



HCC2157

CRL-2340™

Description

HCC2157 is an epithelial cell that was isolated from the mammary gland of a 48-year-old, Black female patient with ductal carcinoma TNM stage IIIA, grade 2. The cell was deposited by AF Gazdar and AK Virmani. The cell line was initiated in 1996 and can be used in cancer research.

Organism: *Homo sapiens*, human

Cell Type: epithelial cell

Tissue: Breast; Mammary gland

Age: 48 years

Gender: Female

Morphology: epithelial

Growth properties: Suspension, multicellular aggregates

Disease: Carcinoma; Ductal; TNM stage IIIA, grade 2

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 ready for use.

Complete medium: RPMI-1640 medium, 90%; fetal bovine serum, 10%

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 $300 \times g$ for 8 minutes.
4. Resuspend cell pellet with the recommended complete medium (see the lot specific information provided on the Certificate of Analysis for the culture recommended dilution ratio) and dispense into an appropriate size culture flask (recommended on the Certificate of Analysis). 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% CO_2 in air atmosphere is recommended if using the medium described on this product sheet.

Subculturing procedure:

Cultures can be maintained by addition of fresh medium. Cultures grow as floating aggregates of large irregular cell clusters.

Partial fluid changes may be performed by setting the flask(s) upright and allowing the cells to settle. Collect a portion of the media which is then centrifuged. Discard the supernatant and resuspend the pellet in fresh media which is then added back

into the flask.

Subculture restrictions:

- Subculture when there are numerous, healthy appearing clusters present in suspension.
- Cells recover slowly from cryopreservation; it may take several weeks for cells to fully recover.
- It is recommended to keep flasks vertical to keep cells in close contact with one another.
- Do not split cells more than 1:1 in volume; doing so may overdilute the cells.

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

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: HCC2157 (ATCC CRL-2340)

References

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

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