



Product Sheet

CHLA-01-MED (ATCC® CRL-3021™)

Please read this FIRST



Storage Temp.
liquid nitrogen
vapor phase



Biosafety Level
1

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

DMEM:F12 Medium (ATCC 30-2006) with 20 ng/mL human recombinant EGF, 20 ng/mL human recombinant basic FGF, and B-27 Supplement (Invitrogen, Cat. No.17504) to a final concentration of 2% (v/v)

Citation of Strain

If use of this culture results in a scientific publication, it should be cited in that manuscript in the following manner: CHLA-01-MED (ATCC® CRL-3021™)

American Type Culture Collection
PO Box 1549
Manassas, VA 20108 USA
www.atcc.org

800.638.6597 or 703.365.2700
Fax: 703.365.2750
Email: Tech@atcc.org

Or contact your local distributor

Description

Organism: *Homo sapiens*, human

Tissue:
brain/brain tumor

Disease: medulloblastoma

Cell Type: Medulloblastoma

Age: 8 years

Gender: male

Morphology: rounded

Growth Properties: single cells and tight clusters in suspension

DNA Profile:

Amelogenin: X,Y

CSF1PO: 11

D13S317: 8, 14

D16S539: 9, 11

D5S818: 11, 13

D7S820: 10

THO1: 6, 9.3

TPOX: 9, 12

vWA: 16, 17

Cytogenetic Analysis: This is a human male cell line with a chromosome range of 42-46. The karyotype contained two consistent chromosome rearrangements, der(17)t(1;17)(q11;q25) and der(18)(17;18)(q11.2;p11.2) and approximately 40% of the examined metaphase spreads contained 11-16 double minutes.

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 submerged 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 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
2. Incubate the flask in an upright position for several hours at 37°C. After the temperature has equilibrated, aseptically remove the entire contents of the flask and centrifuge at 125 x g for 5 to 10 minutes. Remove shipping medium and save for reuse. Resuspend the cell pellet in 10 mL of this medium and transfer to culture flask.
3. Incubate the culture, horizontally, at 37°C in a 5% CO₂ in air atmosphere. Maintain the cell density of the culture as suggested under the subculture procedure.

Subculturing Procedure

Note: Due to large, tight clusters formed by these cells, it may be difficult to accurately measure cell number and viability.


Protocol: Cultures can be maintained by the addition of fresh medium when there are numerous, healthy-appearing clusters present in suspension and pH of medium has decreased. Alternatively, cultures can be established by centrifugation with subsequent resuspension in ¼ volume of the conditioned medium and ¾ volume fresh medium.




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Subcultivation ratio: A subcultivation ratio of 1:2 to 1:3 is recommended.

Medium renewal: Add fresh medium every 4 to 7 days (depending on cell density).



Cryopreservation Medium

Cryoprotectant Medium

Fetal Bovine Serum (FBS), 50%; Cell Freezing Medium-DMSO (Sigma, Cat. No.C-6295), 50%

ATCC tested fetal bovine serum is available as ATCC® Catalog No. 30-2020 (500 ml) and ATCC® Catalog No. 30-2021 (100ml).



Comments

CHLA-01-MED was established from an 8 year old boy with CNS-disseminated medulloblastoma. Cells were isolated from the brain tumor resected at the time of diagnosis and were cultured in neurobasal medium after mechanical disruption. This cell line has been continuously carried in culture for over 18 months, and in neurobasal medium grows as spheres of varying sizes. When placed in medium containing fetal bovine serum the cells attach and spread, but after a period of proliferation they senesce. The generation of this cell line was made possible with the support of the Michael Hoefflin Foundation for Children's Cancer to Childrens Hospital Los Angeles, with the goal of making pediatric brain tumor lines available to the research community. A metastatic recurrent medulloblastoma cell line from this patient is available as CHLA-01R-MED (see ATCC [CRL-3034](#))



References

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



Biosafety Level: 1

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.

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Additional information on this culture is available on the ATCC web site at www.atcc.org.

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