

# 87484<sup>TM</sup>

# **Description**

Expression control clone for microarrays.

Organism: Bacillus subtilis subsp. subtilis (Ehrenberg) Cohn

Clone type: Clone

Host: Escherichia coli DH5

Shipping information: Escherichia coli containing the phagemid in glycerol stock

# **Storage Conditions**

**Product format:** Frozen

Storage conditions: -80°C or colder

#### 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.

### 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

Insert size (kb): 1.98
Insert information:

Gene: thrB

Source: Bacillus subtilis

Genbank accession: X04603 (Nucleotides 1-1982 of the insert correspond to

nucleotides 248-2229 of X04603)

Insert end (5'): XhoI linker

Insert end (3'): BamHI/poly(A)/NotI linker

# **Vector Information**

Construct size (kb): 4.9 Intact vector size: 2.961

Vector name: pBluescript II KS+

**Type of vector:** phagemid **Vector end:** Xhol; Notl

Vector information: Excise insert: Xhol+Notl

Promoter giving the sense strand:T3
Promoter giving the antisense strand: T7

Markers: ampR Promoters: T3; T7

### **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+50  $\mu$ g/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): Notl/XhoI 3.0, 2.0; XhoI 5.0.

-ATCC Staff

#### Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner: pGIBS-THR (ATCC 87484)

### References

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

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