



## Product Sheet

### (HB-8207)

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.

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#### U.S. Patent Number:

4,504,586

#### Technical Information

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#### Product Description

**Organism:** *Mus musculus* (B cell); *Mus musculus* (myeloma), mouse (B cell); mouse (myeloma)

**Isotype:** IgG1

**Cell Type:** hybridoma: B lymphocyte

**Morphology:** lymphoblast

**Growth Properties:** suspension

#### 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

##### Part A. FROZEN CELLS

Vol./Ampule: 1.0 mL

Recommended Handling Upon Receipt: Initiate culture as soon as possible upon receipt. Thaw by rapid agitation in  $37^{\circ}\text{C}$  water bath. See instructions on back.

Dilute ampule contents with culture medium (see batch data above). Add fresh medium as cell density increases every 2-4 days.

#### Handling Procedure for Flask Cultures

##### Part B. FLASK CULTURES

Recommended Handling Upon Receipt:

Suspension Cultures: The culture flasks have been completely filled with medium for shipment. Remove the entire contents of the flask and centrifuge at  $300 \times g$  for 15 minutes. Resuspend the cell pellet as suggested under subculture procedure described above.

#### Subculturing Procedure



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#### Medium Renewal: Twice per week

Cultures can be maintained by addition or replacement of fresh medium. Start cultures at  $5 \times 10^4$  cells/ml and maintain between  $1 \times 10^5$  and  $1 \times 10^6$  cells/ml.

#### Comments

Animals were immunized with purified CSF-I.  
Spleen cells were fused with Sp2/0-Ag14 myeloma cells.  
CSF-I was purified from medium conditioned by MIA PaCa-2 (ATCC CRL-1420).

#### Propagation

#### Complete Growth Medium

HB101 medium (Hana Biologics), 99%; heat-inactivated fetal bovine serum, 1%  
Culture Medium: Iscove's modified Dulbecco's medium, 90%; fetal bovine serum, 10%.

#### 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

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