Genomeditech (Shanghai) Co.,Ltd.
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Product Sheet

H_IL-5 Reporter 293 Cell Line

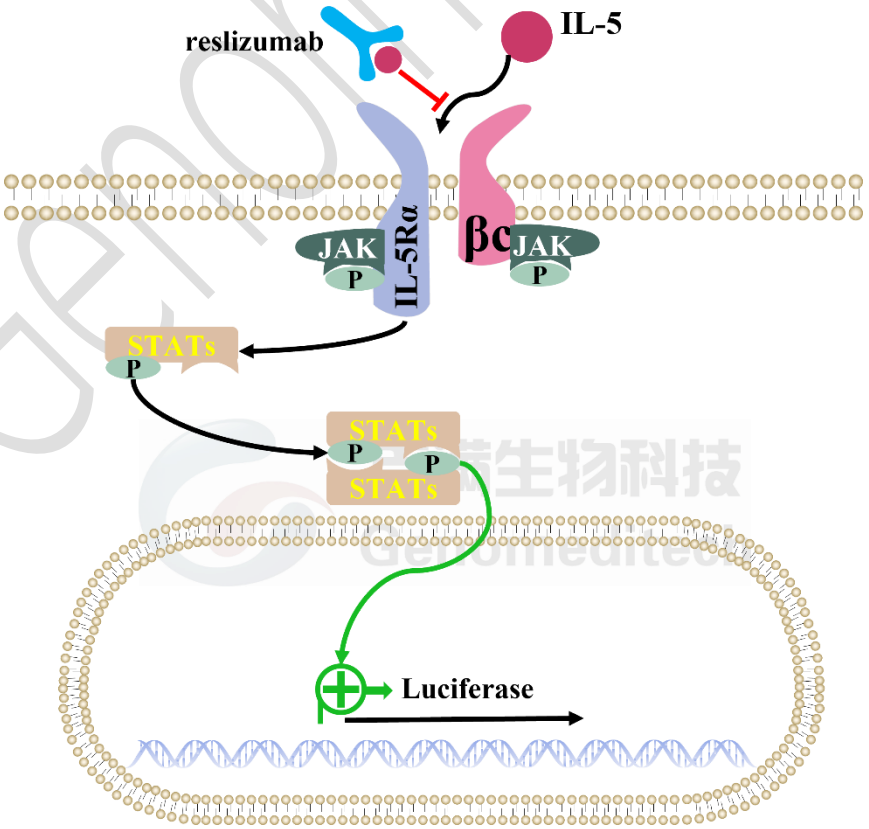
Catalog number: GM-C06723

Version 3.3.1.241225

Interleukin-5 (IL-5) is a cytokine produced by T cells, especially helper T cells, and eosinophils. It regulates the growth, differentiation, and activation of eosinophils, promoting their proliferation and survival. IL-5 is important in immune diseases like allergic reactions and asthma, enhancing eosinophil function and contributing to tissue damage and inflammation.

IL-5 activates signaling pathways by binding to its receptor IL-5R, a dimer of IL-5Rα and β chains. This binding recruits signal transducer and activator of transcription 5 (STAT5), which, upon phosphorylation, moves to the nucleus to regulate genes related to eosinophil proliferation and survival. The IL-5 signaling pathway promotes eosinophil proliferation and enhances their function, playing a key role in allergic and inflammatory responses.

H_IL-5 Reporter 293 Cell Line is a clonal stable cell line constructed using lentiviral technology, constitutive expression of the IL-5Rα and β, along with signal-dependent expression of a luciferase reporter gene. When IL-5 binds to IL-5R, it activates downstream signaling pathways, leading to the expression of luciferase. Blockade antibodies can inhibit this signal transmission. The luciferase activity measurement indicates the activation level of the signaling pathway and can thus be used to evaluate the in vitro effects of drugs related to IL-5.



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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	DMEM+10% FBS+1% P.S
Growth medium	DMEM+10% FBS+1% P.S+4 μg/mL Blasticidin+400 μg/mL G418+125 μg/mL Hygromycin+0.75 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.
DMEM	VivaCell/C3110-0500
Fetal Bovine Serum	Cegrogen biotech/A0500-3010
Pen/Strep	Thermo/15140-122
Blasticidin	Genomeditech/GM-040404
G418	Genomeditech/GM-040402
Hygromycin	Genomeditech/GM-040403
Puromycin	Genomeditech/GM-040401
Human IL-5 / Interleukin 5 Protein	Sino Biological/15673-HNCE
Anti-IL5 hIgG4 Antibody(Reslizumab)	Genomeditech/GM-52408AB
GMOne-Step Luciferase Reporter Gene Assay Kit	Genomeditech/GM-040503

