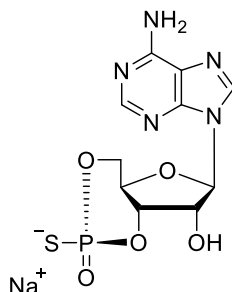


Technical Information about Sp-cAMPS

PDE-resistant activator of cAMP-dependent protein kinases type I and II

Update: June 20, 2017 HU



Abbreviation: Sp-cAMPS

Formula	CAS No.	Molecular Weight	UV	BIOLOG Cat. No.
C ₁₀ H ₁₁ N ₅ O ₅ PS·Na	[71774-13-5]	367.3	λ _{max} 259 nm / ε 15200 / pH 7	A 003 S

Name: Adenosine- 3', 5'- cyclic monophosphorothioate, Sp- isomer (Sp-cAMPS)

Description: Sp-cAMPS is an analogue of the natural signal molecule cyclic AMP in which the axial one of the two exocyclic oxygen atoms in the cyclic phosphate moiety is replaced by sulfur. The suffix "p" indicates that R/S nomenclature refers to phosphorus.

Properties:

- activator of cyclic AMP-dependent protein kinase I and II
- very high metabolic stability towards mammalian cyclic nucleotide- responsive phosphodiesterases
- membrane permeability comparable to 8-bromo-cyclic AMP
- minimal disturbance of cellular functions due to only minor structural differences to natural cyclic AMP

Specification: Crystallized or lyophilized sodium salt. Equal concentrations of Sp-cAMPS can appear very different in volume due to sensitivity of the lyophilized form to humidity. The compound can even contract to small volume droplets. Normally the product is located in the conical bottom of the tube. Micromolar quantities are determined by UV at λ_{max}. The triethyl ammonium salt of Sp-cAMPS is offered as well (Cat. No. A 003 T).

Purity: Typical analysis is > 99% (High Purity Grade) by HPLC / UV / 258 nm. The product is not sterile and has not been tested for endotoxins. BIOLOG's Sp-cAMPS is strictly checked for absence of the inhibitory Rp-cAMPS or cyclic AMP.

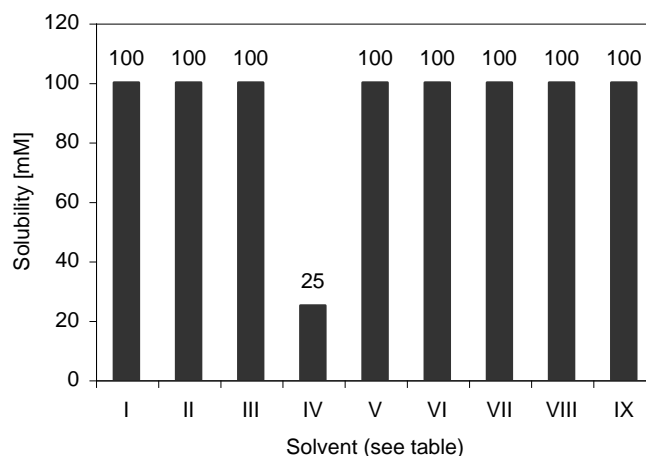
Stability and Storage: Sp-cAMPS has sufficient stability at room temperature and does not need special care during handling or shipment. Nevertheless, we recommend that the compound should be stored in the freezer, for longer storage periods preferably in freeze-dried form, since cyclic AMP can be formed by oxidation processes.

Toxicity and Safety: Since cyclic AMP has multiple tasks in every organism, it is possible that cAMP analogues will interfere with many cell regulation processes *in vivo*. However, due to the rather small quantities to work with no health hazards have been reported. Nevertheless please keep in mind that the *in vivo* properties of this compound are not sufficiently characterized up to now. Avoid skin contact or ingestion and allow only trained personnel to handle the product. Our products are designed, developed and sold for research purposes only. They are intended for *in vitro* and nonhuman *in vivo* laboratory applications. Any other use requires approval of health authorities.

Not for drug, household or related uses!

Solubility: Detailed information on the solubility of Sp-cAMPS in water and various buffers are listed in the solubility chart below. Concentrations have been tested at ambient temperatures and can be considered as minimum concentrations obtainable. When opening the tube please make sure that no substance is lost within the cap. Please rinse tube walls carefully and preferably use ultrasonic or vortex to achieve total and uniform mixing.

No.	Solvent	Solubility [mM]
I	H ₂ O	100
II	DMSO	100
III	DMF	100
IV	Ethanol 96%	25
V	Methanol	100
VI	PBS, pH 7.4	100
VII	100 mM Na ₂ HPO ₄ , pH 7.0	100
VIII	25 mM Hepes/NaOH, pH 7.2	100
IX	25 mM Tris/HCl, pH 7.4	100



Selected References for Sp-cAMPS:

Since its first synthesis by F. Eckstein, Göttingen/Germany, there have been several hundred papers published with Sp-cAMPS, and it is impossible to list them all. However, since we were among the first to offer this structure commercially, we have quite a lot of data and experience with it. Please ask for a search in our data base for articles relevant for your field. For an extended reference list please refer to our website <http://www.biolog.de>.

