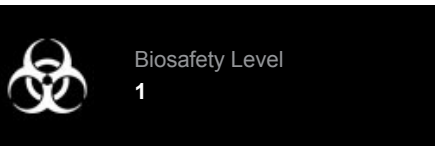
ATCC

Product Sheet

Trypanosoma lewisi (ATCC® 30213™)

Please read this FIRST



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: *Trypanosoma lewisi* (ATCC® 30213™)

Description

Strain Designation: C (Costa Rica)
Deposited Name: *Trypanosoma lewisi* (Kent) Laveran and Mesnil
Depositor: CM Lee
Isolation:
rat, Rattus rattus, San Jose, Costa Rica, 1961

Propagation

Growth Conditions
Duration: in vivo, laboratory rat
Protocol: ATCCNO: 30085 SPEC: Inject the entire thawed contents of the ampule intraperitoneally into an adult rat known to be free of blood parasites. The strain should be monitored, especially after infecting with thawed material. To infect additional rats, draw a drop of tail blood into a 1.0 ml tuberculin syringe containing 0.5 ml of sterile buffered anticoagulant, mix, and inoculate 0.1-0.2 ml of the suspension intraperitoneally/rat. Only a few trypanosomes are needed to infect a non-immune rat. However, do not wait until the blood is swarming with trypanosomes. The time for peak parasitemia may be within 4-5 days if large inocula are given. Peak parasitemia from the thawed material may be slightly sooner or later.

Instructions for Complete Medium
Grown *in vivo* laboratory mouse

Cryopreservation

1. Harvest the parasites according to the protocol for maintenance in vivo.
2. Spin the cell suspension at approximately 50 x g for 3 min, to remove the cellular debris.
3. Transfer the supernatant to a new 15 ml plastic centrifuge tube. Centrifuge at 1300 x g for 10 min.
4. Pool the cell pellets and adjust the concentration to 2.0 - 4.0 x 10^7 cells/ml with a fresh solution of Tyrode's Salt Solution.

*If the concentration is too low centrifuge at 1300 x g for 10 min and resuspend in the volume of Tyrode's Salt Solution required to yield the desired concentration.

5. Mix the cell preparation and 10% DMSO (v/v) Tyrode's Salt Solution in equal portions. The final concentration will be 1.0 - 2.0 x 10^7 cells/ml and 5% DMSO in Tyrode's Salt Solution. The time from the mixing of the cell preparation and the cryoprotective solution before the freezing process begins should be no less than 15 min. and no more than 30 min.
6. Dispense in 0.5 ml aliquots to 1.0-2.0 ml sterile plastic screw-capped cryules (special plastic vials for cryopreservation).
7. Place the vials in a controlled rate freezing unit. From room temperature cool at -1°C/min to -40°C. If the freezing unit can compensate for the heat of fusion, maintain rate at -1°C/min through the heat of fusion. At -40°C plunge 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.)
8. Store in either the vapor or liquid phase of a nitrogen refrigerator.
9. 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 thawed.
10. Immediately after thawing, aseptically remove the contents of the ampule with a syringe and inoculate an uninfected mouse. Follow the protocol for maintenance in vivo.

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.

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

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Fax: 703.365.2750
Email: Tech@atcc.org

Or contact your local distributor

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1

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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.

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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.

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