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# **DATASHEET**

CA200634 CellAura fluorescent adenosine antagonist [XAC]

### **Product overview**

Name CA200634 CellAura fluorescent adenosine antagonist [XAC]

Cat No HB7814

**Biological description** Competitive fluorescent adenosine receptor antagonist (apparent K<sub>D</sub> values are 7.50, 7.37 and 7.30 for

 $A_{2A}$ ,  $A_3$  and  $A_1$  respectively). Antagonizes the activity of NECA, an adenosine receptor agonist. Inhibits cAMP accumulation and stimulates inositol phosphate accumulation (pK<sub>b</sub> values are 6.4 and 6.5

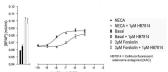
respectively). Exhibits no intrinsic agonist activity.

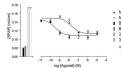
Alternative names Fluorescent Adenosine receptor Antagonist (A-633-AN), A-633-AN, XAC-X-BY630

**Biological action Purity**Antagonist
>97%

**Description** Competitive fluorescent adenosine receptor antagonist

### **Images**











## **Biological Data**

**Application notes** 

For ligand binding; fluorescence imaging; high content analysis; kinetic analysis; cell sorting at adenosine  $A_1 / A_{2A} / A_3$  receptors use solutions up to 100 nM.

Pharmacological validation

The CellAura fluorescent adenosine antagonist [XAC] ligand was shown to antagonize the activity of the adenosine receptor agonist adenosine-5'-N-ethyluronamide (NECA), in three separate recombinant CHO cell lines expressing the human  $A_1$ ,  $A_{2A}$  or  $A_3$  receptor and a cyclic AMP-responsive secreted placental alkaline phosphatase (SPAP) reporter gene. The cyclic AMP-induced expression of SPAP was measured under basal and forskolin-stimulated (maximal) conditions. Addition of CellAura fluorescent adenosine antagonist [XAC] to the basal or forskolin-stimulated cells did not significantly alter basal and stimulated SPAP levels, demonstrating that CellAura fluorescent adenosine antagonist [XAC] has no intrinsic agonist activity. To determine the apparent KD for CellAura fluorescent adenosine antagonist [XAC], cells were treated with varying concentrations of NECA alone, or in the presence of 1 $\mu$ M CellAura fluorescent adenosine antagonist [XAC], and the cyclic AMP-induced expression of SPAP measured. The apparent KD at  $A_1$ ,  $A_{2A}$  and  $A_3$  receptors was calculated from the rightward shift of the agonist response curve in the presence of CellAura fluorescent adenosine antagonist [XAC], compared to the response curve for the agonist alone, for each receptor-expressing cell line.

## **Solubility & Handling**

Storage instructions

-20°C (protect from light)

Solubility overview Soluble in DMSO

Handling After thawing individual aliquots for use, we recommend briefly sonicating the sample to ensure it is

fully dissolved and the solution is homogeneous. We do not recommend using the product after

subjecting it to repetitive freeze-thaw cycles.

Shipping conditions The product, supplied in a dry form, is stable at ambient temperature for periods of up to a few days

and does not require shipping on ice/dry ice.

Important This product is for RESEARCH USE ONLY and is not intended for therapeutic or diagnostic use. Not

for human or veterinary use.

#### **Chemical Data**

Molecular Weight974SourceSyntheticAppearancePurple solidFormulationLyophilized filmExcitation636 nmEmission651 nm