

pC-ACT.2

Description

Clone type: Vector

Host: Escherichia coli DH5

Deposited As: Saccharomyces cerevisiae Hansen, teleomorph

Storage Conditions

Product format: Freeze-dried

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.

BSL₁

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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): 7.40399980545044

Intact vector size: 7.404

Vector name: pC-ACT.2 (plasmid)

Type of vector: plasmid Construction: pRS314 Vector information:

signal peptide coding region: hemagglutinin epitope, 6797-6838

signal peptide coding region: hemagglutinin epitope

Cloning sites: Ndel; Ncol; Avrll; Bglll; Bcll; BamHl; Smal; Sall; Pstl

Coding sequence: bla, <-, 934-1794; bla

Markers: LEU2; crbR

MCS: Ndel..Pstl, ->, 6845-6899

Polylinker sites: Ndel; Ncol; Avrll; Bglll; Bcll; BamHl; Smal; Sall; Pstl

Promoters: PADH, 5666-6375

Replicon: ColE1, 1-362; ARS4, 2066-2442 **Ribosome-binding site:** GAL4, ->, 6383-6791

Terminator: ADC1 transcriptional terminator (TADH), 6917-7107

Transcription terminator: ADC1 transcriptional terminator (TADH), 6917-7107

Growth Conditions

Medium:

ATCC Medium 1637: LB medium (ATCC medium 1065) with 100 mcg/ml carbenicillin

Temperature: 37°C

Notes

Restriction digests of the clone give the following sizes (kb): BamHI--7.4;

pC-ACT.2

PstI--7.4; HindIII--7.4; NdeI--7.4. Useful in two hybrid screening in yeast. A CEN based plasmid with Gal4 activation domain. It has lower copy number than 2 micron based plasmids. Contains unique rstriction sites (BgIII, BcII and BamHI) compatible with 5' GATC overhangs in all three reading frames.

- ATCC staff

Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner: pC-ACT.2 (ATCC 87777)

References

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

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