

# Product Sheet

## H\_CALCR RAMP1(AMY1) Reporter CHO-K1 Cell Line

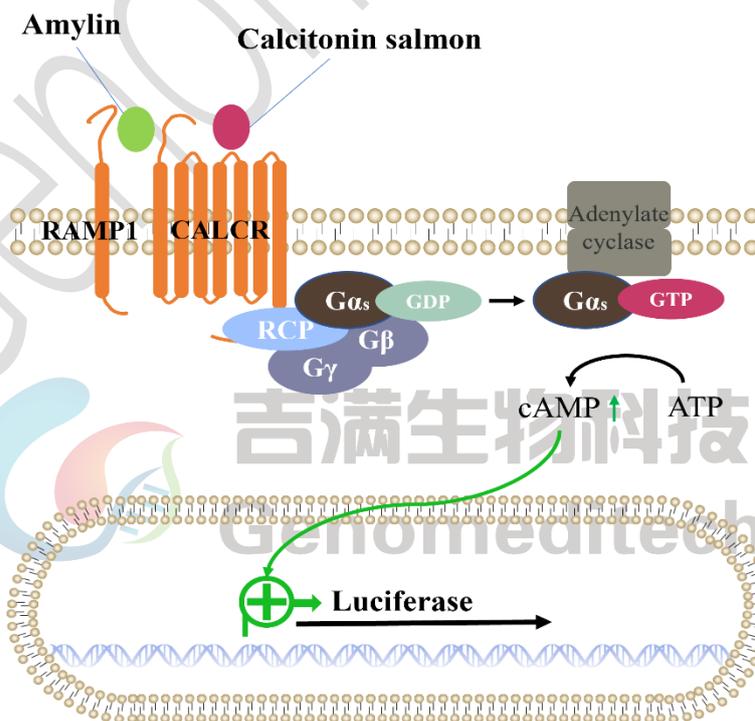
Catalog number: GM-C44040

Version 3.3.1.260316

The Calcitonin Gene Related Peptide (CGRP) family includes several important peptides: Calcitonin (CT), Amylin, and Adrenomedullin (AM). Amylin is produced by the pancreas and is a hormone that regulates nutrient intake. The binding of Amylin to its receptors involves three classes of potential receptors, which are complexes of the Calcitonin Receptor (CALCR) and Receptor Activity Modifying Proteins (RAMPs). RAMPs are a series of type I single-transmembrane proteins that function by forming heterodimers with G protein-coupled receptors. There are three RAMP subtypes: RAMP1, RAMP2, and RAMP3. Although these RAMPs are structurally similar, their amino acid sequences are only about 30% identical.

After binding to the receptor, Amylin can form different complexes: CALCR combined with RAMP1 forms the Amy1 complex, CALCR combined with RAMP2 forms the Amy2 complex, and CALCR combined with RAMP3 forms the Amy3 complex.

The H\_CALCR RAMP1(AMY1) Reporter CHO-K1 Cell Line is a clonal, stable cell line that constitutively expresses human CALCR and human RAMP1, along with a signal-dependent expression of a luciferase reporter gene. The binding of Amylin to the receptor complex activates downstream reporter genes, leading to luciferase expression. The luciferase readout indicates the activation level of the signaling pathway and can thus be used to evaluate the in vitro effects of drugs related to the CALCR RAMP1 (AMY1) complex.



