



VCaP

CRL-2876™

Product Sheet

Description

VCaP is an epithelial cell that was isolated from a White, 59-year-old, male patient with prostate cancer. This cell line was deposited by KJ Pienta in 1997.

Organism: *Homo sapiens*, human

Cell Type: epithelial cell

Tissue: Prostate

Age: 59 years

Gender: Male

Morphology: epithelial

Growth properties: Adherent

Disease: Cancer

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 2

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.

Cells produce the mouse xenotropic retrovirus Bxv-1

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: The base medium for this cell line is ATCC-formulated Dulbecco's Modified Eagle's Medium, Catalog No. 30-2002. To make the complete growth medium, add the following components to the base medium: fetal bovine serum to a final concentration of 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 280 x g for 10 minutes.
4. Resuspend cell pellet with the recommended complete medium (see the specific lot information on the Certificate of Analysis for culture recommended dilution ratio) and dispense into a 25 cm² or a 75 cm² culture flask as 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% CO₂ in air atmosphere is recommended if using the medium described on this product.

Comments

CRL-2876 is a very slow growing cell line and can take up to 48 hours to attach post-thaw and after subcultures. It can routinely take a minimum of 2 weeks or

more for cells to reach approximately 50% confluence with a dense mixture of adherent cells, floating clusters and moderate to heavy debris is always present. The cells may recover better in a T-25 flask compared to a T75 flask. Do not discard any floating cells that may be present during medium changes and subcultures, instead, spin them down using gentle centrifugation and add them back to the adherent population. The cells attach in small tightly formed clusters and some single cells. As the cells attach and start to proliferate and spread, they will grow as flattened epithelial like islands of tightly packed cells.

Subculturing procedure:

Volumes are given for a 75 cm² flask. Increase or decrease the amount of dissociation medium needed proportionally for culture vessels of other sizes. Corning® T-75 flasks (catalog #430641) are recommended for subculturing this product.

1. Remove and discard culture medium.
2. Briefly rinse the cell layer with Hank's Balanced Salt Solution or 0.25% (w/v) Trypsin- 0.53 mM EDTA solution to remove all traces of serum that contains trypsin inhibitor.
3. Add 1.0 to 2.0 mL of Trypsin-EDTA solution to flask and observe cells under an inverted microscope until cell layer is dispersed (usually within 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.
5. Add appropriate aliquots of the cell suspension to new culture vessels.
An inoculum of 2×10^4 to 4×10^4 viable cells/cm² is recommended.
6. Incubate cultures at 37°C.

Interval: Subculture when the cell concentration reaches between 1×10^5 and 2×10^5 cells/cm².

Subcultivation Ratio: 1:3 to 1:4

Material Citation

If use of this material results in a scientific publication, please cite the material in the

following manner: VCaP (ATCC CRL-2876)

References

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

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