

39841TM

Description

Clone type: Vector

Host: Escherichia coli HB101 (ATCC 33694)

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Patent number:

4,824,786

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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BSL₁



pLA2920 39841

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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.

Vector Information

Construct size (kb): 20.0

Vector name: pLA2920 (cosmid) **Construction:** pRK290, pMF517 cos

Enhancer: none
Markers: tetR
Promoters: none
Replicon: oriV

Growth Conditions

Medium:

ATCC Medium 1273: LB medium (ATCC medium 1065) with 20 mcg/ml tetracycline

Temperature: 30°C

Notes



- ATCC staff

Restriction digests of the clone give the following sizes (kb): PstI--20.0; HindIII--20.0; BgIII--20.0; EcoRI--17.0; 2.6. IMPORTANT: To prevent amplification of a rearranged and/or deleted cosmid, we recommend streaking on LB + amp plates at 30C and picking small colonies for liquid culture.

This is a cosmid with a broad host range that can be used to characterize the genome of C1-utilizing microorganisms. This vector transfers efficiently to a broad host range including Escherichia coli HB101, Methylobacterium organophilum DSM 761, Pseudomonas AM1, Pseudomonas putida, and Pseudomonas aeruginosa.

was constructed from pLA2917 (ATCC 39840) by deleting the kanamycin resistance segment from Tn5 to avoid internal rearrangement when the transposon moves and to permit selection of conjugants that have a Tn5 insertion.

- U.S. Pat. 4,824,786 dated Apr. 25, 1989

.patent

This

Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner: pLA2920 (ATCC 39841)

References

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

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