
Product Data Sheet

Product Name: Arphamenine B (hemisulfate)

Cat. No.: GC16829

Chemical Properties

Cas. No. 144110-38-3

Chemical Name α R-[(3S)-3-amino-6-[(aminoiminomethyl)amino]-2-oxohexyl]-4-hydroxybenzenepropanoic acid, hemisulfateSMILES O=C([C@@H](N)CCCNC(N)=N)[C@H](C(O)=O)CC1=CC=C(O)C=C1.OS(O)(=O)=OFormula $C_{16}H_{24}N_4O_4 \cdot 1/2H_2SO_4$

M.Wt 385.4

Solubility ≤ 0.2 mg/ml in DMSOStorage Store at $-20^{\circ}C$ General tips For obtaining a higher solubility, please warm the tube at $37^{\circ}C$ and shake it in the ultrasonic bath for a while. Stock solution can be stored below $-20^{\circ}C$ for several months.

Shipping Condition Evaluation sample solution: ship with blue ice. All other available size: ship with RT, or blue ice upon request.

Structure

Background

Arphamenine B is a specific inhibitor of aminopeptidase B first isolated from bacteria [1]. Aminopeptidase B (Ap-B) is a Zn^{2+} -dependent exopeptidase which selectively removes Arg and/or Lys residues from the N terminus of several peptide substrates.

Aminopeptidase B has been involved in processing events occurring either during its intracellular transport along the secretory pathway or at the plasma membrane level [2].

In vitro: Arphamenine B inhibited the activity of aminopeptidase enzyme with an IC_{50} value of $9.0 \mu M$ [2]. Arphamenine B strongly inhibited transport by the oligopeptide/H⁺ symporter with the EC_{50} values of 15 to $67 \mu M$. Arphamenine at concentration $100 \mu M$ acted as either ineffective or weak inhibitor of membrane-associated hydrolysis [4]. Arphamenine selectively suppressed dipeptide hydrolysis [4].

References:

[1] Umezawa H, AOYAGI T, OHUCHI S, et al. Arphamenines A and B, new inhibitors of

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

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aminopeptidase B, produced by bacteria[J]. The Journal of antibiotics, 1983, 36(11): 1572-1575.

[2] Balogh A, Cadel S, Foulon T, et al. Aminopeptidase B: a processing enzyme secreted and associated with the plasma membrane of rat pheochromocytoma (PC12) cells[J]. Journal of Cell Science, 1998, 111(2): 161-169.

[3] Sajid M, Isaac R E, Harrow I D. Purification and properties of a membrane aminopeptidase from *Ascaris suum* muscle that degrades neuropeptides AF1 and AF2[J]. Molecular and biochemical parasitology, 1997, 89(2): 225-234.

[4] Daniel H, Adibi S A. Functional separation of dipeptide transport and hydrolysis in kidney brush border membrane vesicles[J]. The FASEB journal, 1994, 8(10): 753-759.

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