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# **Product Sheet**

# Cynomolgus\_SEZ6 CHO-K1 Cell Line

Catalog number: GM-C36535

Version 3.3.1.250731

Cynomolgus\_SEZ6 CHO-K1 Cell Line is a clonal stable CHO-K1 cell line that

Description constitutively expresses the cynomolgus SEZ6 gene, constructed using lentiviral

technology.

**Quantity** 5E6 Cells per vial,1 mL

**Product Format** 1 vial of frozen cells

**Shipping** Shipped on dry ice

Storage Conditions Liquid nitrogen immediately upon receipt

Target Cynomolgus\_SEZ6

Gene ID/Uniprot ID A0A2K5WPJ4

Host Cell CHO-K1

**Recovery Medium** F12K+10% FBS+1% P.S

**Growth medium** F12K+10% FBS+1% P.S+4 μg/mL Puromycin

Note None

Freezing Medium 90% FBS+10% DMSO

Growth properties Adherent

**Growth Conditions** 37°C, 5% CO<sub>2</sub>

Mycoplasma Testing The cell line has been screened to confirm the absence of Mycoplasma species.

**Safety considerations** Biosafety Level 2

Note It is recommended to expand the cell culture and store a minimum of 10 vials at an early

passage for potential future use.



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#### **Materials**

Reagent	Manufacturer/Catalogue No.
F12K	BOSTER/PYG0036
Fetal Bovine Serum	ExCell/FSP500
Pen/Strep	Thermo/15140-122
Puromycin	Genomeditech/GM-040401
Anti-SEZ6 hIgG1 Reference Antibody (ABBV-011)	Genomeditech/GM-86892MAB

## **Figures**

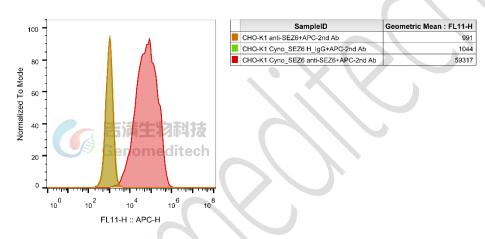


Figure 1 | Cynomolgus\_SEZ6 CHO-K1 Cell Line (Cat. GM-C36535) was determined by flow cytometry using Anti-SEZ6 hIgG1 Reference Antibody (ABBV-011) (Cat. GM-86892MAB). (This clone has undergone two rounds of single-cell separation.)

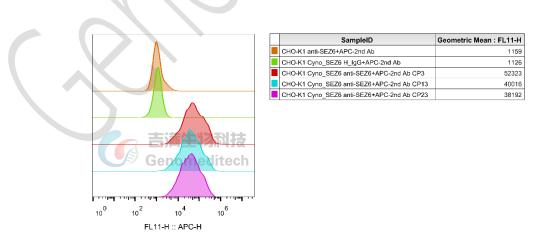


Figure 2 | The passage stability of the Cynomolgus\_SEZ6 CHO-K1 Cell Line (Cat. GM-C36535) was determined by flow cytometry using Anti-SEZ6 hIgG1 Reference Antibody (ABBV-011) (Cat. GM-86892MAB). (This clone has undergone two rounds of single-cell separation.)



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## **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°C. Storage at -70°C will result in loss of viability.

- a) Thaw the vial by gentle agitation in a 37°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).
- b) 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.
- c) Transfer the vial contents to a centrifuge tube containing 5.0 mL complete culture medium and spin at approximately 176 x g for 5 minutes. Discard supernatant.
- d) Resuspend cell pellet with the recommended recovery medium. And dispense into appropriate culture dishes.
- e) Incubate the culture at 37°C in a suitable incubator. A 5% CO<sub>2</sub> in air atmosphere is recommended if using the medium described on this product sheet.

### **Cell Freezing**

Freezing Medium: 90% FBS+10% DMSO

- a) Centrifuge at 176 x g for 3 minutes to collect cells.
- b) Resuspend the cells in pre-cooled freezing medium and adjust the cell density to 5E6 cells/mL.
- c) 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 Puromycin

For the first 1 to 2 passages post-resuscitation, use the recovery medium. Once the cells have stabilized, switch to a growth medium.

- a) Remove and discard culture medium.
- b) Briefly rinse the cell layer with PBS to remove all traces of serum that contains trypsin inhibitor.
- c) 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).
- d) 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.
- e) Add 2.0 mL of growth medium to mix well and aspirate cells by gently pipetting.
- f) After centrifugation, resuspend the pellet and add appropriate aliquots of the cell suspension to new culture vessels.
- g) Incubate cultures at 37°C.



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Subcultivation Ratio: A subcultivation ratio of 1:4 - 1:5 is recommended

Medium Renewal: Every 2 to 3 days

#### **Notes**

a) 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.

#### **Sequence**

#### SEZ6 A0A2K5WPJ4

# **Related Products**

SEZ6	
H_SEZ6 HEK-293 Cell Line	
Anti-SEZ6 hIgG1 Reference Antibody (ABBV-011)	Anti-SEZ6 hIgG1 Antibody(ABBV-011)
Biotinylated Cynomolgus SEZ6 Protein; His-Avi Tag	Cynomolgus SEZ6 Protein; His Tag
Human SEZ6 Protein; His Tag	Human SEZ6 Protein; mFc Tag
Mouse SEZ6 Protein; His Tag	
ADC Related Product	
Anti-DXD Mouse IgG1 Antibody (23E21C5)	Anti-DXD Mouse IgG1 Antibody (4A5A12)
Anti-Dxd Mouse IgG2a Antibody (17D6A4)	Anti-Eribulin Mouse IgG2a Antibody (10F8G4)
Anti-MMAE Mouse IgG1 Antibody (11C10E3)	Anti-MMAE Mouse IgG2a Antibody (17A1K11)
Anti-MMAE Mouse IgG2a Antibody (8F6A3)	Mouse anti Human IgG1-MMAE(Dar4)
Human IgG1 Isotype-DXD (Dar8)	Human IgG1 Isotype-Eribulin (Dar4)
Human IgG1 Isotype-MMAE (Dar4)	
Recombinant DT3C Protein	



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