

Description

This clone encodes the L22F mutant of DHFR that can serve as a dominant selection marker because it retains catalytic activity and also confers methotrexate (MTX) resistance at low MTX concentrations (125 nM). Transfected cells amplify DHFR sequences. Constructed by cloning a HindIII/Smal fragment containing the gene into pSV2-neo from which neoR had been removed. This plasmid was then cut with Smal and BglII to remove 18 G nucleotides at the 3? end. The BglII site was filled in, and ligated to the Smal site, thereby destroying the original BglII and Smal sites. Removing the HindIII/Smal fragment from the pSV2-neo removes all but the 3? 171 bp of neoR.

Organism: Cricetulus griseus, hamster, Chinese

Clone type: Clone

Host: Escherichia coli HB101 (ATCC 33694)

Storage Conditions

Product format: Freeze-dried **Storage conditions:** 2°C to 8°C

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





ATCC determines the biosafety level of a material based on our risk assessment as guided by the current edition of *Biosafety in Microbiological and Biomedical Laboratories* (*BMBL*), U.S. Department of Health and Human Services. It is your responsibility to understand the hazards associated with the material per your organization's policies and procedures as well as any other applicable regulations as enforced by your local or national agencies.

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): 0.709999999999996

Type of DNA: cDNA

Insert source: lung cell line **Insert tissue:** lung cell line

Insert information:

DESCRIPTION OF INSERT COMPONENT:

Insert 5' end: HindIII Insert 3' end: BgIII Cross references:

Genome: hamster, Chinese

Target gene: dihydrofolate reductase **Gene name:** dihydrofolate reductase

Gene product: dihydrofolate reductase [DHFR]

Gene symbol: DHFR

Contains complete coding sequence: Unknown

Vector Information



pSVA3

Construct size (kb): 5.099999904632568

Intact vector size: 5.729 Vector name: pSV2-neo Type of vector: plasmid

Construction: SV40, Tn5, pBR322
Host range: vertebrate cells
Vector end: HindIII; Smal
Vector information:

Cross references: DNA Seq. Acc.: U02434

Cloning sites: EcoRI; BamHI; PvuII; PstI; HindIII

Markers: ampR

Promoters: SV40 early **Replicon:** SV40; pMB1 **Restriction sites:** HindIII

Growth Conditions

Medium:

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

Temperature: 37°C

Handling Procedures

Transfer a loopful to a test tube containing 5 mL LB+50mg/mL of ampicillin broth. A loopful of culture can also be streaked on an LB + amp agar plate. Incubate cultures at 37°C. Isolate DNA using standard plasmid preparation procedures.

Notes

Restriction digests of the clone gave the following sizes (in kb): HindIII/Smal -- 5.2;

BamHI -- 3.5, 1.7; EcoRI -- 2.7, 2.3; Pstl -- 1.9, 1.4, 0.95, 0.9; Pvull -- 4.0, 1.1.

---ATCC staff

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pSVA3

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- ATCC staff

Encodes the L22F mutant of DHFR that can serve as a dominant selection marker because it retains catalytic activity and also confers methotrexate (MTX) resistance at low MTX concentrations (125 nM).

- Gene 112: 179-188, 1992

Transfected cells amplify DHFR sequences.

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Constructed by cloning a HindIII/Smal fragment containing the gene into pSV2-neo from which neoR had been removed. This plasmid was then cut with Smal and BgIII to remove 18 G nucleotides at the 3' end.

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The Bglll site was filled in, and ligated to the Smal site, thereby destroying the original Bglll and Smal sites.

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Removing the HindIII/Smal fragment from pSV2-neo removes all but the 3' 171 bp of neoR.

- Gene 112: 179-188, 1992

Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner: pSVA3 (ATCC 77273)

References





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

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