Genomeditech (Shanghai) Co., Ltd.
Order: +86 021-50432825/50432826
Email: service@genomeditech.com

Product Sheet

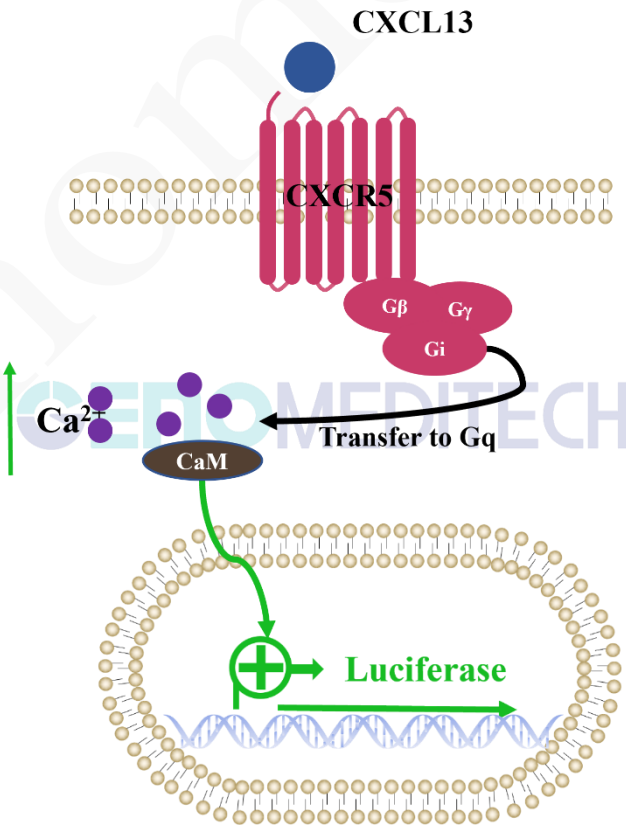
H_CXCR5 Gqi5 Reporter CHO-K1 Cell Line

Catalog number: GM-C49830

Version 4.0.26.924

CXCR5 (C-X-C motif chemokine receptor 5, also known as BLR1 or CD185) is a G protein-coupled receptor (GPCR) primarily expressed in mature B cells, follicular helper T cells (Tfh), and certain dendritic cells. The CXCR5 receptor is involved in various physiological and pathological processes, including the directed migration of lymphocytes to secondary lymphoid follicles, germinal center formation, humoral immune responses, and the development of tumor-associated tertiary lymphoid structures. It specifically recognizes and binds to its sole known ligand, CXCL13 (BLC/BCA-1). CXCR5 is natively coupled to Gi/o proteins and mediates chemotaxis by inhibiting cAMP, activating PI3K, and remodeling the cytoskeleton. It is associated with autoimmune diseases such as systemic lupus erythematosus and rheumatoid arthritis, as well as certain B-cell lymphomas, making it a valuable target in immune-targeted drug development.

H_CXCR5 Gqi5 Reporter CHO-K1 Cell Line is a clonal stable cell line generated by lentiviral technology, co-expressing human CXCR5 and the chimeric Gqi5 protein (Gq backbone with Gi C-terminal tail), together with a Gq-dependent luciferase reporter. Gqi5 redirects native Gi signaling to Gq upon receptor activation, enabling CXCL13-induced luciferase expression. Luciferase activity indicates receptor activation and supports agonist screening and functional profiling of CXCR5 ligands.



For research use only!

Specifications

Quantity	5E6 Cells per vial,1 mL
Product Format	1 vial of frozen cells
Shipping	Shipped on dry ice
Storage Conditions	Liquid nitrogen immediately upon receipt
Recovery Medium	F12K+10% FBS+1% P.S
Growth medium	F12K+10% FBS+1% P.S+4 μg/mL Blasticidin+200 μg/mL G418+4 μg/mL Puromycin
Note	None
Freezing Medium	90% FBS+10% DMSO
Growth properties	Adherent
Growth Conditions	37°C, 5% CO_2
Mycoplasma Testing	The cell line has been screened to confirm the absence of Mycoplasma species.
Safety considerations	Biosafety Level 2
Note	It is recommended to expand the cell culture and store a minimum of 10 vials at an early passage for potential future use.

Materials

Reagent	Manufacturer/Catalogue No.
F12K	BOSTER/PYG0036
Fetal Bovine Serum	ExCell/FSP500
Pen/Strep	Thermo/15140-122
Blasticidin	Genomeditech/GM-040404
Puromycin	Genomeditech/GM-040401
96 well round cell culture plate	PakGent/CL-PT096
96-well White Opaque Plate	Thermo/236108
96-well U-bottom Plate	Saining/1014010
GMOne-Step 2.0 Luciferase Reporter Gene Assay Kit	Genomeditech/GM-040513
BCA-1/CXCL13 Human	Genscript/Z02826-20
PE Rabbit anti-Human CD185/CXCR5 mAb (A26188)	abclonal/A26188

