



## Product Sheet

# HCE-2 [50.B1] (ATCC® CRL-11135™)

### Please read this FIRST



Storage Temp.  
**liquid nitrogen**  
vapor phase

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Biosafety Level  
**2**

### 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

Keratinocyte-Serum Free Medium (Gibco 17005-042) Supplemented with frozen additives included (from GIBCO):

- 1) 0.05 mg/ml bovine pituitary extract (BPE)
- 2) 5 ng/ml epidermal growth factor (EGF).

NOTE: Do not filter EGF

And also supplemented with 500 ng/ml hydrocortisone and 0.005mg/ml insulin (not included).

### Citation of Strain

If use of this culture results in a scientific publication, it should be cited in that manuscript in the following manner: HCE-2 [50.B1] (ATCC® CRL-11135™)

American Type Culture Collection  
PO Box 1549  
Manassas, VA 20108 USA  
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Fax: 703.365.2750  
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## Description

**Organism:** *Homo sapiens*, human

**Tissue:** eye, cornea

**Cell Type:** epithelial Adenovirus 12-SV40 hybrid transformed

**Gender:** male

**Morphology:** epithelial

**Growth Properties:** adherent

**Isoenzymes:**

AK-1, 1

ES-D, 1

G6PD, A

GLO-I, 2

Me-2, 1-2

PGM1, 1

PGM3, 2

**DNA Profile:**

Amelogenin: X,Y

CSF1PO: 10,13

D13S317: 8,11

D16S539: 9,11

D5S818: 11,12

D7S820: 8,10

THO1: 6,9

TPOX: 10,11

vWA: 15,18

**Cytogenetic Analysis:** aneuploid; Y chromosome is present

## 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°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°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. It is recommended that the cryoprotective agent be removed immediately. Transfer the vial contents to a 15 mL centrifuge tube and dilute with the recommended complete culture medium. Centrifuge the cell suspension at approximately 125 x g for 5 to 10 minutes. Discard the supernatant and resuspend the cell pellet in an appropriate amount of fresh growth medium. **NOTE: Seed cells on flasks precoated with a mixture of 0.01 mg/ml fibronectin, 0.03 mg/mL bovine collagen type I and 0.01 mg/mL bovine serum albumin.**
4. Transfer the cells 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 growth medium be placed into the incubator for at least 15 minutes to



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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<sub>2</sub> 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. Also check to determine if the majority of cells are still attached to the bottom of the flask; during shipping the cultures are sometimes handled roughly and many of the cells often detach and become suspended in the culture medium (but are still viable).
- 2. **If the cells are still attached**, aseptically remove all but 5 to 10 mL of the shipping medium. The shipping medium can be saved for reuse. Incubate the cells at 37°C in a 5% CO<sub>2</sub> in air atmosphere until they are ready to be subcultured.
- 3. **If the cells are not attached**, aseptically remove the entire contents of the flask and centrifuge at 125 x g for 5 to 10 minutes. Remove shipping medium and save. Resuspend the pelleted cells in 10 mL of this medium and add to **precoated** 25 cm<sup>2</sup> flask. Incubate at 37°C in a 5% CO<sub>2</sub> in air atmosphere until cells are ready to be subcultured.



**Subculturing Procedure**

Volumes are given for a 75 cm<sup>2</sup> flask. Increase or decrease the amount of dissociation medium needed proportionally for culture vessels of other sizes.

- 1. Remove and discard culture medium.
- 2. Briefly rinse the cell layer with 0.05% (w/v) Trypsin-0.53mM EDTA solution.
- 3. Add 2.0 to 3.0 mL of Trypsin-EDTA solution to flask and observe cells under an inverted microscope until cell layer is dispersed (usually with 5 to 15 minutes).  
Note: To avoid clumping do not agitate the cells by hitting or shaking the flask while waiting for the cells to detach. Cells that are difficult to detach may be placed at 37°C to facilitate dispersal.
- 4. Add 6.0 to 8.0 mL of complete growth medium and aspirate cells by gently pipetting. (Or neutralize with medium containing 10% fetal bovine serum).
- 5. To remove trypsin-EDTA solution, transfer cell suspension to centrifuge tube and spin at approximately 125 x g for 5 to 10 minutes.
- 6. Discard supernatant and resuspend cells in fresh growth medium. Add appropriate aliquots of cell suspension to new culture vessels. Seed cells on flasks precoated with a mixture of 0.01 mg/ml fibronectin, 0.03 mg/mL bovine collagen type I and 0.01 mg/mL bovine serum albumin.
- 7. Incubate cultures at 37°C.

**Subcultivation Ratio:** A subcultivation ratio of 1:3 is recommended

**Medium Renewal:** Twice per week



**Cryopreservation Medium**

85% Growth Medium, 10% FBS, 5% DMSO  
Cell culture tested DMSO is available as ATCC Catalog No. 4-X.



**Comments**

Keratinocyte Serum-Free medium is available from GIBCO, Grand Island, New York.



**References**

References and other information relating to this product are available online at [www.atcc.org](http://www.atcc.org).



**Biosafety Level: 2**

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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### Disclaimers

This product is intended for laboratory research purposes only. It is not intended for use in humans.

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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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