



# pGIKS Dap5 87827™

## Description

This item is one of a set of 11 cloned bacterial and phage genes (1) (set = ATCC No. 87840). *Escherichia coli* DH5a containing pGIKS-Dap5 is provided as a frozen glycerol aliquot. Nucleotides 1316 to 1981 of the gene *dapB* for the putative product dihydropicolinate reductase were directionally cloned from *Bacillus subtilis* genomic DNA into the *Xho* I - *Not* I (5'-3') regions of the pBluescript II KS+ phagemid. The phagemid allows for either the production of an antisense transcript from the T7 promoter or a sense transcript from the T3 promoter. A 720 nucleotide transcript is produced from the T7 promoter when the construct is linearized with *Xho* I.

**Organism:** *Escherichia coli* (Migula) Castellani and Chalmers

**Clone type:** Clone

**Deposited As:** human

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## Storage Conditions

**Product format:** Frozen

**Storage conditions:** -80°C or colder

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## 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 1

ATCC determines the biosafety level of a material based on our risk assessment as

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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](http://www.atcc.org).

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## Insert Information

**Insert size (kb):** 0.7199999999999997

**Type of DNA:** cDNA

**Gene product:** [DapB]

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## Vector Information

**Construct size (kb):** 3.599999904632568

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## Growth Conditions

**Medium:**

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

**Temperature:** 37°C

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## Handling Procedures

The phagemid construct within the *E. coli* DH5a host can be grown in LB + amp (50 mg/mL) at 37°C and then isolated using standard plasmid preparation procedures (2).

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## Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner: pGIKS Dap5 (ATCC 87827)

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## References

References and other information relating to this material are available at [www.atcc.org](http://www.atcc.org).

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## Revision

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