Description
Organism: Homo sapiens, human
Tissue: Peripheral blood
Age: 29 years
Gender: Female
Morphology: hematopoietic
Growth properties: Suspension
Disease: Chronic Myeloid Leukemia At Blast Crisis
Cells per vial: Approximately 8.0 x 10^6 to 1.0 x 10^7
Volume: 1.0 mL

Storage Conditions
Product format: Frozen
Storage conditions: Vapor phase of liquid nitrogen

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.

BSL 1
ATCC determines the biosafety level of a material based on our risk assessment as guided by the current edition of Biosafety in Microbiological and Biomedical Laboratories (BMBL), U.S. Department of Health and Human Services. It is your responsibility to understand the hazards associated with the material per your organization's policies.
and procedures as well as any other applicable regulations as enforced by your local or national agencies.

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
Atmosphere: 95% Air, 5% CO₂

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
Complete medium: The base medium for this cell line is RPMI-1640 (ATCC® 30-2001™). To make the complete medium add the following components:

- 56 mL FBS (ATCC® 30-2020™)
- 5.6 mL L Glutamine (ATCC® 30-2214™)
- 0.5 mL 1 µM Imatinib working stock

To make the 1 µM Imatinib working stock aseptically combine the following:

25 mg Imatinib (Sigma catalog # CDS022173-25MG) and 45 mL Culture grade water (Hyclone catalog # SH30529)

Remaining Imatinib working stock aliquots should be stored at -80°C. Frozen aliquots can be stored for 1 year.

Note: Recommendation: supplement Imatinib at each reseed rather than including in culture medium bottle. The shelflife for Imatinib in culture medium is unknown, so do not use a prepared culture medium bottle for more than 30 days. Additionally, if cell growth appears to slow during cell line expansion, prepare a new bottle of fresh medium.

Handling Procedure: 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°C. Storage at -70°C will result in loss of viability.

1. Thaw the vial by gentle agitation in a 37°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. Transfer the vial contents to a centrifuge tube containing 9.0 mL complete culture medium and spin at approximately 150-400x g for 8 to 12 minutes.
4. Resuspend cell pellet with the recommended complete medium (start up seeding density: 2.5 to 3.5 x 10^5 viable cells/ml) and dispense into a 25 cm² or a 75 cm² culture flask. 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°C in a suitable incubator. A 5% CO₂ in air
atmosphere is recommended if using the medium described on this product
sheet.

Seeding densities for Subculture: Maintain from 2.0 x 10⁵ to 1.0 x 10⁶ viable cells/mL

Subculturing procedure:
Cultures can be maintained by addition or replacement of fresh medium. Start
cultures at 3 X 10⁵ cells/mL and maintain between 3 X 10⁵ and 1 X 10⁶ cells/mL.

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

Material Citation
If use of this material results in a scientific publication, please cite the material in the
following manner: LAMA84-r (ATCC CRL-3348)

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

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formulation and reagents that have been found to be effective for the product.
While other unspecified media and reagents may also produce satisfactory results, a
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Revision

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