



Kosmotoga olearia

BAA-1733™

Description

Strain designation: 19.5.1

Deposited As: *Kosmotoga olearia*

Type strain: No

Storage Conditions

Product format: Freeze-dried

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

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

ATCC highly recommends that appropriate personal protective equipment is always used when handling vials. For cultures that require storage in liquid nitrogen, it is important to note that some vials may leak when submersed in liquid nitrogen and

will slowly fill with liquid nitrogen. Upon thawing, the conversion of the liquid nitrogen back to its gas phase may result in the vial exploding or blowing off its cap with dangerous force creating flying debris. Unless necessary, ATCC recommends that these cultures be stored in the vapor phase of liquid nitrogen rather than submersed in liquid nitrogen.

Certificate of Analysis

For batch-specific test results, refer to the applicable certificate of analysis that can be found at www.atcc.org.

Growth Conditions

Temperature: 65°C

Atmosphere: Anaerobic

Handling Procedures

1. Sterilize the top of the Balch tube by spraying it with 70% ethanol and then flaming the top.
2. Exchange the gas in the test tube for 80% N₂ 20% CO₂, do not pressurize over 5 10 psi. If the tubes are over pressurized inoculating the tubes will prove difficult.
3. Prepare tubes for inoculation: If the medium has been sitting around for more than a two weeks add 0.1 ml of reducing agent (3.0% Cysteine, stock solution) per 10 ml of medium. Let the medium sit at room temperature for at least 1 hour before inoculating.
4. Thaw the frozen vial under a gentle stream of anaerobic gas. Using an anaerobic (see D) 1.0 ml syringe tipped with 22-gauge needle, withdraw the cell suspension

from the vial and transfer to the 1^o tube of ATCC® #2711 medium. Transfer 0.5 ml of the inoculated culture to one or more (2^o) second tube of ATCC® medium #2711.

Plate 0.1 ml of the inoculated culture onto a non-selective medium and incubate the plate aerobically at 37°C. Incubate culture tubes at 65°C.

5. Growth should be detected in the broth within 24–48 hours. No growth should be detected on the aerobic plate.

ANAEROBIC CONDITIONS:

A. Balch tubes (available from Bellco Glass, Vineland, NJ; are specially designed for anaerobic work and use an aluminum crimp cap to hold a rubber stopper in place. Needles can easily be inserted through the stopper, and the tubes can be pressurized to 2 atm. Alternatively, serum vials may be used, or screw cap tubes with butyl rubber stoppers, in the latter case the stopper may be removed and the tube placed under a cannula system that dispenses sterile, oxygen free gas for addition of reducing agents or inoculation.

B. To obtain a fully reduced medium, it is necessary that the medium be anoxic and that a reducing agent be added. Common reducing agents are sodium sulfide, cysteine, dithiothreitol, titanium citrate and Co-enzyme M (see D).

C. We suggest adding the reducing agent to the medium at least one hour before the medium is to be inoculated.

D. Syringes can be made anaerobic by one of two methods.

Notes

Always use freshly prepared anaerobic medium. If there is any question about the anaerobic condition of the medium, then it can be reduced with the addition of 3.0% Cysteine (0.1 ml per each 10 ml of medium).

Material Citation

If use of this material results in a scientific publication, please cite the material in the

following manner: *Kosmotoga olearia* (ATCC BAA-1733)

References

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

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