Proteins

Bucladesine sodium

Cat. No.: HY-B0764 CAS No.: 16980-89-5

Molecular Formula: $C_{18}H_{23}N_5NaO_8P$

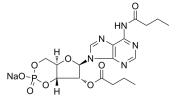
Molecular Weight: 491.37

Target: PKA; Phosphodiesterase (PDE)

Pathway: Stem Cell/Wnt; TGF-beta/Smad; Metabolic Enzyme/Protease

4°C, sealed storage, away from moisture Storage:

* In solvent: -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)



Product Data Sheet

SOLVENT & SOLUBILITY

In Vitro DMSO: 100 mg/mL (203.51 mM; Need ultrasonic)

 $H_2O : \ge 50 \text{ mg/mL} (101.76 \text{ mM})$

* "≥" means soluble, but saturation unknown.

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.0351 mL	10.1756 mL	20.3513 mL
	5 mM	0.4070 mL	2.0351 mL	4.0703 mL
	10 mM	0.2035 mL	1.0176 mL	2.0351 mL

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

In Vivo

- 1. Add each solvent one by one: PBS Solubility: 100 mg/mL (203.51 mM); Clear solution; Need ultrasonic
- 2. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 4.25 mg/mL (8.65 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 4.25 mg/mL (8.65 mM); Clear solution
- 4. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 4.25 mg/mL (8.65 mM); Clear solution

BIOLOGICAL ACTIVITY

Bucladesine sodium salt (Dibutyryl-cAMP sodium salt) is a stabilized cyclic AMP (cAMP) analog and a selective PKA activator. Description

> Bucladesine sodium salt raises the intracellular levels of cAMP. Bucladesine sodium salt is also a phosphodiesterase (PDE) $inhibitor. \ Bucladesine\ sodium\ salt\ has\ anti-inflammatory\ activity\ and\ can\ be\ used\ for\ impaired\ wound\ healing\ [1][2][3][4].$

IC₅₀ & Target

PKA

PDE

In Vitro	Both choline acetyltransferase (ChAT) and vesicular acetylcholine transporter (VAChT) mRNA increased approximately fourfold after treatment of PC12 cells with Bucladesine (dibutyryl cyclic AMP; dbcAMP). ChAT and PKA activity are also increased by Bucladesine ^[5] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	Intrahippocampal infusion of Bucladesine into the CA1 region (male Albino-Wistar rats) can cause an improvement in spatial memory in maze task. Bilateral infusion of $10~\mu M$ and $100~\mu M$ Bucladesine leads to a significant reduction in escape latency and travel distance (showing an improvement in spatial memory). Bucladesine via activation of PKA and induction of cAMP/PKA pathway improved spatial memory retention ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

CUSTOMER VALIDATION

- Cell Stem Cell. 2021 Aug 5;28(8):1362-1379.e7.
- Adv Sci (Weinh). 2021 Oct 31;e2100808.
- Cell Commun Signal. 2022 Apr 12;20(1):52.
- Ecotoxicol Environ Saf. 2022 Aug 17;243:113982.
- Int J Mol Sci. 2023 Feb 8;24(4):3398.

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REFERENCES

- [1]. Sharifzadeh, M., et al., Post-training intrahippocampal infusion of nicotine-bucladesine combination causes a synergistic enhancement effect on spatial memory retention in rats. Eur J Pharmacol, 2007. 562(3): p. 212-20.
- [2]. Mafune, E., M. Takahashi, and N. Takasugi, Effect of vehicles on percutaneous absorption of bucladesine (dibutyryl cyclic AMP) in normal and damaged rat skin. Biol Pharm Bull, 1995. 18(11): p. 1539-43.
- [3]. Rundfeldt, C., et al., The stable cyclic adenosine monophosphate analogue, dibutyryl cyclo-adenosine monophosphate (bucladesine), is active in a model of acute skin inflammation. Arch Dermatol Res, 2012.
- [4]. Salehi F, et al. Effect of bucladesine, pentoxifylline, and H-89 as cyclic adenosine monophosphate analog, phosphodiesterase, and protein kinase A inhibitor on acute pain. Fundam Clin Pharmacol. 2017 Aug;31(4):411-419.
- [5]. Shimojo M, et al. The cholinergic gene locus is coordinately regulated by protein kinase A II in PC12 cells. J Neurochem. 1998 Sep;71(3):1118-26.

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

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