Mycoplasma hyopneumoniae Mare and Switzer

27715™

Description

Strain designation: J (FCX3-line 1)
Deposited As: *Mycoplasma suipneumoniae* Goodwin et al.
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 2

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

Medium:
ATCC Medium 1699: Revised Mycoplasma medium
Temperature: 37°C
Atmosphere: 95% Air, 5% CO_2

Handling Procedures

1. Follow instructions as suggested for the culturing of

Mollicutes:

PROCEDURES FOR PROPAGATING *MOLLICUTES*:

a) Open the vial according to the enclosed instructions.

b) Using a Pasteur or 1.0 ml pipette, withdraw approximately 0.5 to 1.0 ml from a tube containing 5.0 ml. Rehydrate the pellet.

c) Aseptically transfer this aliquot back into the tube. Mix well.

d) Make serial dilutions by transferring 0.5 ml from the

e) Use an uninoculated tube of broth to serve as a control.

f) Plates may be inoculated to check colonial morphology. You can also spot each dilution on the surface of plate (4 or more/plate) to determine the number of colony-forming units. However, not all strains do well on solid medium.

g) Incubate all tubes and plates under the recommended conditions and appropriate temperature. The time necessary for growth will vary from strain to strain. Growth on plates generally requires additional incubation.

h) Depending on the medium used, growth is indicated by increased turbidity, a color change, or both.

i) Tubes are incubated aerobically, plates are incubated under 5% CO_2 or in a *candle jar*. The incubation temperature is 37°C.

3. After broth growth is established, freshly inoculated plates will take three or more days to produce colonies. They vary in size from tiny to small, rough with irregular margin. They do not exhibit the usual "fried egg" appearance.

4. For long term storage of *Mycoplasma hyopneumoniae,* freeze-drying or freezing is recommended. Liquid nitrogen storage is the best method. Optimally grown cells are centrifuged at 9000 rpm for 30 minutes, the supernatant poured off, and the packed cells resuspended in a smaller amount of #1699 broth. To this, add an equal amount of sterile 20% glycerol as a cryoprotectant. This suspension is aliquoted into small plastic vials and stored at -70°C or below.

Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner: *Mycoplasma hyopneumoniae* Mare and Switzer (ATCC 27715)

References

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

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Revision

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Contact Information

ATCC
10801 University Boulevard

Manassas, VA 20110-2209
USA
US telephone: 800-638-6597
Worldwide telephone: +1-703-365-2700
Email: tech@atcc.org or contact your local distributor