

# YCplac22

87585<sup>™</sup>

## 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<sub>1</sub>

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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.

#### Insert Information

Target gene: beta-galactosidase

#### **Vector Information**

Construct size (kb): 4.854000091552734

Intact vector size: 4.854

Vector name: YCplac22 (plasmid)

**Type of vector:** plasmid **Construction:** pUC19

Host range: Saccharomyces cerevisiae; Candida robusta; Escherichia coli

Cloning sites: EcoRI; SacI; KpnI; SmaI; BamHI; XbaI; HincII; AccI; SalI; PstI; SphI; HindIII

Insert detection: lacZ', ->, 216-500

Markers: ampR; TRP1

MCS: HindIII...EcoRI, ->, 234-285

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

Promoters: lac

Replicon: ARS1, 1481-2225; pMB1, 4666-4666

#### **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): Aval--2.9, 1.9;



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HindIII--4.8; XbaI--4.8.

- ATCC staff

One of 9 shuttle vectors (ATCC 87585 - 87593) allowing lacZ detection of cloned inserts and containing the pUC19 MCS with all 10 cloning sites unique. Vectors differ in mode of replication (YE, YC or YI-type) and yeast marker (LEU2, URA3 or TRP1).

- Gene 74: 527-534, 1988

The HindIII, PstI and the Xbal sites from the TRP1 gene of the S. cerevisiae were removed by using oligo-directed mutagenesis.

- Gene 74: 527-534, 1988

#### Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner: YCplac22 (ATCC 87585)

#### 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 2022-10-22

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