

87447TM

Description

This plasmid contains a trimethoprim resistance cassette which is useful as a selectable marker, for cassette mutagenesis of cloned genes, and for the construction of new cloning vectors. The cassette insert was amplified using primers that allowed incorporation of two EcoRI sites, resulting in 3 bp mismatches with respect to the portion of the GenBank record J01774 described below. This vector was constructed by inserting a trimethoprim resistance cassette into the pGEM-3/4 vector. The insert and vector components are described below under ?insert component? and ?vector component?, respectively.

Organism: R388 Clone type: Vector

Shipping information: Escherichia coli 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.

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.

Insert Information

Insert size (kb): 0.61799999999999999

Type of DNA: cDNA

Target gene: dihydrofolate reductase, trimethoprim resistance

Gene product: dihydrofolate reductase, trimethoprim resistance [dhfrIIb]

Vector Information

Construct size (kb): 3.527

Vector name: p34E-Tp (plasmid)

Construction: pGEM-3, pGEM-4, dhfrIIb (tmpR)

Markers: tmpR; ampR

MCS: EcoRI...HindIII; HindIII...EcoRI Promoters: Expression: lac; *In vitro*

Growth Conditions



Medium:

ATCC Medium 2031: LB + 1.5 mg/ml trimethoprim

Temperature: 37°C

Notes

Restriction digests of the clone gave the following sizes (in kb): Aval 2.9, 0.4, 0.2; Bgll 3.5; EcoRl 2.9, 0.6. ATCC Staff

Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner: p34E-Tp plasmid in E. coli (ATCC 87447)

References

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

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Contact Information

ATCC

10801 University Boulevard

Manassas, VA 20110-2209

USA

US telephone: 800-638-6597

Worldwide telephone: +1-703-365-2700

Email: tech@atcc.org or contact your local distributor