## Specifications

<b>Quantity</b>	5E6 Cells per vial, 1 mL
<b>Product Format</b>	1 vial of frozen cells
<b>Shipping</b>	Shipped on dry ice
<b>Storage Conditions</b>	Liquid nitrogen immediately upon receipt
<b>Recovery Medium</b>	F12K+10% FBS+1% P.S
<b>Growth medium</b>	F12K+10% FBS+1% P.S+4 µg/mL Blasticidin+50 µg/mL Bleomycin+4 µg/mL Puromycin
<b>Note</b>	None
<b>Freezing Medium</b>	90% FBS+10% DMSO
<b>Growth properties</b>	Adherent
<b>Growth Conditions</b>	37°C, 5% CO <sub>2</sub>
<b>Mycoplasma Testing</b>	The cell line has been screened to confirm the absence of Mycoplasma species.
<b>Safety considerations</b>	Biosafety Level 2
<b>Note</b>	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.
Puromycin	Genomeditech/GM-040401
Blasticidin	Genomeditech/GM-040404
Bleomycin	Genomeditech/GM-040407
Pen/Strep	Thermo/15140-122
Fetal Bovine Serum	ExCell/FSP500
F12K	BOSTER/PYG0036
GMOne-Step 2.0 Luciferase Reporter Gene Assay Kit	Genomeditech/GM-040513
Amylin, human, amide	GenScript/RP11278CN
Eloralintide	MCE/HY-P10798
Calcitonin salmon (Salmon calcitonin)	GlpBio/GC32851

## Figures

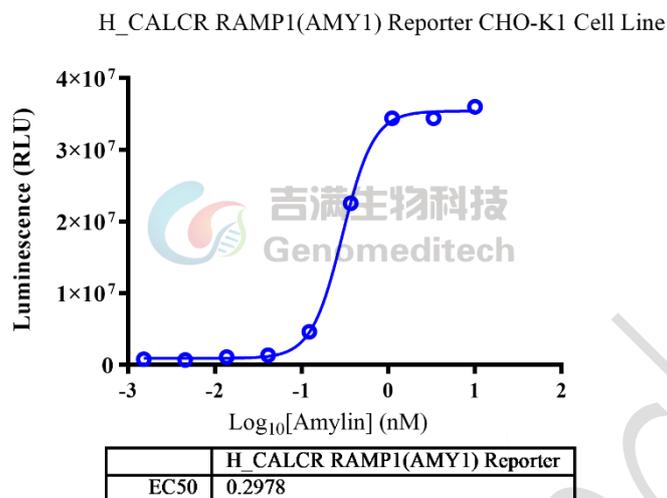


Figure 1 | Response to Amylin, human, amide. The H\_CALCR RAMP1(AMY1) Reporter CHO-K1 Cell Line (Cat. GM-C44040) at a concentration of 1E4 cells/well (96-well format) was stimulated with serial dilutions of Amylin, human, amide (Genscript/RP11278CN) in assay buffer (F12K+1% FBS+1% P.S) for 6 hours. The firefly luciferase activity was measured using the Luciferase Reporter Assay Kit (Genomeditech). The maximum induction fold was approximately [42.4]. Data are shown by drug molar concentration.

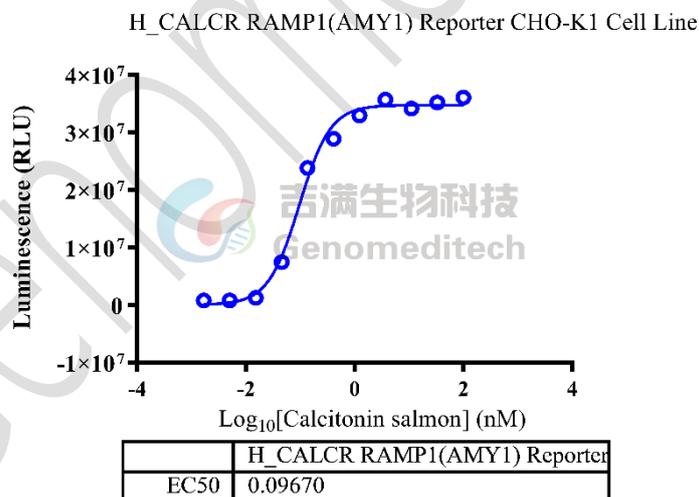


Figure 2 | Response to Calcitonin salmon. The H\_CALCR RAMP1(AMY1) Reporter CHO-K1 Cell Line (Cat. GM-C44040) at a concentration of 1E4 cells/well (96-well format) was stimulated with serial dilutions of Calcitonin salmon (Glpbio/GC32851) in assay buffer (F12K+1% FBS+1% P.S) for 6 hours. The firefly luciferase activity was measured using the Luciferase Reporter Assay Kit (Genomeditech). The maximum induction fold was approximately [43.8]. Data are shown by drug molar concentration.

