

MBA-392[™]

Description

This product is an ATCC manufactured and accessioned progeny of 40379 cited in US Patent Number 4,908,306.

Organism: human papilloma virus type 56

Clone type: Clone

Deposited As: ATCC accessioned progeny of HPV 56 clone 2B purified plasmid DNA

cited in US Patent Number 4,908,306 as 40379. **Shipping information:** Purified plasmid DNA

Storage Conditions

Product format: 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.

BSL₁

ATCC determines the biosafety level of a material based on our risk assessment as guided by the current edition of *Biosafety in Microbiological and Biomedical Laboratories* (*BMBL*), U.S. Department of Health and Human Services. It is your responsibility to understand the hazards associated with the material per your organization's policies



MBA-392

and procedures as well as any other applicable regulations as enforced by your local or national agencies.

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): 5.9000000000000004
Insert source: human vulvar condyloma
Gene name: human papilloma virus type 56

Insert end: BamHI

Vector Information

Construct size (kb): 8.7 Intact vector size: 2.8 Vector name: pT713 Vector end: BamHI Markers: ampR

Promoters: T7 (phi10)

Handling Procedures

Rehydrate plasmid with water or TE buffer. Plasmid can be transformed into a suitable *Escherichia coli* host using standard protocols and then grown on LB + 50 μ g/mL of ampicillin agar. Incubate cultures at 37°C. Isolate DNA using standard plasmid preparation procedures.

MBA-392

Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner: HPV 56 clone 2B purified plasmid DNA (ATCC MBA-392)

References

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

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MBA-392

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Revision

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MBA-392

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