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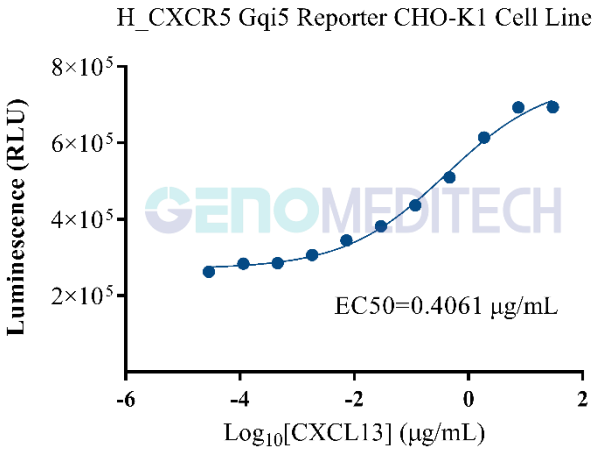
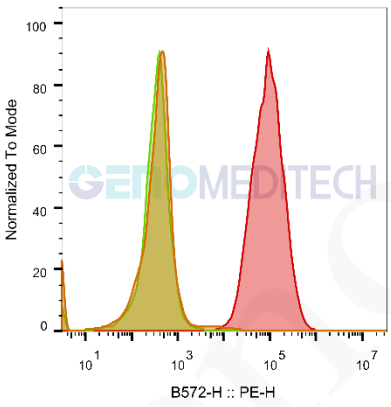
Data

Signaling Pathway



Response to BCA-1/CXCL13 Human. H_CXCR5 Gqi5 Reporter CHO-K1 Cell Line (Cat. GM-C49830) was seeded at a density of 1E4 cells per well in a 96-well plate and incubated overnight. The next day, serial dilutions of the BCA-1/CXCL13 Human (Genscript/Z02826-20) were added to the pre-seeded cells. The mixture was incubated for an additional 6 hours. The firefly luciferase activity was measured using the Luciferase Reporter Assay Kit (Genomeditech). The maximum induction fold was approximately [3.3]. Data are shown by drug mass concentration.

Expression



	SampleID	Geometric Mean : B572-H
	CHO-K1 PE Rabbit anti-Human CXCR5 Ab	325
	H_CXCR5 Gqi5 Reporter CHO-K1 PE Rabbit M_IgG Ab	323
	H_CXCR5 Gqi5 Reporter CHO-K1 PE Rabbit anti-Human CXCR5 Ab	86373

H_CXCR5 Gqi5 Reporter CHO-K1 Cell Line (Cat. GM-C49830) was determined by flow cytometry using PE Rabbit anti-Human CD185/CXCR5 mAb (A26188) (abclonal/A26188).

Cell Recovery

Recovery Medium: F12K+10% FBS+1% P.S

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.

a) 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 - 3 minutes).
b) 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.
c) Transfer the vial contents to a centrifuge tube containing 5.0 mL complete culture medium and spin at approximately 176 x g for 5 minutes. Discard supernatant.
d) Resuspend cell pellet with the recommended recovery medium. And dispense into appropriate culture dishes.
e) Incubate the culture at 37°C in a suitable incubator. A 5% CO_2 in air atmosphere is recommended if using the medium described on this product sheet.

Cell Freezing

Freezing Medium: 90% FBS+10% DMSO

a) Centrifuge at 176 x g for 3 minutes to collect cells.
b) Resuspend the cells in pre-cooled freezing medium and adjust the cell density to 5E6 cells/mL.
c) Aliquot 1 mL into each vial.
d) Place the vial in a controlled-rate freezing container and store at -80°C for at least 1 day, then transfer to liquid nitrogen as soon as possible.

Cell passage

Growth medium: F12K+10% FBS+1% P.S+4 μg/mL Blasticidin+200 μg/mL G418+4 μg/mL Puromycin

For the first 1 to 2 passages post-resuscitation, use the recovery medium. Once the cells have stabilized, switch to a growth medium.

a) Remove and discard culture medium.
b) Briefly rinse the cell layer with PBS to remove all traces of serum that contains trypsin inhibitor.
c) Add 1.0 mL of 0.25% (w/v) Trypsin-EDTA solution to dish and observe cells under an inverted microscope until cell layer is dispersed (usually within 2 to 3 minutes at 37°C).
d) 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.
e) Add 2.0 mL of growth medium to mix well and aspirate cells by gently pipetting.
f) After centrifugation, resuspend the pellet and add appropriate aliquots of the cell suspension to new culture vessels.
g) Incubate cultures at 37°C.

Subcultivation Ratio: A subcultivation ratio of 1:4 - 1:5 is recommended

Medium Renewal: Every 2 to 3 days

Notes

a) After the stabilization of the cell condition, there will be fewer dead cells post-passage, the cell growth rate will tend to stabilize, cell morphology will become uniform, and the cells will appear robust.

Related Products

CXCL13:CXCR5	
Cynomolgus_CXCR5 CHO-K1 Cell Line	Cynomolgus_CXCR5 HEK-293 Cell Line
H_CXCR5 CHO-K1 Cell Line	H_CXCR5 HEK-293 Cell Line
Mouse_CXCR5 HEK-293 Cell Line	H_CXCR5 BaF3 Cell Line
Anti-CXCR5 hIgG1 Antibody(h11G2)	

License Agreement

By purchasing and using this cell line product, the user voluntarily agrees to accept and abide by the following policies:

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