



Product Sheet

pBAD33 (ATCC® 87402™)

Please read this FIRST



Intended Use

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

Citation of Strain

If use of this culture results in a scientific publication, it should be cited in that manuscript in the following manner: pBAD33 (ATCC® 87402™)

American Type Culture Collection
PO Box 1549
Manassas, VA 20108 USA
www.atcc.org

800.638.6597 or 703.365.2700
Fax: 703.365.2750
Email: Tech@atcc.org

Or contact your local distributor

Description

Designation: pBAD33

Distribution Host: *Escherichia coli* DH5alpha

Propagation

Growth Conditions

Temperature: 37°C

Medium

ATCC® Medium 1675: LB medium (ATCC medium 1065) with 10 mcg/ml Chloramphenicol

Vector Information

Intact vector size: 5.356

Type of vector: phagemid

Cloning sites: SacI KpnI SmaI XbaI SalI PstI SphI HindIII

Polylinker sites: NheI EcoRI SacI KpnI SmaI BamHI XbaI SalI AccI PstI SphI HindIII

Construction: pBAD28

Host range: *Escherichia coli*

Features (with orientation and position when available):

regulator: araC, <-, 3452-4330

operator: O2, 4359-4376

promoter: araC, <-, 4481-4509

operator: O1, 4517-4538

other: CAP site, 4560-4573

operator: I2 + I1, 4569-4607

promoter for expression: arabinose BAD, ->, 4606-4633

MCS: NheI...HindIII, ->, 4656-4718

transcription terminator: rrnB T1 + T2, ->, 4719-5144

replicon: M13, 335-793

marker(s): cmLR, <-, 1348-2007

replicon: p15A, 2369-3213

Marker(s): cmIR,araC

References

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

Notes

Restriction digests of the clone give the following sizes (kb): Aval--3.0, 1.3, 1.2; BamHI--5.4; EcoRI--2.95, 2.6.

- ATCC staff

One of several tightly controlled expression vectors (ATCC 87393-87402) regulated by the arabinose operon. The vectors differ in replicon, antibiotic resistance marker, multiple cloning site and mechanism of initiation of translation.

- J. Bacteriol. 177: 4121-4130, 1995

Cultures should be grown in minimal media for more reproducible induction of expression. Expression is induced in glycerol-containing media by addition of arabinose. Expression is repressed by addition of glucose or other catabolites.

- J. Bacteriol. 177: 4121-4130, 1995

Plasmid copy number is low due to the p15A replicon. This vector can be used when reduced gene expression is desirable.

- J. Bacteriol. 177: 4121-4130, 1995

Plasmid is compatible with pBR-derived plasmids and may be used for coexpression of cloned inserts.

- J. Bacteriol. 177: 4121-4130, 1995



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Plasmid contains bla (ampR) sequences following the rrnB terminator which could promoter recombination if this plasmid is used in combination with other compatible ampR plasmids.

- J. Bacteriol. 177: 4121-4130, 1995

Recombination can be avoided by the use of recA host strains, or it can be used to advantage to intentionally exchange markers among plasmids.

- J. Bacteriol. 177: 4121-4130, 1995

The following primers can be used for sequencing of cloned inserts: 5' primer (27 - 8 bp upstream of the NheI site) 5'-CTGTTTCTCCATACCCGTT-3'; and one of two 3' primers: 3' primer 1 (2 - 19 bp downstream of the HindIII site) 5'-CTCATCCGCCAAACAG-3';

- J. Bacteriol. 177: 4121-4130, 1995

3' primer 2 (17 - 33 bp downstream of the HindIII site) 5'-GGCTGAAAATCTTCTCT-3'.

- J. Bacteriol. 177: 4121-4130, 1995



Biosafety Level: 1

Appropriate safety procedures should always be used with this material. Laboratory safety is discussed in the current publication of the *Biosafety in Microbiological and Biomedical Laboratories* from the U.S. Department of Health and Human Services Centers for Disease Control and Prevention and National Institutes for Health.

ATCC Warranty

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Disclaimers

This product is intended for laboratory research purposes only. It is not intended for use in humans.

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Please see the enclosed Material Transfer Agreement (MTA) for further details regarding the use of this product. The MTA is also available on our Web site at www.atcc.org

Additional information on this culture is available on the ATCC web site at www.atcc.org.

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