



pCMV-h240H

99665™

Description

Full length clone of human cholesterol 24-hydroxylase in the pCMV6 vector. The insert for this clone is human cholesterol 24 OH (Genbank number AF094480), nt (-42) to 111 is genomic DNA and from 111 to end is cDNA.

Note: Nt 356 of insert is C, differing from AF094480 which is G. This is probably an RT error and not a polymorphism. This causes an R to T substitution, yet the expressed protein is still active.

-----Personal communication

Organism: *Homo sapiens*, human

Clone type: Clone

Deposited As: human

Shipping information: *Escherichia coli* containing the plasmid

Storage Conditions

Product format: Freeze-dried

Storage conditions: 2°C to 8°C

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 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

Insert size (kb): 2.2000000000000002

Type of DNA: cDNA

Target gene: cholesterol 24 hydroxylase

Gene product: cholesterol 24 hydroxylase [h24OH]

Vector Information

Construct size (kb): 6.900000095367432

Growth Conditions

Medium:

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

Temperature: 37°C

Handling Procedures

Aseptically add 0.3 to 0.4 mL of liquid medium to the freeze-dried pellet and mix well. Transfer 100 μ L to a test tube containing 5 mL LB+50mg/mL of ampicillin. A loopful of culture can also be streaked on an LB + amp agar plate. Incubate cultures at 37⁰ C. Isolate DNA using standard plasmid preparation procedures.

Notes

Restriction digests of the clone gave the following bands (in kb) : EcoRI 4.6, 2.2 ; PstI 4.6, 1.1, 0.7. —ATCC staff

Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner: pCMV-h240H (ATCC 99665)

References

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

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