Product Sheet

PMMTV-STU MYC Plasmid in Escherichia coli

39748[™]

Description

Organism: *Mus musculus*, mouse **Clone type:** Clone **Host:** *Escherichia coli* K-12

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Patent number:

4,736,866

Technical information: ATCC Product Experience does not have technical information on patent deposits that are not produced or characterized by ATCC. Additional information can be found in the corresponding patent available from the patent holder or with the U.S. and/or international patent office.

Storage Conditions

Product format: Freeze-dried

Intended Use

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BSL 1

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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

Type of DNA: genomic Insert information: Genomic copy number: unique Insert 5' end: Stul Insert 3' end: EcoRI Genome: mouse Chromosome: 15 15 Gene name: avian myelocytomatosis viral (v-myc) oncogene homolog Gene product: avian myelocytomatosis viral (v-myc) oncogene homolog(avian myelocytomatosis viral (v-myc) oncogene c-myc) [Myc] Gene symbol: Myc Contains complete coding sequence: Unknown

Vector Information



Product Sheet

Construct size (kb): 15.0 Vector name: pA9 Type of vector: plasmid Host range: Escherichia coli Vector end: Smal; EcoRI Cloning sites: Smal; EcoRI Enhancer: MMTV LTR Markers: ampR

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): PstI--4.4, 4.2, 3.3, 2.7; PvuII--6.6, 4.2, 1.8, 1.4, 0.56; EcoRI--19.0; HindIII--8.0, 4.4, 1.4, 1.2, 0.9.

Constructed by ligating a Stul/EcoRI fragment from a genomic subclone of the gene to a Smal/EcoRI fragment of pA9 (containing the glucocorticoid control region, MMTV promoter, and cap site). The insert begins approximately 3.0 kb 5' of exon 1. It therefore contains the 2 promoters naturally preceding the unactivated gene. The 3' end of the myc sequence in this construct is the HindIII site approximately 1.0 kb 3' of the poly(A) addition site. The EcoRI site used in subcloning was derived from the vector (pBR322). A 5.2 kb BamHI(5')/ClaI(3') fragment containing exons 2 and 3 has been used as a probe. The BamHI site is in intron 1 and the ClaI site is from the flanking pBR322-derived sequences.

- U.S. Pat. 4,736,866 dated Apr. 12, 1988

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.patent

Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner: PMMTV-STU MYC Plasmid in *Escherichia coli* (ATCC 39748)

References

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

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