly repressed by glucose and is induced at a moderate level by galactose.

- Nucleic Acids Res. 22: 5767-5768, 1994

Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner: p423 GALL (ATCC 87335)

References

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

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Revision

This information on this document was last updated on 2025-09-12

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