Eckstein, F.; Eimerl, S.; Schramm, M., *FEBS Lett.*, **64**, 92 - 94 (1976): "Adenosine 3', 5' Cyclic Phosphorothioate: An Effective Inducer of Amylase Secretion in Rat Parotid Slices"

Jarvest, R.L.; Lowe, G.; Baraniak, J.; Stec, W.J., *Biochem. J.*, **203**, 461 - 470 (1982): "A Stereochemical Investigation of the Hydrolysis of Cyclic AMP and the (Sp)- and (Rp)- Diastereomers of Adenosine Cyclic Phosphorothioate by Bovine Heart and Bakers Yeast Cyclic Phosphodiesterases"

Coulson, R.; Baraniak, J.; Stec, W.J.; Jastorff, B., *Life Sci.*, **32**, 1489 - 1498 (1983): "Transport and Metabolism of N6- and C8-Substituted Analogs of Adenosine 3',5'-Cyclic Monophosphate and Adenosine 3',5'-Cyclic Phosphorothioate by the Isolated Perfused Rat Kidney"

Scholübbbers, H.-G.; Van Knippenberg, P.H.; Baraniak, J.; Stec, W.J.; Morr, M.; Jastorff, B.; *Eur. J. Biochem.*, **138**, 101 - 109 (1984): "Investigations on Stimulation of Lac Transcription in Vivo in Escherichia Coli by cAMP Analogs. Biological Activities and Structure-Activity Correlations"

Scheinman, S.J.; Stec, W.J.; Coulson, R., *Miner. Electrolyte Metab.*, **11**, 85 - 90 (1985): "Effects of (Sp)- and (Rp)-Adenosine Cyclic 3',5' Phosphorothioates on Electrolyte Excretion by the Isolated Perfused Rat Kidney"

Hofmann, F.; Gensheimer, H.P.; Landgraf, W.; Hullin, R.; Jastorff, B., *Eur. J. Biochem.*, **150**, 85 - 88 (1985): "Diastereomers of Adenosine 3', 5'- Monothionophosphate (cAMP(S)) Antagonize the Activation of cGMP-Dependent Protein Kinase"

Braumann, T.; Jastorff, B., *J. Chromatogr.*, **350**, 105 - 118 (1985): "Physicochemical Characterization of Cyclic Nucleotides by Reversed Phase High-Performance Liquid Chromatography. II. Quantitative Determination of Hydrophobicity"

Braumann, T.; Jastorff, B.; Richter-Landsberg, C., *J. Neurochem.*, **47**, 912 - 919 (1986): "Fate of Cyclic Nucleotides in PC12 Cell Cultures: Uptake, Metabolism, and Effects of Metabolites on Nerve Growth Factor-Induced Neurite Outgrowth"

Erneux, C.; Van Sande, J.; Jastorff, B.; Dumont, J.E., *Biochem. J.*, **234**, 193 - 197 (1986): "Modulation of Cyclic AMP Action in the Dog Thyroid by its Agonist and Antagonist Sp-and Rp- Adenosine-3',5'-Monophosphorothioate"

Erneux, C.; Miot, F., *Methods Enzymol.*, **159**, 520 - 530 (1988): "Cyclic Nucleotide Analogs Used to Study Phosphodiesterase Catalytic and Allosteric Sites"

Rothermel, J.D.; Botelho, L.H.P., *Biochem. J.*, **251**, 757 - 762 (1988): "A Mechanistic and Kinetic Analysis of the Interactions of the Diastereoisomers of Adenosine 3', 5'- (Cyclic) phosphorothioate with Purified Cyclic AMP-Dependent Protein Kinase"

Dostmann, W.R.G.; Taylor, Susan S.; Genieser, H.-G.; Jastorff, B.; Døskeland, S.O.; Øgreid, D., *J. Biol. Chem.*, **265**, 10484 - 10491 (1990): "Probing the Cyclic Nucleotide Binding Sites of cAMP-Dependent Protein Kinase I and II with Analogs of Adenosine 3', 5'-Cyclic Phosphorothioates"

Van Lookeren Campagne, M.M.; Diaz, F.V.; Jastorff, B.; Winkler, E.; Genieser, H.-G.; Kessin, R.W., *J. Biol. Chem.*, **265**, 5847 - 5854 (1990): "Characterization of the Yeast Low Km cAMP- Phosphodiesterase With cAMP Analogues"

Van Ments-Cohen, M.; Genieser, H.- G.; Jastorff, B.; van Haastert, P.J.M.; Schaap, P., *FEMS Microbiol. Letters* **82**, 9 - 14 (1991): "Kinetics and Nucleotide Specificity of a Surface cAMP Binding Site in Dictyostelium discoideum which is not Down-regulated by cAMP"

