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# Product Data Sheet

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# S-Adenosyl-L-methionine (1,4-butanedisulfonate)

Cat. No.:	HY-B0617B	
CAS No.:	200393-05-1	
Molecular Formula:	$C_{19}H_{32}N_6O_{11}S_3$	H <sub>2</sub> N HO OH
Molecular Weight:	616.69	
Target:	Endogenous Metabolite; Apoptosis	0, О <sub>р</sub> ОН
Pathway:	Metabolic Enzyme/Protease; Apoptosis	HO'S O'O
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.	

<b>BIOLOGICAL ACTIV</b>		
Description	S-Adenosyl-L-methionine ( Adenosyl-L-methionine 1,4 methionine 1,4-butanedisu	S-Adenosyl methionine) 1,4-butanedisulfonate is an orally active methyl group donor. S- I-butanedisulfonate is a dietary supplement with potent antidepressant effects. S-Adenosyl-L- Ilfonate also has anti⊠proliferative, pro⊠apoptotic and anti⊠metastatic roles in cancers. S- I-butanedisulfonate has the potential for, cancer, liver disease and osteoarthritis research <sup>[1][2][3]</sup> .
In Vitro	arrest in Cal-33 and JHU-Se S-Adenosyl-L-methionine ( <sup>[4]</sup> . S-Adenosyl-L-methionine ( expression of DNMTs <sup>[5]</sup> .	300 μM, 24 or 48 h) 1,4-butanedisulfonate induces cell apoptosis, and promotes the cell cycle CC-011 cells <sup>[4]</sup> . 300 μM, 24 h) 1,4-butanedisulfonate decreases the migration of the Cal-33 and JHU-SCC-011 cells 5-40 μg/mL, 48 h) 1,4-butanedisulfonate protects the anticancer effect of 5⊠FU by regulating the y confirmed the accuracy of these methods. They are for reference only.
	Cell Line:	Cal-33 and JHU-SCC-011 cells
	Concentration:	300 μM
	Incubation Time:	24 h (Cal-33) or 48 h (HU-SCC-011)
	Result:	Showed an approximately 10% and 3% of apoptotic cells respectively.
	Cell Cycle Analysis <sup>[4]</sup>	
	Cell Line:	Cal-33 and JHU-SCC-011 cells
	Concentration:	300 μM
	Incubation Time:	24 h (Cal-33) or 48 h (HU-SCC-011)
	Result:	Decreased the expression of cyclin B1, E1 and D1 in the Cal-33 and JHU-SCC-011 cells.
In Vivo	postnatal valproic acid exp	30 mg/kg, p.o., for 3 days) 1,4-butanedisulfonate prevents ASD like behaviors induced by early posure in young mice <sup>[6]</sup> . 50 and 100 mg/kg, p.o.) 1,4-butanedisulfonate shows antiepileptic, memory-enhancing, and

Animal Model:	Valproic acid treated young mice <sup>[6]</sup>
Dosage:	30 mg/kg
Administration:	p.o., for 3 days

### **CUSTOMER VALIDATION**

- J Agric Food Chem. 2021 Jul 30.
- Biochem Pharmacol. 2023 Dec 6:219:115967.
- Int Immunopharmacol. 2021 Mar 22;95:107545.
- Epigenetics Chromatin. 2021 Dec 4;14(1):52.
- J Pharm Biomed Anal. 2024 Jan 20, 115991.

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### REFERENCES

[1]. G M Bressa. S-adenosyl-l-methionine (SAMe) as antidepressant: meta-analysis of clinical studies. Acta Neurol Scand Suppl. 1994;154:7-14.

[2]. Wadie I Najm, et al. S-adenosyl methionine (SAMe) versus celecoxib for the treatment of osteoarthritis symptoms: a double-blind cross-over trial. [ISRCTN36233495]. BMC Musculoskelet Disord. 2004 Feb 26;5:6.

[3]. Shelly C Lu, et al. S-adenosylmethionine in liver health, injury, and cancer. Physiol Rev. 2012 Oct;92(4):1515-42.

[4]. Mosca L, et al. Effects of S@adenosyl@L@methionine on the invasion and migration of head and neck squamous cancer cells and analysis of the underlying mechanisms. Int J Oncol. 2020 May;56(5):1212-1224.

[5]. Ham MS, et al. S-adenosyl methionine specifically protects the anticancer effect of 5-FU via DNMTs expression in human A549 lung cancer cells. Mol Clin Oncol. 2013 Mar;1(2):373-378.

[6]. Ornoy A, et al. S-adenosyl methionine prevents ASD like behaviors triggered by early postnatal valproic acid exposure in very young mice. Neurotoxicol Teratol. 2019 Jan-Feb;71:64-74.

[7]. Dhediya RM, et al. Evaluation of antiepileptic effect of S-adenosyl methionine and its role in memory impairment in pentylenetetrazole-induced kindling model in rats. Epilepsy Behav. 2016 Aug;61:153-157.

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

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