



# ***Naegleria* sp.**

**50353™**

## **Description**

**Strain designation:** SWL NG-256

**Deposited As:** *Naegleria* sp.

**Type strain:** No

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## **Storage Conditions**

**Product format:** Freeze-dried

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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 1**

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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 997: Fresh water ameba medium

**Instructions for complete medium:** ATCC Medium 997 grown with mixed bacteria

**Temperature:** 25-30°C

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## Handling Procedures

### Culture maintenance:

1. Remove an agar block (~5 mm<sup>2</sup>), with trophozoites or cysts, from the edge of an agar plate culture and place it in a test tube containing 1 ml of sterile ATCC medium 1325. Agitate to suspend cells from the agar block. Transfer 0.25 ml of the solution to center of each of two fresh plates and spread evenly with a spread bar.
2. Wrap the entire edge of the plate with parafilm and incubate upright at 25°C.
3. Repeat steps 1-3 at 10-14 d intervals.

### Cryopreservation:

1. Allow the cells to encyst. To detach cysts from the plate flush the surface with 5 ml fresh ATCC medium 1323 (Page's Balanced Salt Solution). Rub the surface of the plate with a spread bar to detach adhering amoebae.
  2. Transfer the cyst suspension to a sterile centrifuge tube.
  3. If the cyst concentration does not exceed  $2 \times 10^6$  cysts/ml adjust the suspension to that concentration. To adjust the concentration, centrifuge at  $600 \times g$  for 5 min and resuspend the pellet in the volume of fresh medium required to yield  $2 \times 10^6$ .
  4. While cells are centrifuging prepare a 15% (v/v) solution of sterile DMSO as follows: Add the required volume of DMSO to a glass screw-capped test tube and place it in an ice bath. Allow the DMSO to solidify. Add the required volume of refrigerated medium. Dissolve the DMSO by inverting the tube several times.
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## Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner: *Naegleria* sp. (ATCC 50353)

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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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