

Human IL-7 R alpha/CD132 (Luc) HEK293 Reporter Cell

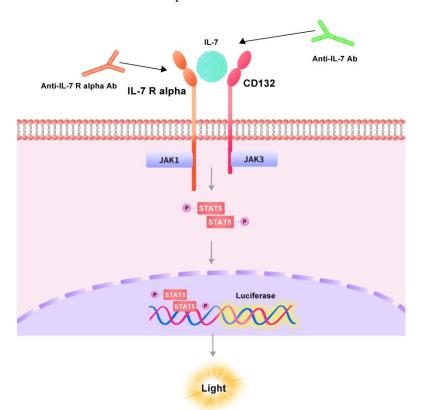
Catalog No.	Size
CHEK-ATF099	$2 \times (1 \text{ vial contains} \sim 5 \times 10^6 \text{ cells})$

• Description

The Human IL-7 R alpha/CD132 (Luc) HEK293 Reporter Cell was engineered to not only express STAT5 signaling response element, but also express the receptors full length human IL-7 R alpha (Gene ID: 3575) and CD132 (Gene ID: 3561). When stimulated with human IL-7 protein, the IL-7 R alpha/CD132 interaction drives STAT5-mediated luminescence. Neutralization of biological effect of human IL-7 protein by corresponding antibody results in a decrease in luminescence.

• Application

- Screen for neutralizing antibodies blocking the stimulation of human IL-7 protein.
- Bioactivity detection of human IL-7 fusion protein



Human IL-7 R alpha/CD132 (Luc) HEK293 Reporter Cell



• Cell Line Profile

Cell line	Human IL-7 R alpha/CD132 (Luc) HEK293 Reporter Cell
Host Cell	HEK293
Property	Adherent
Complete Growth Medium	DMEM + 10% FBS
Selection Marker	Puromycin (2 μg/mL) + Hygromycin (40 μg/mL) + G418 (200 μg/mL)
Incubation	37°C with 5% CO ₂
Doubling Time	22-24 hours
Transduction Technique	Lentivirus

• Materials Required for Cell Culture

- DMEM medium (Gibco, Cat.No.11965-092)
- Fetal bovine serum (CellMax, Cat.No.SA211.02)
- Puromycin (InvivoGen, Cat.No.ant-pr-5b)
- Hygromycin B (Invitrogen, Cat.No.10687010)
- G418 (Invivogen, Cat.No.ant-gn-5)
- Complete Growth Medium: DMEM + 10% FBS
- Culture Medium: DMEM + 10% FBS, Puromycin (2 μg/mL), Hygromycin (40 μg/mL), G418 (200 μg/mL)
- Freeze Medium: 90% FBS, 10% (V/V) DMSO
- T-75 Culture flask (Corning, 430641)
- Cryogenic storage vials (SARSTEDT, 72.379.007)
- Thermostat water bath
- Centrifuge
- Luna cell counter (Logos Biosystems, LUNA-II)
- CO₂ Incubator (Thermo, 3111)
- Biological Safety Cabinet (Thermo, 1389)



• 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.



• 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



• Receptor Assay

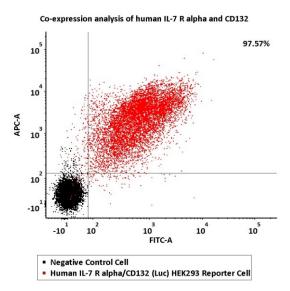


Fig1. Co-expression analysis of human IL-7 R alpha and CD132 on Human IL-7 R alpha/CD132 (Luc) HEK293 Reporter Cell by FACS. Cell surface staining was performed on Human IL-7 R alpha/CD132 (Luc) HEK293 Reporter Cell or negative control cell using FITC-labeled anti- IL-7 R alpha antibody and APC-labeled anti-CD132 antibody.

• Application



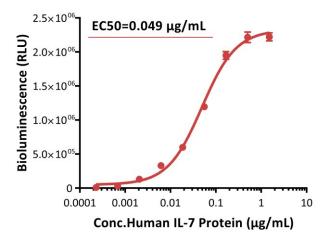


Fig2. Response to human IL-7 protein (RLU). The Human IL-7 R alpha/CD132 (Luc) HEK293 Reporter Cell was stimulated with serial dilutions of human IL-7 protein (Cat.No.IL7-H4219). The EC50 was approximately 0.049 μg/mL.



Human IL-7 Protein Stimulation (FOLD)

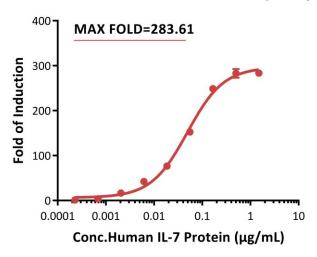


Fig3. Response to human IL-7 protein (FOLD). The Human IL-7 R alpha/CD132 (Luc) HEK293 Reporter Cell was stimulated with serial dilutions of human IL-7 protein (Cat.No.IL7-H4219). The max induction fold was approximately 283.61.



• Passage Stability

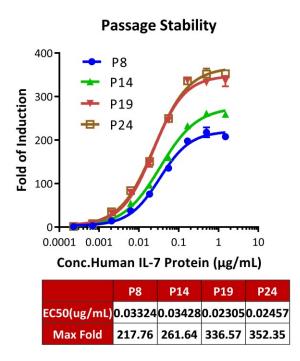


Fig4. Passage stability analysis by Signaling Bioassay. The continuously growing Human IL-7 R alpha/CD132 (Luc) HEK293 Reporter Cell was stimulated with serial dilutions of human IL-7 protein. Human IL-7 protein stimulated response demonstrates passage stabilization (fold induction and EC50) across passage 8-24.



• License Disclosure

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Related Products

<u>Products</u> <u>Cat.No.</u>

Human IL-7 Protein, premium grade IL7-H4219