

Product sheet

CHO-CXCR4 CHO-CXCR4 | 305411MH

Product Overview

Description CHO-CXCR4-Medium-High is a CHO cell line (Chinese hamster ovary) genetically engineered for CXCR4 surface expression (medium-high expression level). The cells are stably transfected with a CXCR4 cDNA construct under the control of a CMV promoter. The cells are grown in DMEM supplemented with 10% FBS. The cells are characterized by high CXCR4 surface expression and are suitable for antibody screening, CXCR4-targeted therapy development, HIV entry research, hematopoietic stem cell biology, and flow cytometry.

Organism Mammalia
Tissue Epithelial cells

Disease Chinese hamster ovary, non-neoplastic; genetically engineered for CXCR4 surface expression (medium-high expression level)

Applications Antibody screening; CXCR4-targeted therapy development; HIV entry research; hematopoietic stem cell biology; flow cytometry

Synonyms CHO-CXCR4

Characteristics

Age 1-3 months

Gender Male

Morphology Epithelial cells

Cell type Epithelial cells

Growth properties Adherent / Suspension

References

Citation CHO-CXCR4 (GenBank: U08888) (ATCC: CRL-2739) (305411MH)

Biosafety level 1

NCBI_TaxID 10029

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Thawing and Culturing Cells

1. Thaw the vial rapidly in a 37°C water bath. Remove the vial and centrifuge at 300 x g for 3 minutes. Discard the supernatant and resuspend the cells in 10 ml of complete medium. Seed the cells into a T75 flask containing 50 ml of complete medium.
2. Incubate the cells at 37°C in a humidified atmosphere of 5% CO₂. Once the cells have reached confluence, passage them into a new T75 flask.
3. For long-term preservation, harvest the cells by trypsinization and resuspend them in 1 ml of cryopreservation medium. Seed the cells into a cryovial containing 0.5 ml of cryopreservation medium.
4. Store the cryovial in liquid nitrogen vapor phase. Thaw the cryovial rapidly in a 37°C water bath. Remove the vial and centrifuge at 300 x g for 3 minutes. Discard the supernatant and resuspend the cells in 10 ml of complete medium. Seed the cells into a T75 flask containing 50 ml of complete medium.
5. Incubate the cells at 37°C in a humidified atmosphere of 5% CO₂. Once the cells have reached confluence, passage them into a new T75 flask.
6. For long-term preservation, harvest the cells by trypsinization and resuspend them in 1 ml of cryopreservation medium. Seed the cells into a cryovial containing 0.5 ml of cryopreservation medium.
7. Store the cryovial in liquid nitrogen vapor phase. Thaw the cryovial rapidly in a 37°C water bath. Remove the vial and centrifuge at 300 x g for 3 minutes. Discard the supernatant and resuspend the cells in 10 ml of complete medium. Seed the cells into a T75 flask containing 50 ml of complete medium.
8. Incubate the cells at 37°C in a humidified atmosphere of 5% CO₂. Once the cells have reached confluence, passage them into a new T75 flask.

Incubation Atmosphere

37°C, 5% CO₂, humidified atmosphere.

Shipping Conditions

Cryopreserved cell lines are shipped on dry ice in validated, insulated packaging with sufficient refrigerant to maintain approximately -78 °C throughout transit. On receipt, inspect the container immediately and transfer vials without delay to appropriate storage.

Storage Conditions

For long-term preservation, place vials in vapor-phase liquid nitrogen at about -150 to -196 °C. Storage at -80 °C is acceptable only as a short interim step before transfer to liquid nitrogen.

Genotyping / HLA

Sterility

Genotyping (PCR) and sterility testing are performed on all cell lines. The results of the genotyping and sterility testing are available upon request.