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Data Sheet

ICOSL-CHO Recombinant Cell Line Cat. #: 79635

Product Description

Recombinant CHO-K1 cells constitutively expressing human ICOSL (also known as human secreted B7 homolog 2 (B7-H2), CD275, ICOSLG, and B7-related protein (B7RP), GenBank Accession No. NM 001283050.

Background

ICOSL is the Ligand for the T-cell-specific cell surface receptor ICOS. When engaged by the ICOSL, the ICOS initiates a costimulatory signal for T-cell proliferation and cytokine production. It also induces B-cell proliferation and differentiation into plasma cells. The ICOS/ICOSL is a major regulator of the adaptive immune responses and also plays an important role in mediating local tissue responses to inflammatory conditions.

Applications

This cell line is useful for screening for ICOSL antibodies through flow cytometry, immunocytochemistry or Western blot.

Format

Each vial contains ~2 x 106 cells in 1 ml of 10% DMSO

Storage

Store in liquid nitrogen immediately upon receipt.

Mycoplasma Testing

This cell line has been screened using the MycoAlert™ Mycoplasma Detection Kit (Lonza, #LT07-118) to confirm the absence of Mycoplasma contamination.

Thaw Medium 3 (BPS Bioscience #60186): F-12K Medium supplemented with 10% FBS, 1% Penicillin/Streptomycin.

Growth Medium 3D (BPS Bioscience #79539): F-12K Medium supplemented with 10% FBS, 1% Penicillin/Streptomycin plus 1 mg/ml of Geneticin to ensure recombinant expression.

Cells should be grown at 37°C with 5% CO₂ using Growth Medium 3D.



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Recommended Culture Condition

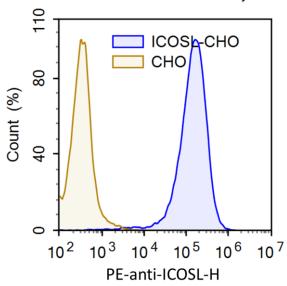
It is recommended to quickly thaw the frozen cells from liquid nitrogen in a 37°C water-bath, transfer to a tube containing 10 ml of Thaw Medium 3 (no Geneticin), spin down cells, resuspend cells in pre-warmed Thaw Medium 3 (no Geneticin), transfer re-suspended cells to T25 flask and culture in 37°C CO₂ incubator overnight. The next day, replace the medium with fresh Thaw Medium 3 (no Geneticin), and continue growing culture in a CO₂ incubator at 37°C until the cells are ready to be split. Cells should be split before they reach complete confluence. At first passage switch to Growth Medium 3D (contains 1 mg/ml Geneticin).

To passage the cells, rinse cells with phosphate buffered saline (PBS), detach cells from culture vessel with 0.05% Trypsin/EDTA, add Growth Medium 3D and transfer to a tube, spin down cells, re-suspend cells and seed appropriate aliquots of cell suspension into new culture vessels.

To freeze down the cells, rinse cells with phosphate buffered saline (PBS), and detach cells from culture vessel with 0.05% Trypsin/EDTA. After detachment, add Thaw Medium 3 (no Geneticin) and count the cells, then transfer to a tube, spin down cells, and resuspend in 4° C Freezing Medium (10% DMSO + 90% FBS) at \sim 2 x 10^{6} cells/ml. Dispense 1 ml of cell aliquots into cryogenic vials. Place vials in an insulated container for slow cooling and store at -80°C overnight. Transfer to liquid nitrogen the next day for storage. It is recommended to expand the cells and freeze down more than 10 vials of cells for future use at early passage.

Functional Validation and Assay Performance

Expression of human ICOSL in CHO-K1 cells was confirmed by FACS.



ICOSL-CHO-K1 cells (blue) and control CHO-K1 cells (yellow) were incubated with PE-labeled anti-ICOSL and detected by flow cytometry.



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Sequence

Human ICOSL sequence (accession number NM_001283050)

MRLGSPGLLFLLFSSLRADTQEKEVRAMVGSDVELSCACPEGSRFDLNDVYVYWQTSESKTV VTYHIPQNSSLENVDSRYRNRALMSPAGMLRGDFSLRLFNVTPQDEQKFHCLVLSQSLGFQEV LSVEVTLHVAANFSVPVVSAPHSPSQDELTFTCTSINGYPRPNVYWINKTDNSLLDQALQNDTV FLNMRGLYDVVSVLRIARTPSVNIGCCIENVLLQQNLTVGSQTGNDIGERDKITENPVSTGEKNA ATWSILAVLCLLVVVAVAIGWVCRDRCLQHSYAGAWAVSPETELTGHV

Related Products

<u>Product</u>	<u>Cat. #</u>	<u>Size</u>
B7-H2, Avi-His-Tag, Biotin-Labeled B7-H2, Avi-His-Tag	79300 79119	50 μg 100 μg
B7-H2 (CD275, ICOSL), Fc fusion (Human) HiP™ ICOS (CD278), Fc fusion (Human)	71130 71179	100 μg 100 μg
Thaw Medium 3 Growth Medium 3D	60186 79539	100 ml 500 ml

License Disclosure:

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