




Product Sheet

# *Oxytricha trifallax* (ATCC® 50532™)

Please read this **FIRST**

Storage Temp.  
**Frozen: -70°C or colder**  
**Freeze-Dried: 2°C to 8°C**  
**Live Culture: See Protocols Section**

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 Biosafety Level  
**1**

## Intended Use

This product is intended for research use only. It is not intended for any animal or human therapeutic or diagnostic use.

## Citation of Strain

If use of this culture results in a scientific publication, it should be cited in that manuscript in the following manner: *Oxytricha trifallax* (ATCC® 50532™)

American Type Culture Collection  
PO Box 1549  
Manassas, VA 20108 USA  
[www.atcc.org](http://www.atcc.org)

800.638.6597 or 703.365.2700  
Fax: 703.365.2750  
Email: [Tech@atcc.org](mailto:Tech@atcc.org)

Or contact your local distributor

## Description

**Strain Designation:** SLC 89  
**Deposited Name:** *Oxytricha trifallax*  
**Depositor:** GA Herrick  
**Isolation:** Unknown

## Propagation

### Growth Conditions

**Temperature:** 20°C to 25°C  
**Culture system:** Xenic, with *Enterobacter aerogenes* ATCC 13048 and unidentified microalgae as food sources

### Medium

ATCC® Medium 802: Sonneborn's Paramecium medium

### Instructions for Complete Medium

**Media:** ATCC Medium 802 inoculated with *Klebsiella pneumoniae* subsp. *pneumoniae* (ATCC® 700831™) or *Enterobacter aerogenes* (ATCC® 13048™). Culture also contains unidentified microalgae as food source.

## Protocols

### Handling of Live Culture

This strain is routinely shipped as a growing culture in a glass 16 x 125 mm screw-capped test tube. The volume of the cell suspension is approximately 5 mL. When the culture arrives remove it promptly from the shipping container. **Do not store the culture at refrigeration temperatures before handling.** To assure viability, immediately loosen the test tube cap and incubate upright at 25°C for at least one hour before observing the culture. There should be numerous active trophozoites in suspension. If the numbers are low or cysts are present, the culture may have been exposed to temperature extremes in transit. Regardless of the state of the culture, aseptically transfer a 0.5-1.0 mL aliquot to a 16 x 125 mm screw-capped test tube containing 5 mL of bacterized ATCC Medium 802. Incubate the parent and daughter cultures upright with the caps on loosely at 25°C.

### Culture Maintenance

Subculture every week to a fresh tube of bacterized medium in the following manner:

1. Transfer 0.5-1.0 mL from a growing culture to 5.0 mL of bacterized ATCC medium 802.
2. Add 1.0 mL of bacterized ATCC medium 802 twice weekly. When the tube is filled to within one inch of the top, aspirate from the bottom of the tube and reduce the volume to 5.0 mL.
3. Incubate upright at 25°C with the cap on loosely.

## Cryopreservation

### Reagents

#### Cryoprotective Solution

DMSO, 2.0 mL

Fresh growth medium w/o bacteria, 8.0 mL

### Harvest and Preservation


1. Mix the components in the order listed. When the medium is added to the DMSO the solution will warm up due to chemical heat.
2. Harvest cells from a culture that is at or near peak density by centrifugation at 125 x g for 5 min.
3. Adjust the concentration of cells at least  $1 \times 10^5$ /mL in fresh medium.
4. Mix the cell preparation and the cryoprotective solution in equal portions.
5. Dispense in 0.5 mL aliquots into 1.0 - 2.0 mL sterile plastic screw-capped cryovials (special plastic vials for cryopreservation).
6. Place vials in a controlled rate freezing unit. From room temperature cool at -1°C/min to -40°C. If freezing unit can compensate for the heat of fusion, maintain rate at -1°C/min through heat of fusion. At -40°C plunge ampules into liquid nitrogen.
7. Ampules are stored in either the vapor or liquid phase of a nitrogen refrigerator.
8. To establish a culture from the frozen state add 1.0 mL bacterized ATCC medium 802 to the frozen



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- ampule and place it in a 35°C water bath. Immerse the vial to a level just above the surface of the frozen material. Do not agitate the vial.
9. Immediately after thawing, do not leave in water bath, aseptically remove the contents of the ampule and inoculate onto the surface of an ATCC medium 919 (non-nutrient agar) plate containing an overlay of 15.0 mL of bacterized ATCC medium 802.
  10. Incubate at 25°C with the cap on loosely.
  11. Once the culture is established, transfer 0.5 mL to 5.0 mL of bacterized ATCC medium 802.
  12. Follow the protocol for maintenance of culture.



## References

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



## Biosafety Level: 1

Appropriate safety procedures should always be used with this material. Laboratory safety is discussed in the current publication of the *Biosafety in Microbiological and Biomedical Laboratories* from the U.S. Department of Health and Human Services Centers for Disease Control and Prevention and National Institutes for Health.

## ATCC Warranty

The viability of ATCC® products is warranted for 30 days from the date of shipment, and is valid only if the product is stored and cultured according to the information included on this product information sheet. ATCC lists the media formulation that has been found to be effective for this strain. While other, unspecified media may also produce satisfactory results, a change in media or the absence of an additive from the ATCC recommended media may affect recovery, growth and/or function of this strain. If an alternative medium formulation is used, the ATCC warranty for viability is no longer valid.

## Disclaimers

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Please see the enclosed Material Transfer Agreement (MTA) for further details regarding the use of this product. The MTA is also available on our Web site at [www.atcc.org](http://www.atcc.org)

Additional information on this culture is available on the ATCC web site at [www.atcc.org](http://www.atcc.org).

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