



# *Entamoeba histolytica* Schaudinn

30459™

## Description

*Entamoeba histolytica* strain HM-1:IMSS [ABRM] is a parasitic protozoan that was isolated from the colonic biopsy of a rectal ulcer from an adult male with amebic dysentery. This strain can be used in enteric and infectious disease research.

**Strain designation:** HM-1:IMSS [ABRM]

**Deposited As:** *Entamoeba histolytica* Schaudinn

**Type strain:** No

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## Storage Conditions

**Product format:** Frozen

**Storage conditions:** -80°C or colder for 1 week, vapor phase of liquid nitrogen for long-term storage

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## 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.

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## 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

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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.

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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](http://www.atcc.org).

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### **Growth Conditions**

**Medium:**

ATCC Medium 2463: LYI-S-2 medium, modified

**Instructions for complete medium: Media:** ATCC Medium 2463.

**Alternate Media:** ATCC Medium PRA-2154 modified with an additional 5% heat-inactivated bovine serum may also be used; quality controlled freeze-dried lots of this medium are commercially available from ATCC.

**Temperature:** 35°C

**Atmosphere:** Microaerophilic

**Culture system:** Axenic

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## Handling Procedures

### Storage and Culture Initiation

Frozen ampules packed in dry ice should either be thawed immediately or stored in liquid nitrogen. If liquid nitrogen storage facilities are not available, frozen ampules may be stored at or below  $-70^{\circ}\text{C}$  for approximately one week. **Do not under any circumstance store frozen ampules at refrigerator freezer temperatures (generally  $-20^{\circ}\text{C}$ ).** Storage of frozen material at this temperature will result in the death of the culture.

1. To thaw a frozen ampule, place in a  $35^{\circ}\text{C}$  water bath, until thawed (2-3 min). Immerse the ampule just sufficiently to cover the frozen material. Do not agitate the ampule.
2. Immediately after thawing, aseptically transfer contents to a glass screw-capped tube containing 13 mL ATCC Medium 2463. Screw cap on tightly and incubate on a  $15^{\circ}$  horizontal slant at  $35^{\circ}\text{C}$ .

### Culture maintenance:

1. Ice culture at or near peak density for 10 min.
2. Gently invert culture 20 times.
3. Aseptically transfer a 0.1 and 0.25 mL aliquot to freshly prepared (no older than 7-10 d) tubes of ATCC medium 2463.
4. Screw caps on tightly and incubate at a  $15^{\circ}$  horizontal slant at  $35^{\circ}\text{C}$ .
5. Subculture when many trophozoites are observed (typically every 2-4 days). The transfer interval will depend on the quantity of the inoculum and the quality of the medium. This should be empirically determined by examining the culture on a daily basis until the growth cycle has stabilized. Do not allow the culture to overgrow. The culture crashes soon after reaching peak density.

### Reagents for cryopreservation: CPMB-5 Cryoprotective Solution

DMSO, 1.0 mL

2.5 M Sucrose, 0.8 mL

L-Cysteine/ Ascorbic Acid Solution, 0.2 mL

CPMB-2 Basal Solution, 6.0 mL

HIBS, 2.0 mL

### CPMB-2 Basal Solution

Yeast Extract, 60.0 g  
K<sub>2</sub>HPO<sub>4</sub>, 1.0 g  
KH<sub>2</sub>PO<sub>4</sub>, 0.6 g  
NaCl, 2.0 g  
Distilled water, 1.0 L  
Autoclave for 15 minutes

#### L-Cysteine/ Ascorbic Acid Solution

L-Cysteine-HCl, 1.0 g  
Ascorbic Acid, 0.1 g  
Distilled water, 10.0 mL

Add 9.0 mL of distilled water to a 20 mL beaker and dissolve the first two components. While stirring, adjust the pH to 7.2 with 10N NaOH (approximately 0.7 mL). Adjust final volume to 10 mL with distilled water and filter sterilize. Solution should be used soon after preparation. Discard any unused solution.

#### **Cryopreservation:**

#### **Harvest and Preservation**

1. Harvest cells from several cultures that are in the late logarithmic to early stationary phase of growth. Place culture vessels on ice for 10 min.
2. Invert tubes 20 times and centrifuge at 200 x g for 5 min.
3. While cells are centrifuging, prepare the cryoprotective solution.
  - a. Place 1.0 mL of DMSO in a 16 x 125 mm screw-capped test tube and ice until solidified.
  - b. Add 0.8 mL of the 2.5 M Sucrose solution, remove from ice and invert until the DMSO is liquefied. Return to ice bath.
  - c. Add 0.2 mL of the L-Cysteine/Ascorbic Acid Solution to the DMSO solution and mix.
  - d. Add 6.0 mL of the CPMB-2 Basal Solution and mix.
  - e. Add 2.0 mL HIBS and mix.
4. Resuspend the cell pellets and pool to a final volume of approximately 10 mL with the supernatant. Make a determination of the cell density and adjust the concentration of the cells between 5 x 10<sup>5</sup>/mL - 1 x 10<sup>6</sup>/mL using fresh medium. If the cell concentration is below 5 x 10<sup>5</sup>/mL, centrifuge the cell suspension and resuspend the pellet in a volume that will yield the desired

concentration.

5. After the cell concentration is adjusted, centrifuge as in step 2.
6. Remove as much supernatant as possible and determine the volume removed.
7. Resuspend the cell pellet with a volume of the cryoprotective solution equal to the volume of the supernatant removed. Invert the tube several times to obtain a uniform cell density.
8. Dispense 0.5 mL aliquots into 1.0 - 2.0 mL plastic sterile cryules (special plastic vials for cryopreservation).
9. Place the vials in a controlled rate freezing unit. Use the following cooling cycle: From room temperature cool at  $-10^{\circ}\text{C}/\text{min}$  to the heat of fusion; from the heat of fusion to  $-40^{\circ}\text{C}$ , cool at  $-1^{\circ}\text{C}/\text{min}$ . At  $-40^{\circ}\text{C}$  plunge into liquid nitrogen. The cooling cycle should be initiated no less than 15 and no more than 30 minutes after the addition of DMSO to the cell preparation.
10. Store ampules in a liquid nitrogen refrigerator until needed.
11. To establish a culture from the frozen state, place an ampule in a  $35^{\circ}\text{C}$  water bath, until thawed (2-3 min). Immerse the vial just sufficiently to cover the frozen material. Do not agitate the ampule.
12. Transfer contents of thawed ampule to a 16 x 125 mm screw-capped borosilicate glass test tube containing 13 mL of ATCC medium 2463.
13. Screw cap on tightly and incubate at a  $15^{\circ}$  horizontal slant at  $35^{\circ}\text{C}$ . Observe the culture daily and transfer when many trophozoites are observed.

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## Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner: *Entamoeba histolytica* Schaudinn (ATCC 30459)

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## References

References and other information relating to this material are available at [www.atcc.org](http://www.atcc.org).

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## Warranty

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