ATCC

Product Sheet

Salpingoeca rosetta (ATCC® PRA-390™)

Please read this FIRST

Storage Temp.

Frozen Cultures:

-70°C for 1 week; liquid N_2 vapor for long term storage

Freeze-dried Cultures:

2-8°C

Live Cultures:

See Protocols section for handling information

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: *Salpingoeca rosetta* (ATCC® PRA-390™)

American Type Culture Collection
PO Box 1549
Manassas, VA 20108 USA
www.atcc.org

800.638.6597 or 703.365.2700
Fax: 703.365.2750
Email: Tech@atcc.org

Or contact your local distributor

Description

Strain Designation: SrEpac
Depositor: T Levin
Isolation: Monoxenic culture derived from ATCC® PRA-366™

Notes

This culture should only be fed a single species of feeder bacterium, *Echinicola pacifica* KCTC 12368, which is included in the culture.

Propagation

Growth Conditions
Temperature: 25°C
Culture System: Monoxenic culture with *Echinicola pacifica* KCTC 12368

Medium
ATCC® Medium 1525: Seawater 802 medium
ATCC® Medium 1405: HESNW medium

Instructions for Complete Medium
Media: ATCC Medium 1525 uninoculated with any feeder bacteria
Alternate Media: ATCC Medium 1525 and ATCC Medium 1405 HESNW medium, combined in equal parts, uninoculated with any feeder bacteria

Note about growth media: Some xenic (bacterized) protist cultures may exhibit better growth if bacterial density in the culture is reduced to some degree. Cerophyl media can be diluted to "partial-strength" to reduce bacterial density by mixing it 1:1 or 1:2 with a suitable buffer medium such as ATCC medium 2348 or ATCC medium 1323 for freshwater cultures, or such as ATCC medium 1405 for marine cultures.

Protocols

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°C for approximately one week. **Do not under any circumstance store frozen ampules at refrigerator freezer temperatures (generally -20°C).** Storage of frozen material at this temperature will result in the death of the culture.

1. To thaw a frozen ampule, place it in a 35°C water bath such that the lip of the ampule remains above the water line. Thawing time is approximately 2 to 3 minutes. Do not agitate the ampule. Do not leave ampule in water bath after it is thawed.
2. Add the thawed contents to a T-25 flask containing 10 mL of uninoculated ATCC medium 1525.
3. Incubate with the cap tightly sealed at 25°C.

Culture Maintenance
Subculture every two weeks to a fresh T-25 flask of medium in the following manner:

1. Vigorously agitate the flask (or scrape the flask bottom using a sterile cell scraper) and aseptically transfer 0.5 mL from a growing culture to a T-25 tissue culture flask containing 10.0 mL of uninoculated ATCC medium 1525.
2. Incubate flask at 25°C with the cap on tightly.

Cryopreservation

Reagents
Cryoprotective Solution
DMSO, 2.0 mL
Fresh growth medium w/o bacteria, 8.0 mL

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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 filtration and centrifugation at 800 x g for 5 min.
3. Adjust the concentration of cells to at least 2 x 10^6/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 cryules (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. Alternatively, place the vials in a Nalgene 1°C freezing apparatus. Place the apparatus at -80°C for 1.5 to 2 hours and then plunge ampules into liquid nitrogen. (The cooling rate in this apparatus is approximately -1°C/min.)
7. Ampules are stored in either the vapor or liquid phase of a nitrogen refrigerator.
8. To establish a culture from the frozen state place the vial 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. Immediately after thawing, do not leave in water bath, aseptically remove the contents of the ampule and inoculate into a T-25 tissue culture flask containing 10 mL of uninoculated ATCC medium 1525.
9. Incubate at 25°C with the cap screwed on tightly.
10. Once the culture is established, vigorously agitate the flask and aseptically transfer 0.5 mL to 10.0 mL of uninoculated ATCC medium 1525.
11. Follow the protocol for maintenance of culture.

References

References and other information relating to this product are available online at 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

This product is intended for laboratory research purposes only. It is not intended for use in humans.

While ATCC uses reasonable efforts to include accurate and up-to-date information on this product sheet, ATCC makes no warranties or representations as to its accuracy. Citations from scientific literature and patents are provided for informational purposes only. ATCC does not warrant that such information has been confirmed to be accurate.

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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

Additional information on this culture is available on the ATCC web site at www.atcc.org.