



# *Entamoeba histolytica* Schaudinn

50412™

## Description

**Strain designation:** SAW 891 R Clone B

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

**Type strain:** No

## Storage Conditions

**Product format:** Frozen

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

## Intended Use

This product is intended for laboratory research use only. It is not intended for any animal or human therapeutic, 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 *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 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 vials that require storage in liquid nitrogen, it is important to note that some vials may leak when submerged in liquid nitrogen. The vials will slowly fill with liquid nitrogen. Upon thawing, the conversion of the liquid nitrogen back to its gas phase may cause the vial to expand and potentially leak.

vial exploding or blowing off its cap with dangerous force creating flying debris. Unless necessary, 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 2154: LYI Entamoeba medium

### Instructions for complete medium:

ATCC Medium PRA-2154

(Quality controlled freeze-dried lots of this medium are commercially available from ATCC).

**Temperature:** 35°C

**Atmosphere:** Anaerobic

**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 facilities are not available, frozen ampules may be stored at or below -70°C for approximately one week. **Do not under any circumstance store frozen ampules at refrigerator freezer temperatures (generally -20°C).** Storage of frozen ampules at this temperature will result in the death of the culture.

1. To thaw a frozen ampule, place in a 35°C water bath, until thawed (2-3 min). Immerse the ampule just sufficient to cover the frozen material. Do not agitate the ampule.

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2. Immediately after thawing, aseptically transfer contents to a glass screw-capped tube containing 2154. Screw cap on tightly and incubate on a 15° horizontal slant at 35°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 7-10d) tubes of ATCC medium.
4. Screw caps on tightly and incubate at a 15° horizontal slant at 35°C.
5. Subculture every 10-14 days.

**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 7.0 with 10N NaOH (approximately 0.7 ml). Adjust final volume to 10 ml with distilled water and filter sterilize. Solution is stable for 1 month. Discard any unused solution.

**Cryopreservation:**

1. Harvest cells from several cultures that are in the late logarithmic to early stationary phase of growth. Harvest into 1.5 ml microcentrifuge vessels on ice for 10 min.

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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. Re
  - 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  $5 \times 10^5/\text{ml}$  -  $1 \times 10^6/\text{ml}$  in the medium. If the cell concentration is below  $5 \times 10^5/\text{ml}$ , 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 cryovials (special plastic vials for cryopreservation).
9. Place the vials in a controlled rate freezing unit. Use the following cooling cycle: From room temperature to  $-10^\circ\text{C}/\text{min}$  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 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). Add a vial just sufficient 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 ATCC medium 2154.
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 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 50412)

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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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Product Sheet

Please see the material transfer agreement (MTA) for further details regarding the use of this product at [www.atcc.org](http://www.atcc.org).

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