## Inhibitors



## KI-7

Cat. No.: HY-131032 CAS No.: 1489263-00-4 Molecular Formula:  $C_{23}H_{18}N_{2}O_{2}$ Molecular Weight: 354.4

Target: Adenosine Receptor Pathway: GPCR/G Protein

Storage: Powder -20°C 3 years

> 2 years In solvent -80°C 6 months

-20°C 1 month

**Product** Data Sheet

## **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 250 mg/mL (705.42 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.8217 mL	14.1084 mL	28.2167 mL
	5 mM	0.5643 mL	2.8217 mL	5.6433 mL
	10 mM	0.2822 mL	1.4108 mL	2.8217 mL

Please refer to the solubility information to select the appropriate solvent.

In Vivo

1. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (5.87 mM); Clear solution

## **BIOLOGICAL ACTIVITY**

Description

KI-7 is an A2B adenosine receptor positive allosteric modulator. KI-7 potentiates the cAMP accumulation induced by the  $non-selective~A2B~adenosine~receptor~agonist~NECA~(EC_{50}=445.8~nM).~KI-7~also~potentiates~the~cAMP~accumulation~induced$ by the selective A2B adenosine receptor agonist BAY 60-6583 as well as by adenosine with EC<sub>50</sub>s of 2390 nM and 2550 nM, respectively<sup>[1][2]</sup>.

In Vitro

KI-7 (1 μM; 5-21 days; mesenchymal stem cells) induces a significant increase in mRNA expression of Runx2 and Osterix<sup>[1]</sup>. KI-7 (1  $\mu$ M; 15-21 days) induced a significant increase in cell viability in both differentiation stages<sup>[1]</sup>. KI-7, as A2B adenosine receptor positive allosteric modulator in MSCs, demonstrating it is able to potentiate the effects of

either adenosine and synthetic orthosteric A2B adenosine receptor agonists in mediating osteoblast differentiation in vitro. NECA, BAY 60-6583 and KI-7 induce a strong increase in IL-6 production. KI-7 is able to potentiate the effects of orthosteric agonists in both differentiation stages, even if the effect became significant only at 21 days [1].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Caution: Product has not been fully validated for medical applications. For research use only.  Tel: 609-228-6898 Fax: 609-228-5909 E-mail: tech@MedChemExpress.com  Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA	ncavelli ML, et al. O	losteric modulators of human A2B adenosine is steoblast differentiation and survival: A role for		nodulators. Biochim Biophys Acta. 2014;1843(12)	2957-
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REFERENCES

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