

## CHO/Human uPAR Stable Cell Line

| Catalog No.  | Size   |
|--------------|--|
| SCCHO-ATP152 | $2 \times (1 \text{ vial contains } \sim 5 \times 10^6 \text{ cells})$ |

## • Description

The CHO/Human c-MET Stable Cell Line was engineered to express the receptor full length human uPAR (Gene ID: 5329), used to mimic cancer target cells. Surface expression of human uPAR was confirmed by flow cytometry.

## • Application

• Useful for cell-based uPAR binding assay

## • Cell Line Profile

| Cell line              | CHO/Human uPAR Stable Cell Line |  |
|------------------------|---------------------------------|--|
| Host Cell              | СНО                             |  |
| Property               | Adherent                        |  |
| Complete Growth Medium | F-12K + 10% FBS                 |  |
| Selection Marker       | Puromycin (2 μg/mL)             |  |
| Incubation             | 37°C with 5% CO <sub>2</sub>    |  |
| Doubling Time          | 22-24 hours                     |  |
| Transduction Technique | Lentivirus                      |  |



## • Materials Required for Cell Culture

- F-12K Nutrient Mixture (Gibco, Cat.No.21127-022)
- Fetal bovine serum (CellMax, Cat.No.SA211.02)
- Puromycin (InvivoGen, Cat.No.ant-pr-5b)
- Complete Growth Medium: F-12K + 10% FBS
- Culture Medium: F-12K + 10% FBS, Puromycin (2 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)
- CO<sub>2</sub> 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% CO<sub>2</sub> 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 3 mL of 0.25% trypsin to cell culture flask. Place the flask at 37°C for 5-7 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% CO<sub>2</sub> incubator.

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

**Medium Renewal:** Every 2 to 3 days.



### • Cryopreservation

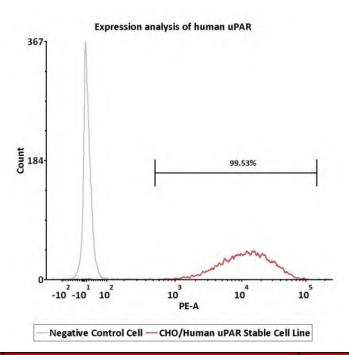
- 1. Remove and discard spent medium.
- 2. Detach cells from the cell culture flasks with 0.25% trypsin.
- 3. Centrifuge at 1000 rpm for 5 min at RT to pellet cells.
- 4. Resuspend the cell pellets with complete growth medium and count viable cells.
- 5. Centrifuge at 1000 rpm for 5 min at RT and resuspend cells in freezing medium to a concentration of  $5 \times 10^6$  to  $1 \times 10^7$  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



| Catalog No.  | Stable Cell Line                | MFI for uPAR (PE) |
|--------------|---------------------------------|-------------------|
| NA           | Negative Control Cell           | 7.08              |
| SCCHO-ATP152 | CHO/Human uPAR Stable Cell Line | 11466.08          |

**Fig1.** Expression analysis of human uPAR on CHO/Human uPAR Stable Cell Line by FACS. Cell surface staining was performed on CHO/Human uPAR Stable Cell Line or negative control cell using PE-labeled antihuman uPAR antibody.



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### • Related Products

| <u>Products</u>   | Cat.No.      |
|---|--------------|
| HEK293/Human CEACAM5 Stable Cell Line                       | CHEK-ATP083  |
| HEK293/Human DLL3 Stable Cell Line                          | CHEK-ATP090  |
| HEK293/Human ROR1 Stable Cell Line                          | CHEK-ATP084  |
| HEK293/Human TL1A Stable Cell Line                          | CHEK-ATP142  |
| HEK293/Human NAPI-IIb Stable Cell Line                      | CHEK-ATP116  |
| HEK293/Human Cadherin-6 Stable Cell Line                    | CHEK-ATP127  |
| HEK293/Human ENPP3 Stable Cell Line                         | CHEK-ATP122  |
| HEK293/Human FOLR1 Stable Cell Line                         | CHEK-ATP091  |
| HEK293/Human Glypican-3 (GPC3) Stable Cell Line             | CHEK-ATP092  |
| CHO/Human Mesothelin Stable Cell Line Development Service   | SCCHO-ATP120 |
| CHO/Human Glypican-3 (GPC3) Stable Line Development Service | SCCHO-ATP112 |
| CHO/Human c-MET Stable Cell Line Development Service        | SCCHO-ATP141 |