Genomeditech (Shanghai) Co.,Ltd.
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Product Sheet

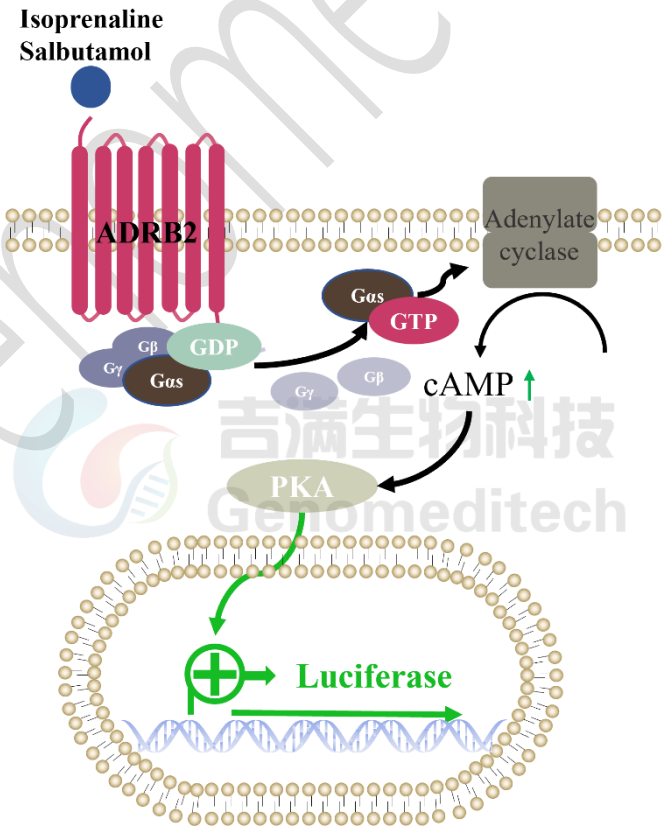
H_ADRB2 Gs Reporter CHO-K1 Cell Line

Catalog number: GM-C42210

Version 3.3.1.260408

ADRB2 is a key member of the neurokinin receptor family within the G protein–coupled receptor (GPCR) superfamily. It is primarily distributed in tissues such as bronchial smooth muscle, vascular smooth muscle, cardiac muscle, uterine smooth muscle, and immune cells (e.g., mast cells, T cells). Upon Isoprenaline binding, H_ADRB2 can couple to Gs, activating adenylyl cyclase (AC) to convert ATP into cyclic adenosine monophosphate (cAMP). Increased cAMP activates protein kinase A (PKA), which phosphorylates downstream targets to regulate gene expression, metabolism, and neuronal excitability. Dysregulation of the Gs/cAMP/PKA pathway is linked to various pathological states and may influence airway smooth muscle tone, intestinal secretion and motility, and neuromodulator release associated with anxiety and depression.

H_ADRB2 Gs Reporter CHO-K1 Cell Line is a clonal stable CHO-K1 cell line constructed using lentiviral technology, constitutive expression of the ADRB2 gene, along with signal-dependent expression of a luciferase reporter gene. When Isoprenaline、Salbutamol、ATR-258 binds to the receptor, the downstream signaling pathway is activated, thereby inducing luciferase expression. 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 ADRB2.



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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	F12K+10% FBS+1% P.S
Growth medium	F12K+10% FBS+1% P.S+4 μg/mL Blasticidin+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
Isoprenaline	Selleck/E8311
Salbutamol	MCE/HY-B1037
ATR-258	MCE/HY-176742

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Figures

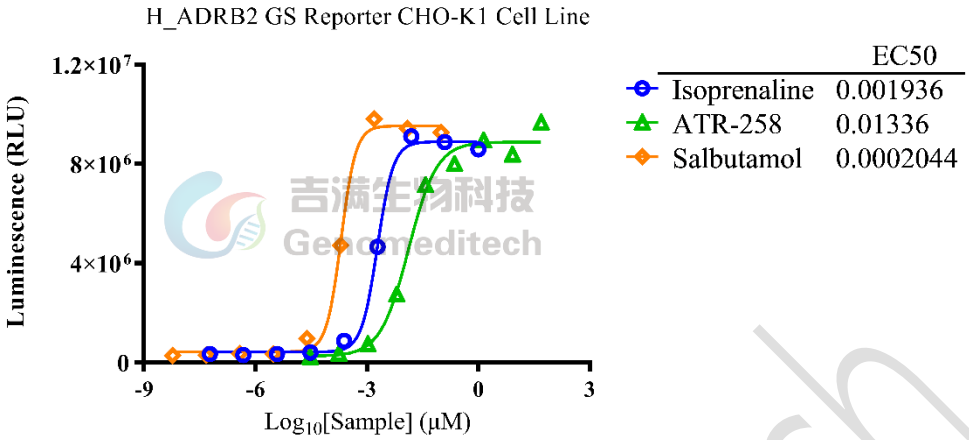

Figure 1 | Response to Isoprenaline, Salbutamol, ATR-258. The H_ADRB2 Gs Reporter CHO-K1 Cell Line (Cat. GM-C42210) at a concentration of 1E4 cells/well (96-well format) was stimulated with serial dilutions of drugs in assay buffer (F12K+1% FBS+1% P.S) for 6 hours. The firefly luciferase activity was measured using the Luciferase Reporter Assay Kit (Genomeditech).Data are shown by drug molar concentration.

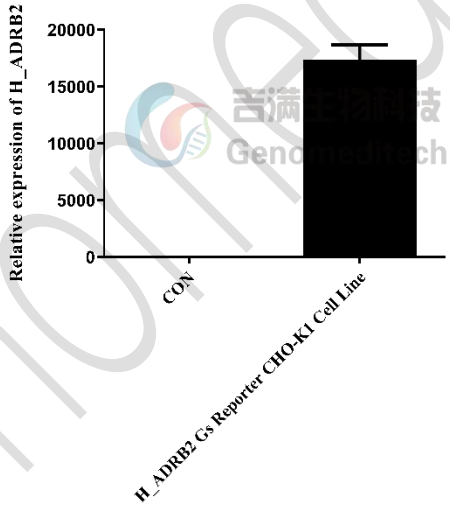

Figure 2 | The mRNA expression levels of H_ADRB2 in the H_ADRB2 Gs Reporter CHO-K1 Cell Line (Cat. GM-C42210) were determined by RT-qPCR.

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+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.

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