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产品名称: **20-羟基二十碳-5Z,8Z,11Z,14Z-四烯酸**  
产品别名: **20-HETE; 20-hydroxy Arachidonic Acid; 20-羟花生四烯酸**

生物活性:				
Description	20-HETE(20-hydroxy Arachidonic Acid) is a potent vasoconstrictor produced in vascular smooth muscle (VSM) cells. It depolarizes VSM by blocking the open-state probability of Ca <sup>2+</sup> -activated K <sup>+</sup> -channels. IC <sub>50</sub> Value: Target: 20-Hydroxyeicosatetraenoic acid (20-HETE) is a cytochrome P450-derived arachidonic acid metabolite that has been shown to increase smooth muscle contractions and proliferation, stimulate endothelial dysfunction and activation and promote hypertension. in vitro: Addition of 20-HETE to the bath (1-100 nM), reduced the frequency of opening of the large-conductance Ca(2+)-activated K <sup>+</sup> channel recorded using cell-attached patches on VSM [1]. In kidney, 20-HETE induces diuresis by inhibiting Na <sup>+</sup> -K <sup>+</sup> -ATPase in proximal tubules and Na <sup>+</sup> /K <sup>+</sup> /Cl <sup>-</sup> cotransporter in the thick ascending limb of Henle's loop [2]. in vivo: In Cyp4a14(-/-) mice, which display androgen-driven and 20-HETE-dependent hypertension, treatment with 20-HETE antagonist abolished remodeling of renal resistance arteries measured as media thickness (24±1 vs. 15±1µm) and M/L (0.29±0.03 vs. 0.17±0.01) [4]. The transgenic mice had overexpressed hepatic CYP4F2, high hepatic 20-HETE and fasting plasma glucose levels but normal insulin level. The GP activity was increased and the cAMP/PKA-PhK-GP pathway was activated in the transgenic mice compared with wild-type mice [5]. Clinical trial: Mechanisms of Response to Diesel Exhaust in Subjects With Asthma. Phase not specified			
IC <sub>50</sub> & Target	Human Endogenous Metabolite			
Solvent&Solubility	<b>In Vitro:</b> <b>Ethanol : 6.67 mg/mL (20.81 mM; Need ultrasonic)</b> <b>DMSO : ≥ 3.2 mg/mL (9.99 mM)</b>  * "≥" means soluble, but saturation unknown.			
		Solvent Mass Concentration	1 mg	5 mg
	Preparing	1 mM	3.1204 mL	15.6021 mL
	Stock Solutions	5 mM	0.6241 mL	3.1204 mL
		10 mM	0.3120 mL	1.5602 mL
*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 (protect from light, stored under argon)。 -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。				
References	[1]. Zou AP, Fleming JT, Falck JR, 20-HETE is an endogenous inhibitor of the large-conductance Ca(2+)-activated K <sup>+</sup> channel in renal arterioles. Am J Physiol. 1996 Jan;270(1 Pt 2):R228-37. [2]. Schwartzman M, Ferreri NR, Carroll MA, Renal cytochrome P450-related arachidonate metabolite inhibits (Na <sup>+</sup> + K <sup>+</sup> )ATPase. Nature. 1985 Apr 18-24;314(6012):620-2. [3]. Ding Y, Wu CC, Garcia V, 20-HETE INDUCES REMODELING OF RENAL RESISTANCE ARTERIES INDEPENDENT OF BLOOD PRESSURE ELEVATION IN HYPERTENSION. Am J Physiol Renal Physiol.			



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[4]. Lai G, Wu J, Liu X, 20-HETE induces hyperglycemia through the cAMP/PKA-PhK-GP pathway. Mol Endocrinol. 2012 Nov;26(11):1907-16.



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