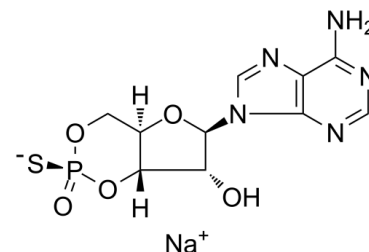


## Data Sheet

<b>Product Name:</b>	Sp-cAMPS (sodium salt)
<b>Cat. No.:</b>	CS-0107086
<b>CAS No.:</b>	142439-95-0
<b>Molecular Formula:</b>	C <sub>10</sub> H <sub>11</sub> N <sub>5</sub> NaO <sub>5</sub> PS
<b>Molecular Weight:</b>	367.25
<b>Target:</b>	Phosphodiesterase (PDE); PKA
<b>Pathway:</b>	Metabolic Enzyme/Protease; Stem Cell/Wnt
<b>Solubility:</b>	DMSO : 100 mg/mL (272.29 mM; Need ultrasonic)



### BIOLOGICAL ACTIVITY:

Sp-cAMPS sodium salt, a cAMP analog, is potent activator of cAMP-dependent **PKA I** and **PKA II**. Sp-cAMPS sodium salt is also a potent, competitive **phosphodiesterase (PDE3A)** inhibitor with a **K<sub>i</sub>** of 47.6  $\mu$ M. Sp-cAMPS sodium salt binds the **PDE10 GAF domain** with an **EC<sub>50</sub>** of 40  $\mu$ M<sup>[1][2][3]</sup>. **In Vitro:** Treatment of hepatocytes with Sp-cAMPS, the stimulatory diastereomer of adenosine cyclic 3',5'-phosphorothioate, mimics the response seen with glucagon. The glucagon-stimulated increases in the level of Ca<sup>2+</sup> can be mimicked by Sp-cAMPS<sup>[4]</sup>. **In Vivo:** In chronic alcohol consumption (CAC) mice, direct infusion of the Sp-cAMPS (1  $\mu$ g/ $\mu$ L) into the prefrontal cortex significantly improves or impairs, respectively, working memory performance in withdrawn and water animals<sup>[5]</sup>.

### References:

- [1]. Su H Hung, et al. A new nonhydrolyzable reactive cAMP analog, (Sp)-adenosine-3',5'-cyclic-S-(4-bromo-2,3-dioxobutyl)monophosphorothioate irreversibly inactivates human platelet cGMP-inhibited cAMP phosphodiesterase. *Bioorg Chem.* 2002 Feb;30(1):16-31.
- [2]. L Y Wang, et al. Regulation of kainate receptors by cAMP-dependent protein kinase and phosphatases. *Science.* 1991 Sep 6;253(5024):1132-5.
- [3]. Ronald Jäger, et al. Activation of PDE10 and PDE11 phosphodiesterases. *J Biol Chem.* 2012 Jan 6;287(2):1210-9.
- [4]. P A Connelly, et al. A study of the mechanism of glucagon-induced protein phosphorylation in isolated rat hepatocytes using (Sp)-cAMPS and (Rp)-cAMPS, the stimulatory and inhibitory diastereomers of adenosine cyclic 3',5'-phosphorothioate. *J Biol Chem.* 1987 Mar 25;262(9):4324-32.
- [5]. G Dominguez, et al. Rescuing prefrontal cAMP-CREB pathway reverses working memory deficits during withdrawal from prolonged alcohol exposure. *Brain Struct Funct.* 2016 Mar;221(2):865-77.

### CAIndexNames:

Adenosine, cyclic 3',5'-[hydrogen [P(S)]-phosphorothioate], sodium salt (1:1)

### SMILES:

O[C@H]1[C@@H](O[C@@]2([H])[C@@]1([H])O[P@](OC2)([S])=O)N3C4=C(C(N)=NC=N4)N=C3.[Na+].[-]

**Caution: Product has not been fully validated for medical applications. For research use only.**

Tel: 610-426-3128

Fax: 888-484-5008

E-mail: sales@ChemScene.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA