

HEK293/Human GIPR Stable Cell Line (High Expression) Data Sheet

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HEK293/Human GIPR Stable Cell Line (High Expression) Data Sheet

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| Catalog No. | Size |
|-------------|--|
| CHEK-ATP206 | 2 × (1 vial contains ~5×10 ⁶ cells) |

• *Description*

The HEK293/Human GIPR Stable Cell Line was engineered to express the receptor full length human GIPR (Gene ID: 2696), with different levels of GIPR expression (High, Medium, Low). Surface expression of human GIPR was confirmed by flow cytometry.

• *Application*

- Useful for cell-based GIPR binding assay
- Screen for human GIPR agonists based on cAMP accumulation assay

• *Cell Line Profile*

| | |
|------------------------|--|
| Cell line | HEK293/Human GIPR Stable Cell Line (High Expression) |
| Host Cell | HEK293 |
| Property | Adherent |
| Complete Growth Medium | DMEM + 10% FBS |
| Selection Marker | Hygromycin B (20 µg/mL) |
| Incubation | 37°C with 5% CO ₂ |
| Doubling Time | 22-24 hours |
| Transduction Technique | Lentivirus |

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• *Materials Required for Cell Culture*

- DMEM Medium (BasalMedia, Cat. No. L120KJ)

Note: If you are unable to obtain the specified DMEM medium (BasalMedia, Cat. No. L120KJ) in China, you may use an alternative DMEM medium (Gibco, Cat. No. 11965-092) or another suitable medium for culturing.

- Fetal bovine serum (CellMax, Cat. No. SA211.02)
- Hygromycin B (Invitrogen, Cat. No. 10687010)
- 0.25% Trypsin-EDTA (1X), Phenol Red (Gibco, Cat. No. 25200-056)
- Penicillin-Streptomycin (Gibco, Cat. No. 15140-122)
- Phosphate Buffered Saline (1X) (HyClone, Cat. No. SH30256.01)
- Complete Growth Medium: DMEM + 10% FBS, 1%P/S
- Culture Medium: DMEM + 10% FBS, Hygromycin B (20 µg/mL), 1%P/S
- Freeze Medium: 90% FBS, 10% (V/V) DMSO
- T-75 Culture flask (Corning, Cat. No. 430641)
- Cryogenic storage vials (SARSTEDT, Cat. No. 72.379.007)
- Thermostat water bath
- Centrifuge (Cence, Model: L550)
- Cell counter (MONWEI, Model: SmartCell200A Plus)
- CO₂ Incubator (Thermo, Model: 3111)
- Biological Safety Cabinet (Thermo, Model: 1389)

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• *Recovery*

1. Thaw the vial by gentle agitation in a 37°C water bath. To reduce the possibility of contamination, keep the cap out of the water. Thawing should be rapid (approximately 2 minutes).
2. Remove the vial from the water bath as soon as the contents are thawed, and decontaminate by spraying with 70% ethanol. All the operations from this point on should be carried out under strict aseptic conditions.
3. Transfer the vial contents to a centrifuge tube containing 4.0 mL complete growth medium and spin at approximately 1000 rpm for 5 minutes.
4. Resuspend cell pellet with 5 mL **complete growth medium** and transfer the cell suspension into T-75 flask containing 10-15 mL of pre-warmed complete growth medium.
5. Incubate at 37°C with 5% CO₂ incubator until the cells are ready to be split.

• *Subculture*

1. Remove and discard culture medium.
2. Wash the cells once with sterile PBS.
3. Add 2 mL of 0.25% trypsin to cell culture flask. Place the flask at 37°C for 2-3 minutes, until 90% of the cells have detached.
4. Add 6.0 to 8.0 mL of **culture medium** and aspirate cells by gently pipetting.
5. Add appropriate aliquots of the cell suspension to new culture vessel.
6. Incubate at 37°C with 5% CO₂ incubator.

Subcultivation Ratio: A subcultivation ratio of 1:6 to 1:10 is recommended.

Medium Renewal: Every 2 to 3 days.

Note: After recovery for 1-2 generations with the complete growth medium not containing the selection marker, if the cell state is well, changing to the culture medium containing the selection marker.

