



上海源叶生物科技有限公司  
Shanghai yuanye Bio-Technology Co., Ltd  
电话: 021-61312973 传真: 021-55068248  
网址: [www.shyuanye.com](http://www.shyuanye.com)  
邮箱: [shyysw@sina.com](mailto:shyysw@sina.com)

产品名称: 阿维菌素 B1a  
产品别名: Avermectin B1a ; Abamectin B1a

生物活性:				
Description	Avermectin B1a is an antiparasitic agent that paralyzes nematodes without causing hypercontraction or flaccid paralysis.			
In Vitro	<p>[<sup>3</sup>H]AVM B1a preferentially binds to synaptic membranes from several regions of rat brain. [<sup>3</sup>H]AVM B1a specific binding to intact monolayers of granule cells increases rapidly with time of incubation and reaches equilibrium after approximately 20 min at 24°C. Higher concentrations of [<sup>3</sup>H]AVM B1a leads to markedly greater nonspecific binding, 60% at 25 nM. Various AVM analogs also produce concentration-dependent inhibition of [<sup>3</sup>H]AVM B1a binding in intact cerebellar neurons. AVM B1a and moxidectin are similar in potency (IC<sub>50</sub> values, 120 and 126 nM, respectively)[3]. AVMB1a-stimulated chloride efflux from mouse brain synaptic vesicles results from the activation of GABA-insensitive chloride channels and that this action is distinct from their previously documented effects on GABA-gated chloride channels in mouse brain preparations[4].</p>			
In Vivo	<p>Bacteria are significantly inhibited when the AVM B1a concentration is higher than 83.3 mg/kg, while fungi are less impaired in soil. Soil respiration is also inhibited by high concentration AVM B1a, which differs with soil types. The half lethal dosage (LD<sub>50</sub>) of AVM B1a to soil earthworm is estimated as 4.63 mg × cm<sup>2</sup> in filter paper contact test, and as 24.13 and 17.06 mg/kg, respectively after treated 7 and 14 days in artificial soil[1]. In artificial soil, the LC50 of AVM B1a on earthworms are 24.1 mg/kg and 17.1 mg/kg, respectively, for 7 and 14 days. About 80.0% and 94.8% of the accumulated AVM B1a are eliminated respectively in two groups within 1 day after they are exposed to AVM B1a-free soil, but a trace amount of AVM B1a is found for a relative long time in earthworms[2].</p>			
Solvent&Solubility	<p><b>In Vitro:</b> DMSO : ≥ 100 mg/mL (114.54 mM) H<sub>2</sub>O : &lt; 0.1 mg/mL (insoluble) * "≥" means soluble, but saturation unknown.</p>			
		Solvent Mass Concentration	1 mg	5 mg
	Preparing	1 mM	1.1454 mL	5.7269 mL
	Stock Solutions	5 mM	0.2291 mL	1.1454 mL
		10 mM	0.1145 mL	0.5727 mL
	<p>*请根据产品在不同溶剂中的溶解度选择合适的溶剂配制储备液; 一旦配成溶液, 请分装保存, 避免反复冻融造成的产品失效。 储备液的保存方式和期限 -80°C, 6 months; -20°C, 1 month。 -80°C 储存时, 请在 6 个月内使用, -20°C 储存时, 请在 1 个月内使用。 <b>In Vivo:</b> 请根据您的实验动物和给药方式选择适当的溶解方案。以下溶解方案都请先按照 In Vitro 方式配制澄清的储备液, 再依次添加助溶剂: ——为保证实验结果的可靠性, 澄清的储备液可以根据储存条件, 适当保存; 体内实验的工作液, 建议您现用现配, 当天使用; 以下溶剂前显示的百分比是指该溶剂在您配制终溶液中的体积占比; 如在配制过程中出</p>			



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	<p>现沉淀、析出现象，可以通过加热和/或超声的方式助溶</p> <p>1.请依序添加每种溶剂： 10% DMSO→40% PEG300 →5% Tween-80 → 45% saline</p> <p>Solubility: <math>\geq 2.5</math> mg/mL (2.86 mM); Clear solution</p> <p>此方案可获得 <math>\geq 2.5</math> mg/mL (2.86 mM, 饱和度未知) 的澄清溶液。</p> <p>以 1 mL 工作液为例，取 100 <math>\mu</math>L 25.0 mg/mL 的澄清 DMSO 储备液加到 400 <math>\mu</math>L PEG300 中，混合均匀向上述体系中加入 50 <math>\mu</math>L Tween-80，混合均匀；然后继续加入 450 <math>\mu</math>L 生理盐水定容至 1 mL。</p> <p>2.请依序添加每种溶剂： 10% DMSO →90% corn oil</p> <p>Solubility: <math>\geq 2.5</math> mg/mL (2.86 mM); Clear solution</p> <p>此方案可获得 <math>\geq 2.5</math> mg/mL (2.86 mM, 饱和度未知) 的澄清溶液，此方案不适用于实验周期在半个月以上的实验。</p> <p>以 1 mL 工作液为例，取 100 <math>\mu</math>L 25.0 mg/mL 的澄清 DMSO 储备液加到 900 <math>\mu</math>L 玉米油中，混合均匀。</p>
References	<p>[1]. Sun Y, et al. [Effects of avermectin B1a on soil microorganism and earthworm (<i>Eisenia fetida</i>)]. Ying Yong Sheng Tai Xue Bao. 2005 Nov;16(11):2140-3.</p> <p>[2]. Sun Y, et al. Bioaccumulation and elimination of avermectin B1a in the earthworms (<i>Eisenia fetida</i>). Chemosphere. 2005 Jul;60(5):699-704.</p> <p>[3]. Huang J, et al. Avermectin B1a binds to high- and low-affinity sites with dual effects on the gamma-aminobutyric acid-gated chloride channel of cultured cerebellar granule neurons. J Pharmacol Exp Ther. 1997 Apr;281(1):261-6.</p> <p>[4]. Payne GT, et al. Activation of gamma-aminobutyric acid insensitive chloride channels in mouse brain synaptic vesicles by avermectin B1a. J Biochem Toxicol. 1991 Winter;6(4):283-92.</p>

源叶生物