Sjöholm, A., *FEBS Lett.*, **289**, 249 - 252 (1991): "Inhibition of Fetal Rat Pancreatic β -Cell Replication by Interleukin-1 in Vitro is Not Mediated Through Pertussis Toxin-sensitive G-Proteins, a Decrease in Cyclic AMP, or Protease Activation"

Wang, L.- Y.; Salter, M.W.; MacDonald, J.F., *Science*, **253**, 1132 - 1135 (1991): "Regulation of Kainate Receptors by cAMP-Dependent Protein Kinase and Phosphatases"

- Naß, N.; Colling, C.; Cramer, M.; Genieser, H.-G.; Butt, E.; Winkler, E.; Jaenicke, L.; Jastorff, B., *Biochem. J.*, **285**, 129 - 136 (1992): "Mapping of the Epitope/Paratope Interactions of a Monoclonal Antibody Directed Against Adenosine- 3', 5'-monophosphate"
- Hochner, B.; Kandel, E.R., *Proc. Natl. Acad. Sci. USA*, **89**, 11476 - 11480 (1992): "Modulation of a Transient K⁺ Current in the Pleural Sensory Neurons of Aplysia by Serotonin and cAMP: Implications for Spike Broadening"
- Laycock, J.F.; Hubbard, J.I.; Schwartz, J.H.; Stanton, B.A.; Valtin, H., *Neurohypophysis: A Window on Brain Function* (Series: Ann. N.Y. Acad. Sci., **689**, 606 - 608 (1993)): "The cAMP Agonist Sp-cAMPS Stimulates Osmotic Water Transport Across Rat Inner Medullary Collecting Duct Cell"
- Pinsky, D.; Oz, M.; Liao, H.; Morris, S.; Brett, J.; Sciacca, R.; Karakurum, M.; van Lookeren Campagne, M.; Platt, J.; Nowygrod, R.; Koga, S.; Stern, D., *J. Clin. Invest.*, **92**, 2994 - 3002 (1993): "Restoration of the cAMP Second Messenger Pathway Enhances Cardiac Preservation for Transplantation in a Heterotopic Rat Model"
- Schaap, P.; van Ments-Cohen, M.; Soede, R.D.M.; Brandt, R.; Firtel, R.A.; Dostmann, W.; Genieser, H.-G.; Jastorff, B.; van Haastert, P.J.M., *J. Biol. Chem.*, **268**, 6323 - 6331 (1993): "Cell-permeable Non-hydrolyzable cAMP Derivatives as Tools for Analysis of Signaling Pathways Controlling Gene Regulation in Dictyostelium"
- Nguyen, P.V.; Abel, T.; Kandel, E.R., *Science* **256**, 1104 - 1107 (1994): "Requirement of a Critical Period of Transcription for Induction of a Late Phase of LTP"
- Bang, Y.J.; Pirnia, F.; Fang, W.-G.; Kang, W.K.; Sartor, O.; L. Whitesell, M.J. Ha.; Tsokos, M.; Sheahan, M.D.; Nguyen, P.; Niklinski, W.T.; Myers, C.E.; Trepel, J.B., *Proc. Natl. Acad. Sci. USA*, **91**, 5330 - 5334 (1994): "Terminal Neuroendocrine Differentiation of Human Prostate Carcinoma Cells in Response to Increased Intracellular Cyclic AMP"
- Gjertsen, B.T.; Mellgren, G.; Otten, A.; Maronde, E.; Genieser, H.-G.; Jastorff, B.; Vintermyr, O.; McKnight, G. S.; Døskeland, S.O., *J. Biol. Chem.*, **270**, 20599 - 20607 (1995): "Novel (Rp)-cAMPS Analogs as Tools for Inhibition of cAMP-Kinase in Cell Culture"
- Kramer, R.H.; Tibbs, G.R., *J. Neurosci.*, **16**, 1285 - 1293 (1996): "Antagonists of Cyclic Nucleotide-gated Channels and Molecular Mapping of their Site of Action"
- Kaji, H.; Sugimoto, T.; Kanatani, M.; Nishiyama, K.; Nasu, M.; Chihara, K., *J. Cell. Physiol.*, **172**, 55-62 (1997): "Insulin-Like Growth Factor-1 Mediates Osteoclast-Like Cell Formation Stimulated By Parathyroid Hormone"
- Brown, R.L.; Strassmeier, T.; Brady, J.D.; Karpen, J.W., *Curr. Pharmaceut. Design*, **12**, 3597 - 3613 (2006): "The Pharmacology of Cyclic Nucleotide-gated Channels: Emerging from the Darkness"
- Scott, S.-P.; Shea, P.W.; Dryer, S.E., *Biochemistry*, **46**, 9417 - 9431 (2007): "Mapping Ligand Interactions with the Hyperpolarization Activated Cyclic Nucleotide Modulated (HCN) Ion Channel Binding Domain Using a Soluble Construct"
- Jäger, R.; Russwurm, C.; Schwede, F.; Genieser, H.-G.; Koesling, D.; Russwurm, M., *J. Biol. Chem.*, **287**, 1210 - 1219 (2012): „Activation of PDE10 and PDE11 Phosphodiesterases“