H\_CALCR RAMP1(AMY1) Reporter CHO-K1 Cell Line

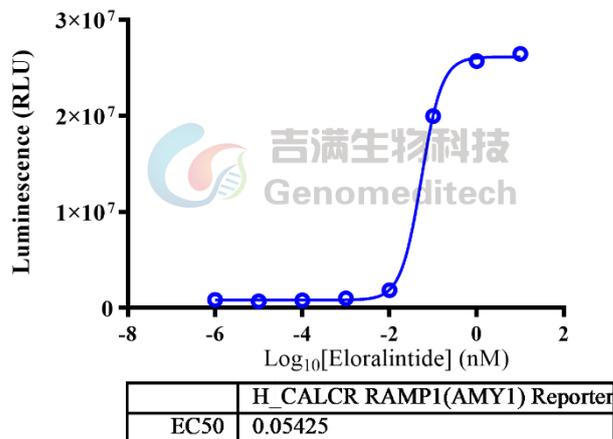


Figure 3 | Response to Eloralintide. The H\_CALCR RAMP1(AMY1) Reporter CHO-K1 Cell Line (Cat. GM-C44040) at a concentration of 1E4 cells/well (96-well format) was stimulated with serial dilutions of Eloralintide (MCE/HY-P10798) in assay buffer (F12K+1% FBS+1% P.S) for 6 hours. The firefly luciferase activity was measured using the Luciferase Reporter Assay Kit (Genomeditech). The maximum induction fold was approximately [31.1]. Data are shown by drug molar concentration.

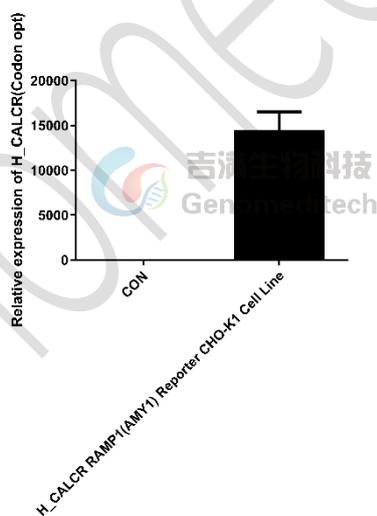


Figure 4 | The mRNA expression levels of H\_CALCR in the H\_CALCR RAMP1(AMY1) Reporter CHO-K1 Cell Line (Cat. GM-C44040) were determined by RT-qPCR.

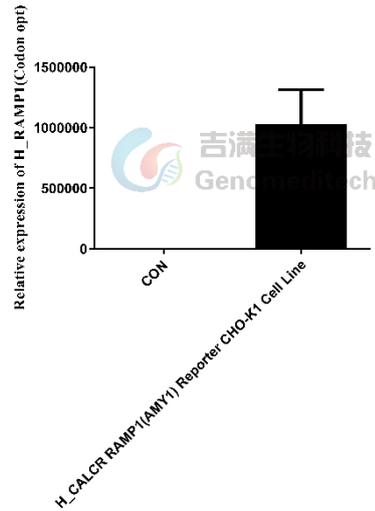


Figure 5 | The mRNA expression levels of H\_RAMP1 in the H\_CALCR RAMP1(AMY1) Reporter CHO-K1 Cell Line (Cat. GM-C44040) were determined by RT-qPCR.

## 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  $-70^{\circ}\text{C}$ . Storage at  $-70^{\circ}\text{C}$  will result in loss of viability.

