

## 14-518ACL: GPC3 Stable Cell Line

**Application :** Functional Assay

### Description

GPC3 Stable Cell Line is a stably transfected CHO-K1 cell line which expresses human Glypican 3 (GPC3, also known as GTR2-2, OCI-5 and MXR7).

**Sequence data:** hGPC3 (accession number NP\_004475)

MAGTVRTACLVVAMLLSLDFPGQAQPPPPPDATCHQVRSFFQR  
LQPLKWWPETPVPGLQVCLPKGPTCCSRKMEEKYQLTARLNMEQLQSASMEKLF  
LIIQNAAVFQEAFAIVVRHAKNYTNAMFKNNYPSLTPQAFEFVGEFFTDVSLYLGS  
INVDDMVNELFDSLFPVIYQLMNPGLPDSALDINECLRGARRDLKVFGNFKLIMTQ  
VSKSLQVTRIFLQALNLGIEVINTTDHLKFSKDCGRMLTRMWYCSYQGLMMVKPCGG  
YCNVVMQGCMAAGVVEIDKYWREYILSLEELVNGMYRIYDMENVLLGLFSTIHDSIQYV  
QKNAGKLTTTIGKCAHSQQRQYRSAYYPEDLFIDKKVLKVAHVEHEETLSSRRRELI  
QKLKSFISFYALPGYICSHSPAENDTLCWNGQELVERYSQKAARNGMKNQFNHLEL  
KMKGPPEVVSQIIDKLKHINQLLRTMSMPKGRVLDKNLDEEGFESGDCGDEDECIGG  
SGDGMKVKQLRFLAELAYDLVDVDPAGNSQQATPKDNEISTFHNLGNVHSPLKLLT  
SMAISVVCFFFLVH

### Product Info

**Amount :** 1 Vial  
**Content :** Each vial contains  $2 \sim 3 \times 10^6$  cells in 1 ml of 90% FBS + 10% DMSO.  
**Storage condition :** Immediately upon receipt, store in liquid nitrogen.

### Application Note

#### Application:

- Screen for antibodies of human GPC3 through Flow Cytometry.

#### Culture conditions:

Cells should be grown at 37°C with 5% CO<sub>2</sub> using DMEM medium (w/ L-Glutamine, 4.5g/L Glucose and Sodium Pyruvate) supplemented with 10% heat-inactivated FBS and 1% Pen/Strep, plus 10 µg/ml of Blasticidin.

It is recommended to quickly thaw the frozen cells upon receipt or from liquid nitrogen in a 37°C water-bath, transfer to a tube containing 10 ml of growth medium without Blasticidin, spin down cells, resuspend cells in pre-warmed growth medium without Blasticidin, transfer resuspended cells to T25 flask and culture in 37°C-CO<sub>2</sub> incubator.

Leave the T25 flask in the incubator for 1~2 days without disturbing or changing the medium until cells completely recover viability and become adherent. Once cells are over 90% adherent, remove growth medium and passage the cells through trypsinization and centrifugation. At first passage, switch to growth medium containing Blasticidin. Cells should be split before they reach complete confluence.

To passage the cells, detach cells from culture vessel with Trypsin/EDTA, add complete growth medium and transfer to a tube, spin down cells, resuspend cells and seed appropriate aliquots of cells suspension into new culture vessels. Subcultivation ration = 1:10 to 1:20 weekly. To achieve satisfactory results, cells should not be passaged over 16 times.

#### LIMITED USE RESTRICTIONS:

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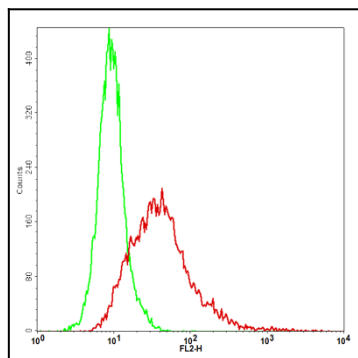


Fig-1: Detection of human GPC3 in the CHO-K1/GPC3 stable cell line by Flow Cytometry [Cell surface staining]. CHO-K1 cells (Green); CHO-K1/GPC3 cells (Red).