**Product Sheet** 

# R2-10F6 [R2-10F6.3C10.3F9] CRL-2358<sup>™</sup>

## Description

**Organism:** *Rattus norvegicus* (B cell); *Mus musculus* (myeloma), rat (B cell); mouse (myeloma) **Cell Type:** hybridoma: b lymphocyte **Growth properties:** Suspension

## **Storage Conditions**

Product format: Frozen

## 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.

### **BSL1**

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www.atcc.org

Page 1 of 5

## R2-10F6 [R2-10F6.3C10.3F9] CRL-2358

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.

## **Certificate of Analysis**

For batch-specific test results, refer to the applicable certificate of analysis that can be found at www.atcc.org.

## **Growth Conditions**

Temperature: 37°C

## Handling Procedures

#### Unpacking and 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°C, preferably in liquid nitrogen vapor, until ready for use.

**Complete medium:** Iscove's Modified Dulbecco's Medium with 4 mM L-glutamine adjusted to contain 1.5 g/L sodium bicarbonate supplemented with 0.05 mM 2-mercaptoethanol and 100 U/ml recombinant human IL-6, 90%; fetal bovine serum, 10%.

## R2-10F6 [R2-10F6.3C10.3F9]

CRL-2358

**Handling Procedure:** Recommended Handling Upon Receipt: Initiate culture as soon as possible upon receipt. Thaw by rapid agitation in 37°C water bath. See instructions on back. Dilute ampule contents with culture medium (see batch data above). Add fresh medium as density increases.

#### Subculturing procedure:

Medium Renewal: Every 2 to 3 days

Cultures can be maintained by addition or replacement of fresh medium. Alternately, cultures can be established by centrifugation with subsequent resuspension in fresh culture medium.

Establish new cultures at 2 X 10 exp5 viable cells/ml and maintain at between 1 X 10 exp5 and 1 X 10 exp6 viable cells/ml.

**Reagents for cryopreservation:** Complete growth medium supplemented with 5% (v/v) DMSO (ATCC 4-X)

## **Material Citation**

If use of this material results in a scientific publication, please cite the material in the following manner: R2-10F6 [R2-10F6.3C10.3F9] (ATCC CRL-2358)

## References

References and other information relating to this material are available at www.atcc.org.

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## R2-10F6 [R2-10F6.3C10.3F9] CRL-2358

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**Product Sheet** 

## R2-10F6 [R2-10F6.3C10.3F9] CRL-2358

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## Revision

This information on this document was last updated on 2024-10-25

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