

## 14-522ACL: A2AR Stable Cell Line

**Application :** Functional Assay

### Description

A2AR Stable Cell Line is a stably transfected CHO-K1 cell line which expresses human adenosine A2A receptor (A2AR, also known as ADORA2A).

**Sequence data:** hA2AR (accession number AAH13780)

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MPIMGSSVYITVELAIAVLAILGNVLCWAVWLNSNLQNVTNYF
VVSLLAAADIAVGVLAIIPFAITISTGFCAACHGCLFIACFVLVLTQSSIFSLLAIAIDR
YIAIRIPLRYNGLVTGTRAKGIIAICWVLSFAIGLTPMLGWNNCGQPKEGKNHSQCGG
EGQVACLFEDEVVPMNYMVYFNFFACVLVPLLLMLGVYLRIFLAARRQLKQMESQPLPG
ERARSTLQKEVHAAKSLAIIVGLFALCWLPLHIINCFTFFCPDCSHAPLWLMYLAIVL
SHTNSVVNPFYAYRIREFRQTRFRKIIRSHVLRQQEPFKAAGTSARVLAAHGSDGEQV
SLRLNGHPPGVWANGSAPHPERRPNGYALGLVSGGSAQESQNTGLPDVELLSHELKG
VCPEPPGLDDPLAQDGAGVS
```

### Product Info

**Amount :** 1 Vial  
**Content :** Each vial contains 2 ~ 3 x 10<sup>6</sup> 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 A2AR 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.

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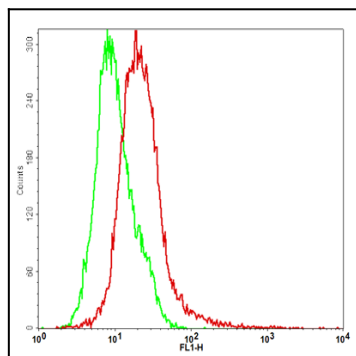


Fig-1: Detection of human A2AR in the CHO-K1/A2AR stable cell line . CHO-K1 cells (Green); CHO-K1/A2AR (Red).