- Thaw the vial by gentle agitation in a  $37^{\circ}\text{C}$  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).
- 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.
- Transfer the vial contents to a centrifuge tube containing 5.0 mL complete culture medium and spin at approximately  $176 \times g$  for 5 minutes. Discard supernatant.
- Resuspend cell pellet with the recommended recovery medium. And dispense into appropriate culture dishes.
- Incubate the culture at  $37^{\circ}\text{C}$  in a suitable incubator. A 5%  $\text{CO}_2$  in air atmosphere is recommended if using the medium described on this product sheet.

## Cell Freezing

Freezing Medium: 90% FBS+10% DMSO

- Centrifuge at  $176 \times g$  for 3 minutes to collect cells.
- Resuspend the cells in pre-cooled freezing medium and adjust the cell density to  $5 \times 10^6$  cells/mL.
- 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+50 µg/mL Bleomycin+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.

- Remove and discard culture medium.
- Briefly rinse the cell layer with PBS to remove all traces of serum that contains trypsin inhibitor.
- 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).
- 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.
- Add 2.0 mL of growth medium to mix well and aspirate cells by gently pipetting.
- After centrifugation, resuspend the pellet and add appropriate aliquots of the cell suspension to new culture vessels.
- 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

- 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

GCGR	
<a href="#">H_GCGR Reporter CHO-K1 Cell Line</a>	<a href="#">H_GCGR Reporter HEK-293 Cell Line</a>
<a href="#">H_GCGR Reporter HEK-293 DDX35TM Cell Line</a>	<a href="#">Cynomolgus_GCGR HEK-293 Cell Line</a>
<a href="#">H_GCGR CHO-K1 Cell Line</a>	<a href="#">H_GCGR HEK-293 Cell Line</a>
<a href="#">Mouse_GCGR HEK-293 Cell Line</a>	
<a href="#">Anti-H_GCGR hIgG2 Antibody(volagidemab)</a>	
GLP1R	
<a href="#">H_GLP1R Reporter CHO-K1 Cell Line</a>	<a href="#">H_GLP1R Reporter HEK-293 Cell Line</a>
<a href="#">H_GLP1R Reporter HEK-293 DDX35TM Cell Line</a>	<a href="#">H_GLP1R β-Arrestin Reporter CHO-K1 Cell Line</a>
<a href="#">Cynomolgus_GLP1R GIPR CHO-K1 Cell Line</a>	<a href="#">Cynomolgus_GLP1R HEK-293 Cell Line</a>
<a href="#">H_GLP1R CHO-K1 Cell Line</a>	<a href="#">H_GLP1R GIPR CHO-K1 Cell Line</a>
<a href="#">H_GLP1R HEK-293 Cell Line</a>	<a href="#">Mouse_GLP1R GIPR CHO-K1 Cell Line</a>
<a href="#">Mouse_GLP1R HEK-293 Cell Line</a>	<a href="#">Rat_GLP1R HEK-293 Cell Line</a>
<a href="#">Anti-GLP1R hIgG1 Antibody(mAb-36986)</a>	<a href="#">Anti-H_GLP1R hIgG1 Antibody(glutazumab)</a>