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• *Cryopreservation*

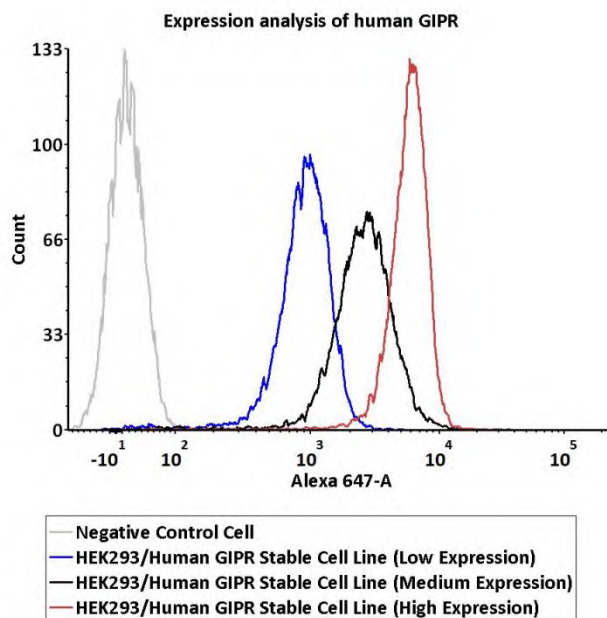
1. Remove and discard spent medium.
2. Detach cells from the cell culture flasks with 0.25% trypsin.
3. Centrifuge at 1000 rpm for 5 min at RT to pellet cells.
4. Resuspend the cell pellets with complete growth medium and count viable cells.
5. Centrifuge at 1000 rpm for 5 min at RT and resuspend cells in freezing medium to a concentration of 5×10^6 to 1×10^7 cells/mL.
6. Aliquot into cryogenic storage vials. Place vials in a programmable cooler or an insulated box placed in a -80°C freezer overnight, then transferring to liquid nitrogen storage.

• *Storage*

- **Product format:** Frozen
- **Storage conditions:** Liquid nitrogen immediately upon receipt

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• *Receptor Assay*



| Catalog No. | Stable Cell Line | MFI for GIPR (Alexa Fluor® 647) |
|--------------------|---|------------------------------------|
| CHEK-ATP208 | HEK293/Human GIPR Stable Cell Line (Low Expression) | 897.15 |
| CHEK-ATP207 | HEK293/Human GIPR Stable Cell Line (Medium Expression) | 2495.80 |
| CHEK-ATP206 | HEK293/Human GIPR Stable Cell Line (High Expression) | 5716.90 |

Fig1. Expression analysis of human GIPR on HEK293/Human GIPR Stable Cell Line by FACS. Cell surface staining using Alexa Fluor® 647-labeled anti-human GIPR antibody was performed on HEK293/Human GIPR Stable Cell Line with different expression levels: HEK293/Human GIPR Stable Cell Line (Low Expression); HEK293/Human GIPR Stable Cell Line (Medium Expression); HEK293/Human GIPR Stable Cell Line (High Expression).

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Products

HEK293/Human GIPR Stable Cell Line (Medium Expression)
 HEK293/Human GIPR Stable Cell Line (Low Expression)
 HEK293/Human GLP-1R Stable Cell Line (High Expression)
 HEK293/Human GLP-1R Stable Cell Line (Medium Expression)
 HEK293/Human GLP-1R Stable Cell Line (Low Expression)
 HEK293/Human ASGR1 Stable Cell Line
 Human GLP-1R (Luc) HEK293 Reporter Cell
 Human GCGR (Luc) HEK293 Reporter Cell
 Human GIPR (Luc) HEK293 Reporter Cell
 Human FGF-21 (Luc) HEK293 Reporter Cell
 Human Activin RII (Luc) HEK293 Reporter Cell
 HEK293/Human ASGR1&ASGR2 Stable Cell Line
 HEK293/Human GPR75 Stable Cell Line
 Human THRB (Luc) HEK293 Reporter Cell
 Human THRA (Luc) HEK293 Reporter Cell
 HEK293/Human GLP-1R&GIPR Stable Cell Line

Cat.No.

CHEK-ATP207
 CHEK-ATP208
 CHEK-ATP160
 CHEK-ATP161
 CHEK-ATP162
 CHEK-ATP080
 CHEK-ATF096
 CHEK-ATF103
 CHEK-ATF104
 CHEK-ATF163
 CHEK-ATF164
 CHEK-ATP172
 CHEK-ATP174
 CHEK-ATF181
 CHEK-ATF180
 CHEK-ATP205