Product sheet

CHO-TACD2 Cells | 305415

General information

Description	**Disclaimer: The prices displayed for cell lines are exclusively for academic/not-for-profit customers. For commercial entities the price is approximately €6,250. If you represent a commercial entity or are unsure which category applies, please contact us.** The CHO-TACD2 cell line is a stable recombinant CHO (Chinese Hamster Ovary) cell line engineered to express the TACD2 receptor at a medium-high level, approximately 12,600 molecules per cell. This cell line was developed using an innovative landing pad technology, ensuring precise and reproducible integration of the TACD2 gene at a specific, pre-validated genomic locus. TACD2, also known as TROP2 or GA733-1, is a tumor-associated calcium signal transducer. It plays a critical role in intracellular calcium signaling, which is crucial for various cellular processes, including growth, division, and differentiation. Overexpression of TACD2 has been observed in various carcinomas, such as colorectal, gastric, and pancreatic cancers, making it a potential target for antibody-drug conjugates and immunotherapy. The expression of TACD2 (TROP2) in this cell line was confirmed using flow cytometry.
Organism	Chinese hamster
Tissue	Ovary
Disease	Chinese hamster ovary, non-neoplastic; genetically engineered for TACD2/TROP2 (GA733-1) surface expression at medium-high level
Applications	Antibody screening; ADC development; TROP2-targeted therapy development; colorectal/gastric/pancreatic cancer research; flow cytometry

Characteristics

Age	Adult
Gender	Female
Morphology	Epithelial-like
Cell type	Epithelial cells
Growth properties	Adherent/suspension

Regulatory Data

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Citation	CHO-TACD2 (Cytion catalog number 305415)
Biosafety level	1
NCBI_TaxID	10029
CellosaurusAccession	CVCL_A8X3
GMO Status	GMO-S1: This CHO cell line contains a TACD2 expression cassette supporting receptor-function analyses. This classification applies only within Germany and may differ elsewhere.

Biomolecular Data

Receptors expressed	TACD2 (TROP2 or GA733-1)

Handling

Culture Medium	For adherent cultures: DMEM:Ham's F12 (1:1), w: 3.1 g/L Glucose, w: 2.5 mM L-Glutamine, w: 15 mM HEPES, w: 0.5 mM Sodium pyruvate, w: 1.2 g/L NaHCO3 (Cytion article number 820400a) For suspension cultures: CHO Growth Medium A (from InSCREENeX; InSCREENeX catalog number INS-ME-1039)
Supplements	For adherent cultures: Supplement the medium with 5% FBS. Add Geneticin (G418-Sulfat) to achieve a final concentration of 0.5 mg/mL.
Dissociation Reagent	For adherent cultures: Trypsin-EDTA
Doubling time	approx. 14-16 hours
Subculturing	For routine adherent cell culture: Aspirate the old culture medium from the adherent cells, and wash them with PBS to remove any remaining medium. After aspirating the PBS, add the appropriate volume of Trypsin/EDTA solution based on the culture vessel size (e.g., 1 ml for a T25 flask, 3 ml for a T75 flask) and incubate at room temperature or 37°C for 5-10 minutes, or until the cells detach. Monitor detachment under a microscope, and gently tap the vessel if necessary to release the cells. Once detached, add complete medium to inactivate the Trypsin/EDTA, gently resuspend the cells, and transfer an aliquot of the cell suspension into a new culture vessel containing fresh medium. Place the vessel in an incubator set to 37°C with 5% CO_2, and change the medium every 2-3 days.
Split ratio	1 to 5

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Seeding density	2 to 5 x 10^4 cells/cm^2
Fluid renewal	2 to 3 times per week
Post-Thaw Recovery	After thawing, split the cells at a ratio of 1:2 to 1:3 in T25 flasks and allow the cells to recover from the freezing process and to adhere (for adherent cultures) for at least 24 hours.
Freeze medium	As a cryopreservation medium, we use complete growth medium (including FBS) + 10% DMSO for adequate post-thaw viability, or CM-1 (Cytion catalog number 800100), which includes optimized osmoprotectants and metabolic stabilizers to enhance recovery and reduce cryo-induced stress.
Thawing and Culturing Cells	1. Confirm that the vial remains deeply frozen upon delivery, as cells are shipped on dry ice to maintain optimal temperatures during transit. 2. Upon receipt, either store the cryovial immediately at temperatures below -150°C to ensure the preservation of cellular integrity, or proceed to step 3 if immediate culturing is required. 3. For immediate culturing, swiftly thaw the vial by immersing it in a 37°C water bath with clean water and an antimicrobial agent, agitating gently for 40-60 seconds until a small ice clump remains. 4. Perform all subsequent steps under sterile conditions in a flow hood, disinfecting the cryovial with 70% ethanol before opening. 5. Carefully open the disinfected vial and transfer the cell suspension into a 15 ml centrifuge tube containing 8 ml of room-temperature culture medium, mixing gently. 6. Centrifuge the mixture at 300 x g for 3 minutes to separate the cells and carefully discard the supernatant containing residual freezing medium. 7. Gently resuspend the cell pellet in 10 ml of fresh culture medium. For adherent cells, divide the suspension between two T25 culture flasks; for suspension cultures, transfer all the medium into one T25 flask to promote effective cell interaction and growth. 8. Adhere to established subculture protocols for continued growth and maintenance of the cell line, ensuring reliable experimental outcomes.
Incubation Atmosphere	37°C, 5% CO_2, 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.

Quality Control & Molecular Analysis

Sterility	Mycoplasma contamination is excluded using both PCR-based assays and luminescence-based mycoplasma detection methods. To ensure there is no bacterial, fungal, or yeast contamination, cell cultures are subjected to daily visual inspections.