FGFR1	
H_FGF21 Reporter HEK-293 Cell Line	
Human FGF-21 Protein; His Tag	
CALCA(CGRP):CALCRL RAMP	
H_CALCRL RAMP1 Reporter HEK-293 Cell Line	H_CALCRL RAMP1 Reporter HEK-293 DDX35TM Cell Line
Cynomolgus_CALCRL RAMP1 HEK-293 Cell Line	H_CALCRL RAMP1 CHO-K1 Cell Line
H_CALCRL RAMP1 HEK-293 Cell Line	
Anti-CALCRL RAMP1 hIgG2 Antibody(Erenumab)	
GPR75	
H_GPR75 HEK-293 Cell Line	
GIPR	
H_GIPR Reporter CHO-K1 Cell Line	H_GIPR Reporter HEK-293 Cell Line
H_GIPR Reporter HEK-293 DDX35TM Cell Line	Cynomolgus_GIPR CHO-K1 Cell Line
Cynomolgus_GIPR HEK-293 Cell Line	H_GIPR CHO-K1 Cell Line
H_GIPR HEK-293 Cell Line	Mouse_GIPR CHO-K1 Cell Line
Mouse_GIPR HEK-293 Cell Line	
Anti-H_GIPR hIgG1 Antibody(AMG-133)	
ACVR2A:ACTRIIB:Active A	
ACVR2A KO HEK-293 Cell Line	ACVR2B KO HEK-293 Cell Line
Activin A Reporter Cell Line	BRE Reporter 293 Cell Line
H_ACVR2A Reporter Cell Line	H_ACVR2B Reporter Cell Line
H_ACVR2B Reporter DDX35TM Cell Line	H_ACVR2A HEK-293(ACVR2B KO) Cell Line
H_ACVR2B CHO-K1 Cell Line	H_ACVR2B HEK-293(ACVR2A KO) Cell Line
Anti-ACVR2A hIgG1 Antibody(LAE-102)	Anti-ACVR2B hIgG1 Antibody(Bimagrumab)
Anti-ACVR2B hIgG1 Antibody(Fab-17G05)	Anti-ACVR2B mIgG2a Antibody(Bimagrumab)
Anti-GDF8 hIgG4 Reference Antibody (Aptibio)	Anti-H_ACVR2B hIgG1 Reference Antibody(Bimbio)
Biotinylated Human ACVR2A Protein; His-Avi Tag	Biotinylated Human ACVR2B Protein; His-Avi Tag
Biotinylated Mouse ACVR2A Protein; His-Avi Tag	Biotinylated Mouse ACVR2B Protein; His-Avi Tag
Human Activin A Protein; His Tag	Human Activin A Protein; His Tag (CHO)
Human Activin B Protein; His Tag	Human ACVR2A Protein; hFc Tag
Human ACVR2A Protein; hFc Tag (Sotatercept)	Human ACVR2A Protein; His Tag
Human ACVR2B Protein; hFc Tag	Human ACVR2B Protein; His Tag
Human latent GDF-8 Protein; His Tag	Mouse ACVR2A Protein; His Tag
Mouse ACVR2B Protein; His Tag	
AMY:CALCR RAMP	
H_CALCR RAMP3(AMY3) Reporter CHO-K1 Cell Line	H_CALCR RAMP3(AMY3) $\beta$ -Arrestin Reporter CHO-K1 Cell Line
H_CALCR Reporter CHO-K1 Cell Line	H_CALCR $\beta$ -Arrestin Reporter CHO-K1 Cell Line
Rat_CALCR RAMP3(AMY3) Reporter COS-7 Cell Line	Rat_CALCR Reporter COS-7 Cell Line
THRβ	
H_THR $\beta$ Reporter HEK-293 Cell Line	

MC4R	
<a href="#">H_MC4R Reporter HEK-293 Cell Line</a>	
ASGR1	
<a href="#">H_ASGR1 CHO-K1 Cell Line</a>	<a href="#">H_ASGR1 HEK-293 Cell Line</a>
<a href="#">Anti-ASGR1 hIgG1 Antibody(4A2.001)</a>	
<a href="#">Cynomolgus ASGR1 Protein; His Tag</a>	<a href="#">Human ASGR1 Protein; hFc Tag</a>
<a href="#">Human ASGR1 Protein; His Tag</a>	<a href="#">Mouse ASGR1 Protein; His Tag</a>

## License Agreement:

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

- This cell line product is restricted to research use only and shall not be used for any commercial purposes.
- This product is strictly prohibited from being used in the diagnosis or treatment of human or animal diseases, and shall not be directly used in experiments involving humans.
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