

# CHO/Human c-MET Stable Cell Line Development Service Data Sheet

## CHO/Human c-MET Stable Cell Line

| Catalog No.  | Size   |
|--------------|--|
| SCCHO-ATP141 | 2 × (1 vial contains ~5×10 <sup>6</sup> cells) |

### • *Description*

The CHO/Human c-MET Stable Cell Line was engineered to express the receptor full length human c-MET (Gene ID: 4233), used to mimic cancer target cells. Surface expression of human c-MET was confirmed by flow cytometry.

### • *Application*

- Useful for cell-based c-MET binding assay

### • *Cell Line Profile*

|                        |                                  |
|------------------------|----------------------------------|
| Cell line              | CHO/Human c-MET Stable Cell Line |
| Host Cell              | CHO                              |
| 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                       |

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### • *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)

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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% 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.

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## • *Cryopreservation*

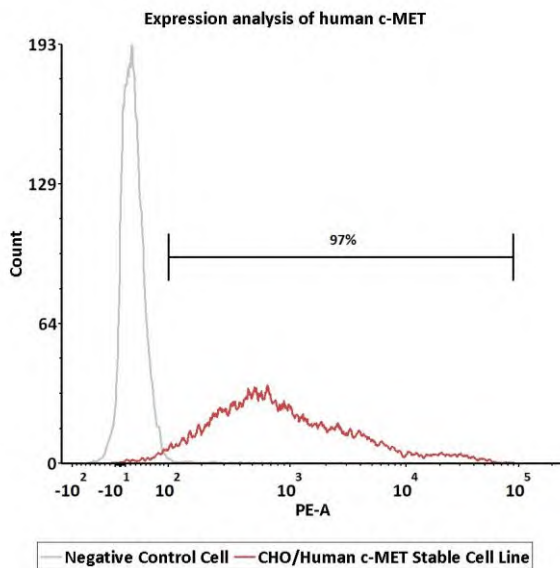
1. Remove and discard spent medium.
2. Detach cells from the cell culture flasks with 0.25% trypsin.
3. Centrifuge at 1000 rpm for 5 min at RT to pellet cells.
4. Resuspend the cell pellets with complete growth medium and count viable cells.
5. Centrifuge at 1000 rpm for 5 min at RT and resuspend cells in freezing medium to a concentration of  $5 \times 10^6$  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  $-80^\circ\text{C}$  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 c-MET (PE) |
|---------------------|---|--------------------|
| NA                  | Negative Control Cell                   | 41.62              |
| <b>SCCHO-ATP141</b> | <b>CHO/Human c-MET Stable Cell Line</b> | <b>2769.73</b>     |

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

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## • *Related Products*

### Products

### 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 uPAR Stable Cell Line Development Service         | SCCHO-ATP152 |