



Product Sheet

# LADMAC (ATCC® CRL-2420™)

Please read this **FIRST**



Storage Temp.  
**liquid nitrogen**  
vapor phase

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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.

## Complete Growth Medium

The base medium for this cell line is ATCC-formulated Eagle's Minimum Essential Medium, Catalog No. 30-2003. To make the complete growth medium, add the following components to the base medium: fetal bovine serum to a final concentration of 10%.

## Citation of Strain

If use of this culture results in a scientific publication, it should be cited in that manuscript in the following manner: LADMAC (ATCC® CRL-2420™)

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

**Organism:** *Mus musculus*, mouse  
**Strain:** C3H  
**Tissue:** bone marrow  
**Cell Type:** macrophage, monocyte  
**Age:** adult  
**Morphology:** lymphoblast  
**Growth Properties:** suspension, with some loosely adherent cells

## Batch-Specific Information

Refer to the Certificate of Analysis for batch-specific test results.

## SAFETY PRECAUTION

ATCC highly recommends that protective gloves and clothing always be used and a full face mask always be worn when handling frozen vials. It is important to note that some vials 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 vessel exploding or blowing off its cap with dangerous force creating flying debris.

## Unpacking & Storage Instructions

1. Check all containers for leakage or breakage.
2. Remove the frozen cells from the dry ice packaging and immediately place the cells at a temperature below  $-130^{\circ}\text{C}$ , preferably in liquid nitrogen vapor, until ready for use.

## Handling Procedure for Frozen Cells

To insure the highest level of viability, thaw the vial and initiate the culture as soon as possible upon receipt. If upon arrival, continued storage of the frozen culture is necessary, it should be stored in liquid nitrogen vapor phase and not at  $-70^{\circ}\text{C}$ . Storage at  $-70^{\circ}\text{C}$  will result in loss of viability.

1. Thaw the vial by gentle agitation in a  $37^{\circ}\text{C}$  water bath. To reduce the possibility of contamination, keep the O-ring and cap out of the water. Thawing should be rapid (approximately 2 minutes).
2. Remove the vial from the water bath as soon as the contents are thawed, and decontaminate by dipping in or spraying with 70% ethanol. All of the operations from this point on should be carried out under strict aseptic conditions.
3. It is recommended that the cryoprotective agent be removed immediately. Centrifuge the cell suspension at approximately  $125 \times g$  for 5 to 10 minutes. Discard the supernatant and resuspend the cell pellet in an appropriate amount of fresh growth medium.
4. Transfer the cell pellet to an appropriate size vessel. It is important to avoid excessive alkalinity of the medium during recovery of the cells. It is suggested that, prior to the addition of the vial contents, the culture vessel containing the complete growth medium be placed into the incubator for at least 15 minutes to allow the medium to reach its normal pH (7.0 to 7.6).
5. Incubate the culture at  $37^{\circ}\text{C}$  in a suitable incubator. A 5%  $\text{CO}_2$  in air atmosphere is recommended if using the medium described on this product sheet.

## Handling Procedure for Flask Cultures

The flask was seeded with cells (see specific batch information), grown, and completely filled with medium at ATCC to prevent loss of cells during shipping.

1. Upon receipt visually examine the culture for macroscopic evidence of any microbial contamination. Using an inverted microscope (preferably equipped with phase-contrast optics), carefully check for any evidence of microbial contamination.
2. Incubate the flask in an upright position for several hours at  $37^{\circ}\text{C}$ . After the temperature has equilibrated, aseptically remove the entire contents of the flask and centrifuge at  $125 \times g$  for 5 to 10 minutes. Remove shipping medium and save for reuse. Resuspend the cell pellet in 10 mL of this medium.
3. From this cell suspension remove a sample for a cell count and viability.
4. Incubate the culture, horizontally, at  $37^{\circ}\text{C}$  in a 5%  $\text{CO}_2$  in air atmosphere. Maintain the cell density of the culture as suggested under the subculture procedure.



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## Subculturing Procedure

Cultures can be maintained by the addition of fresh medium or replacement of medium. Alternatively, cultures can be established by centrifugation with subsequent resuspension at 1 to 2 x 10<sup>5</sup> viable cells/mL. Maintain cell density between 1 x 10<sup>5</sup> and 1 x 10<sup>6</sup> viable cells/mL. Attached cells may be subcultured by tapping the sides of the flask until cells are dispersed.

**Medium Renewal:** Add fresh medium every 2 to 3 days (depending on cell density).

### Generation of conditioned Medium

LADMAC conditioned medium is made from LADMAC cells (ATCC CRL-2420). LADMAC cells secrete the growth factor colony stimulating factor 1 (CSF-1). CSF-1 is capable of supporting the in vitro proliferation of mouse bone marrow macrophages. The Pannell-Milstein roller bottle apparatus may be used to produce high concentrations of CSF-1.[Ref](#)

1. Allow cells to become confluent.
2. After 5 to 7 days, collect supernatant, centrifuge at 125 x g for 5 to 10 minutes
3. Filter (200 nM filter)
4. Store aliquots at -20°C.



## Cryopreservation Medium

Complete growth medium described above supplemented with 5% (v/v) DMSO.  
Cell culture tested DMSO is available as ATCC Catalog No. 4-X.



## Comments

The cell line has monocyte-like morphology; contains nonspecific esterase; is phagocytic for latex beads; secretes lysozyme, and bears the Mac-1 antigen.

A minority of cells are Fc receptor positive and an appreciable number of cells are complement receptor 1 positive.

The cells are tumorigenic in nu+, nu+ mice but not in syngenic mice. The cells are not phagocytic for antibody or complement-coated particles; they do not constitutively secrete Interleukin-1.

LADMAC cells secrete the growth factor colony stimulating factor 1 (CSF-1). CSF-1 is capable of supporting the in vitro proliferation of mouse bone marrow macrophages.

The Pannell-Milstein roller bottle apparatus may be used to produce high concentrations of CSF-1.



## 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

ATCC® products are warranted for 30 days from the date of shipment, and this warranty is valid only if the product is stored and handled according to the information included on this product information sheet. If the ATCC® product is a living cell or microorganism, ATCC lists the media formulation that has been found to be effective for this product. 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 product. If an alternative medium formulation is used, the ATCC warranty for viability is no longer valid.

## Disclaimers

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This product is sent with the condition that you are responsible for its safe storage, handling, and use. ATCC



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

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