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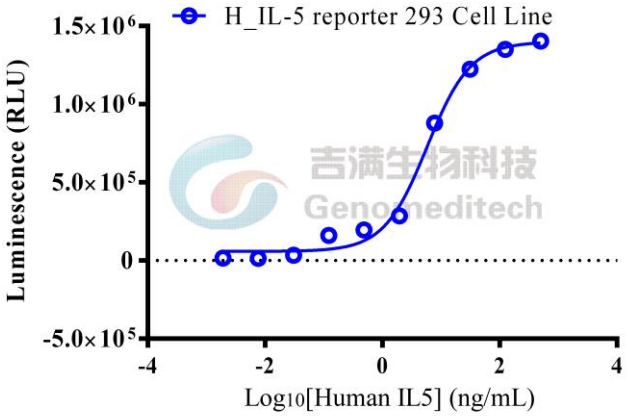
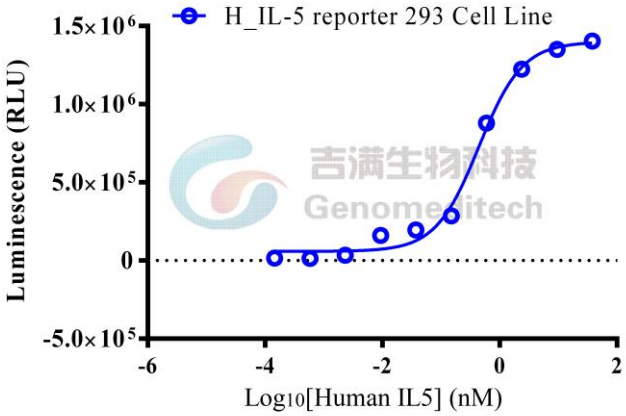
Figures



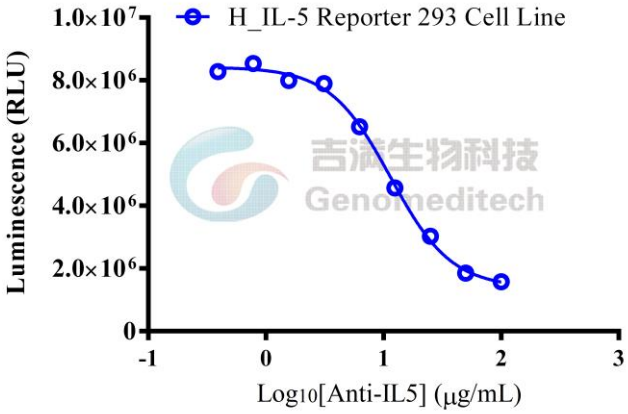
	H_IL-5 reporter 293 Cell Line
EC50	5.805

Figure 1 | Response to Human IL5 / Interleukin 5 Protein. The H_IL-5 Reporter 293 Cell Line (Cat. GM-C06723) at a concentration of 1.5E4 cells/well (96-well format) was stimulated with serial dilutions of Human IL5 / Interleukin 5 Protein (Sino Biological/15673-HNCE) in assay buffer (DMEM + 1% FBS + 1% P.S) for 16 hours. The firefly luciferase activity was measured using the GMOne-Step Luciferase Reporter Gene Assay Kit (Cat. GM-040503). The maximum induction fold was approximately [90.5]. Data are shown by drug mass concentration.



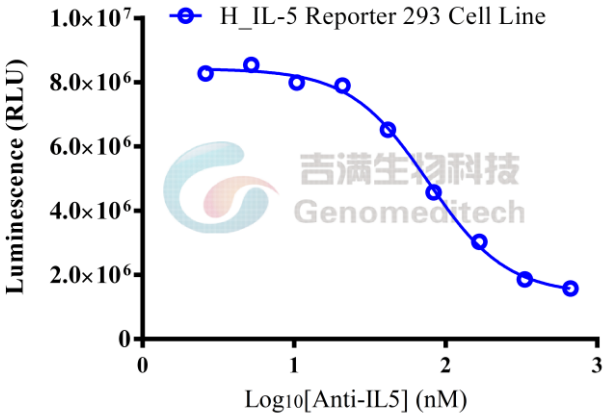
	H_IL-5 reporter 293 Cell Line
EC50	0.4431

Figure 2 | Response to Human IL5 / Interleukin 5 Protein. The H_IL-5 Reporter 293 Cell Line (Cat. GM-C06723) at a concentration of 1.5E4 cells/well (96-well format) was stimulated with serial dilutions of Human IL5 / Interleukin 5 Protein (Sino Biological/15673-HNCE) in assay buffer (DMEM + 1% FBS + 1% P.S) for 16 hours. The firefly luciferase activity was measured using the GMOne-Step Luciferase Reporter Gene Assay Kit (Cat. GM-040503). The maximum induction fold was approximately [90.5]. Data are shown by drug molar concentration.



