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丹酚酸B (98%, HPLC)

产品编号	产品名称	包装
SM1015-5mg	丹酚酸 B (98%, HPLC)	5mg
SM1015-25mg	丹酚酸 B (98%, HPLC)	25mg
SM1015-100mg	丹酚酸 B (98%, HPLC)	100mg

产品简介:

▶ 化学信息:

ET ET/LEAD				
丹酚酸B				
Salvianolic acid B				
丹参酚酸B; 丹酚酸乙; 紫草酸B				
Lithospermic acid B; Danshensuan B				
丹参 <i>Salvia miltiorrhiza</i> Bunge				
苯丙素类(Phenylpropanoids)>苯丙酸类				
C ₃₆ H ₃₀ O ₁₆ 718.61 121521-90-2 98%, HPLC				
			Water: 50 mg/ml (69.58 mM; adjust pH to 3 with	
			HCl); DMSO: 25 mg/ml (34.79 mM)	
			10mg加入1.39ml DMSO,或者每7.19mg加入1ml	
DMSO,配制成10mM溶液。				

生物信息

		disease, cerebrovascular disease, and diabetic vascular complication.					
产	^E 品描述						
信	言号通路						
	靶点	MAPK	-	-	-	-	
	IC ₅₀	1 μΜ	-	-	-	-	
		Salvianolic acid B (SA-B) 1 and 10 micromol/L decrease the cell active TGF-beta1 secretion by					
	体外研究	63.3 % and 15.6 % of the control, down-regulat pro-collgen alpha1(I) mRNA expression to 77.0%					
		and 51.8% respec	tively (P<0.05). SA-B	1 and 10 micromol/L	also inhibit MAPK a	ctivity by 1 to 2 fold	
		respectively. The degradation of Salvianolic acid B is temperature dependent. It was stable at 4°C					
存		for 30 h in aqueous solution. However, decomposition of Salvianolic acid B aqueous solution					
		occurres automatically at 25°C, and is enhanced at 37, 65 and 100°C. On the other hand, Salvianolic					
		acid B is also stable at 4, 25 and 37°C for 30 h in TPA (total phenolic acids). Salvianolic acid B is					
		stable for 30 h in buffered phosphate aqueous solutions at pH 1.5, 3.0 and 5.0. With an increase of					
		pH from the neutral, the stability of Sal B decreased.					
		Salvianolic acid	B (SalB) (5 mg · kg	g-1 · h-1) significant	tly attenuates LPS-i	nduced pulmonary	
	体内研究	microcirculatory disturbance, including the increase in leukocyte adhesion and albumin leakage.					
		In addition, LPS increases pulmonary tissue wet-to-dry weight ratio and tumor necrosis factor					
ſ.		[alpha] and interleukin 8 levels in plasma and bronchoalveolar lavage fluid enhances the					
ľ		expression of E-selectin, intercellular adhesion molecule 1, myeloperoxidase, MMP-2, and MMP-9,					
		whereas it decreases the expression of AQP-1 and AQP-5 in pulmonary tissue, all of which are					
		attenuated by SalB pretreatment1. SalB administration (10 mg/kg) significantly ameliorate the					
		Aβ25-35 peptide-induced memory impairment in the passive avoidance task (P<0.05). SalB					

	treatment also reduced the number of activated microglia and astrocytes that are observed during		
the inflammatory reaction after the administration of the Aβ25-35 peptide. Moreover, S			
markedly reduce inducible nitric oxide synthase and cyclooxygenase-2 expression levels a			
thiobarbituric acid reactive substances, which are increased by the administration of the Aβ25-3			
	peptide. Furthermore, SalB administration significantly rescue the Aβ25-35 peptide-induce		
	decrease of choline acetyltransferase and brain-derived neurotrophic factor protein levels.		
临床实验	N/A		

参考文献:

- 1. Lee YW, et al. Eur J Pharmacol. 201,704(15):70-7.
- 2. Liu P, et al. Acta Pharmacol Sin. 2002,23(8):733-8.
- 3. Lin, Fang, et al. Shock. 2013,39(3):317-25.
- 4. Man Xu, et al. Natural Product Communications. 2008,3(5):669-676.

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-	说明书	1 份

保存条件:

-20℃保存,至少一年有效。固体粉末4℃保存,至少一个月有效。如果溶于非DMSO溶剂,建议分装后-80℃保存,预计6个月内 有效。

注意事项:

- > 本产品可能对人体有一定的毒害作用,请注意适当防护,以避免直接接触人体或吸入体内。
- ▶ 本产品仅限于专业人员的科学研究用,不得用于临床诊断或治疗,不得用于食品或药品,不得存放于普通住宅内。
- ▶ 为了您的安全和健康,请穿实验服并戴一次性手套操作。

使用说明:

- 1. 收到产品后请立即按照说明书推荐的条件保存。使用前可以在2,000-10,000g离心数秒,使粉末充分沉降至管底后再开盖使用。
- 2. 该产品在溶液中不稳定、建议现用现配。具体的最佳工作浓度请参考本说明书中的体外、体内研究结果或其它相关文献、或者根 据实验目的,以及所培养的特定细胞和组织,通过实验进行摸索和优化。
- 3. 不同实验动物依据体表面积的等效剂量转换表请参考如下网页: https://www.beyotime.com/support/animal-dose.htm

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