

39757<sup>™</sup>

## Description

Clone type: Clone Host: Escherichia coli

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#### Patent number:

4.666.848

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## Storage Conditions

Product format: Freeze-dried

### 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.



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### 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

Type of DNA: cDNA

Insert information: Genomic copy number: unique

**Genome:** human **Chromosome:** 4

4 q26-q27

Gene name: interleukin 2

Gene product: interleukin 2 [IL2]

Gene symbol: IL2

Contains complete coding sequence: Unknown

## **Vector Information**

Construct size (kb): 0.0 Intact vector size: 4.363 Vector name: pBR322 Type of vector: plasmid



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Construction: pBR313
Host range: Escherichia coli
Vector end: EcoRI; PstI
Vector information:

Cross references: DNA Seq. Acc.: J01749

Cloning sites: EcoRI; ClaI; HindIII; EcoRV; BamHI; SphI; SalI; XmaIII; NruI; BspMI; BsmI; StyI; AvaI; BalI; BspMII; PvuII; Tth111I; NdeI; AfIIII; PpaI; PstI; PvuI; ScaI; SspI; AatII

Markers: ampR; tetR Replicon: pMB1

#### **Growth Conditions**

Temperature: 37°C

#### Notes

The cry sequence enhances expression of upstream coding regions.

- U.S. Pat. 4,865,974 dated Sept. 12, 1989

Constructed by inserting a 0.4 kb BamHI/EcoRI fragment containing the 3' sequences of the Bacillus thuringiensis crystal protein gene (cry) into the Stul site of pLW1 (ATCC 39405).

- U.S. Pat. 4,666,848 dated May 19, 1987

The order of the major features in this plasmid is: pBR322 - trp promoter - ribosome binding sites - IL2 - BamHI - cry sequence - EcoRI.

- U.S. Pat. 4,865,974 dated Sept. 12, 1989

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#### Material Citation



If use of this material results in a scientific publication, please cite the material in the following manner: PHCW701 Plasmid in *Escherichia coli* (ATCC 39757)

#### References

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

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## Revision



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