	H_IL-5 Reporter 293 Cell Line
IC50	11.50

Figure 3 | Response to Anti-IL5 hIgG4 Antibody(Reslizumab). Serial dilutions of Anti-IL5 hIgG4 Antibody(Reslizumab) (Cat. GM-52408AB) was incubated with 0.581 ng/well of Human IL5 / Interleukin 5 Protein (Sino Biological/15673-HNCE) for 1 hour in assay buffer (DMEM + 1% FBS + 1% P.S). After pre-incubation, add the mixture to the H_IL-5 Reporter 293 Cell Line (Cat. GM-C06723) at a density of 1.5E4 cells/well in a 96-well format, and incubate for 16 hours. Firefly luciferase activity is then measured using the GMOne-Step Luciferase Reporter Gene Assay Kit (Cat. GM-040503). The results indicated maximum blocking folds of approximately [5.2]. Data are shown by drug mass concentration.



	H_IL-5 Reporter 293 Cell Line
IC50	76.64

Figure 4 | Response to Anti-IL5 hIgG4 Antibody(Reslizumab). Serial dilutions of Anti-IL5 hIgG4 Antibody(Reslizumab) (Cat. GM-52408AB) was incubated with 0.581 ng/well of Human IL5 / Interleukin 5 Protein (Sino Biological/15673-HNCE) for 1 hour in assay buffer (DMEM + 1% FBS + 1% P.S). After pre-incubation, add the mixture to the H_IL-5 Reporter 293 Cell Line (Cat. GM-C06723) at a density of 1.5E4 cells/well in a 96-well format, and incubate for 16 hours. Firefly luciferase activity is then measured using the GMOne-Step Luciferase Reporter

Gene Assay Kit (Cat. GM-040503). The results indicated maximum blocking folds of approximately [5.2]. Data are shown by drug molar concentration.

Cell Recovery

Recovery Medium: DMEM+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: DMEM+10% FBS+1% P.S+4 μg/mL Blasticidin+400 μg/mL G418+125 μg/mL Hygromycin+0.75 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) Subculturing is necessary when the cell density reaches 80%. It is recommended to perform subculturing at a ratio of 1:3 to 1:4 every 2-3 days. Ensure that the density does not exceed 80%, as overcrowding can lead to reduced viability due to compression.
b) Remove and discard culture medium.
c) Briefly rinse the cell layer with PBS to remove all traces of serum that contains trypsin inhibitor.

d) 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 30 to 60 seconds at 37°C).

e) 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.

f) Add 2.0 mL of growth medium to mix well and aspirate cells by gently pipetting.

g) After centrifugation, resuspend the pellet and add appropriate aliquots of the cell suspension to new culture vessels.

h) Incubate cultures at 37°C.

Subcultivation Ratio: A subcultivation ratio of 1:3 - 1:4 is recommended

Medium Renewal: Every 2 to 3 days

Notes

a) Upon initial thawing, a higher number of dead cells is observed, which is a normal phenomenon. Significant improvement is seen after adaptation. Once the cells reach a stable state, the number of dead cells decreases after subculturing and the cell growth rate becomes stable.

b) Ensure that the cell density does not exceed 80%, as overcrowding may lead to reduced viability due to compression.

Related Products

IL-4/IL-13	
IL-4 Reporter Cell Line	IL-4/IL-13 Reporter 293 Cell Line
IL-4/IL-13 Reporter 293 DDX35TM Cell Line	Cynomolgus_IL4R CHO-K1 Cell Line
H_IL4R CHO-K1 Cell Line	
Anti-IL-4R hIgG1 Antibody(12B5)	Anti-IL4R hIgG4 Antibody(Dupilumab)
Anti-IL4R hIgG4 Reference Antibody (Dupbio)	
Human IL-4R alpha Protein; mFc Tag	
TSLP:TSLPR	
H_TSLP Reporter Cell Line	H_TSLPR CHO-K1 Cell Line
Anti-H_TSLPR hIgG1 Antibody	Anti-TSLP hIgG2 Reference Antibody(Tezbio)
Anti-TSLP hIgG2 Antibody(Tezepelumab)	
Cynomolgus TSLP Protein; His Tag	Human TSLP Protein; His Tag
IL-5	
H_IL-5RA CHO-K1 Cell Line	H_IL-5RA HEK-293 Cell Line
Anti-IL5 hIgG4 Antibody(Reslizumab)	Anti-IL-5R hIgG1 Antibody(Benralizumab)

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