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产品名称: INT-767
产品别名: INT-767

生物活性:						
Description		INT-767 is a dual farnesoid X receptor (FXR)/TGR5 agonist with mean EC50s of 30 and 630 nM, respectively[1][2].				
In Vitro		INT-767 does not show cytotoxic effects in HepG2 cells, does not inhibit cytochrome P450 enzymes, is highly stable to phase I and II enzymatic modifications, and does not inhibit the human ether-a-go-go-related gene potassium channel[2].				
In Vivo		INT-767 (10-20 mg/kg; i.p.; daily for 2 weeks) decreases plasma total cholesterol and triglyceride levels in db/m and db/db mice [2].				
		Animal Model:	Male 8-week old C57BKS/J db/db mice, control nondiabetic db/m mice[2]			
		Dosage:	10, 20 mg/kg			
		Administration:	Intraperitoneal injection; daily for 2 weeks			
		Result:	Decreased plasma total cholesterol and triglyceride levels.			
Solvent&Solubility		In Vitro: DMSO : ≥ 205.5 mg/mL (415.44 mM) * "≥" means soluble, but saturation unknown.				
		<div>Preparing Stock Solutions</div>	<div>Solvent / Mass / Concentration</div>	1 mg	5 mg	10 mg
			1 mM	2.0216 mL	10.1080 mL	20.2159 mL
			5 mM	0.4043 mL	2.0216 mL	4.0432 mL
			10 mM	0.2022 mL	1.0108 mL	2.0216 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℃, 6 months; -20℃, 1 month。 -80℃ 储存时, 请在 6 个月内使用, -20℃ 储存时, 请在 1 个月内使用。				
		源叶生物				
References		<p>[1]. Baghdasaryan A, et al. Dual farnesoid X receptor/TGR5 agonist INT-767 reduces liver injury in the Mdr2-/- (Abcb4-/-) mouse cholangiopathy model by promoting biliary HCO3- output. Hepatology. 2011 Oct;54(4):1303-1312.</p> <p>[2]. Rizzo G, et al. Functional characterization of the semisynthetic bile acid derivative INT-767, a dual farnesoid X receptor andTGR5 agonist. Mol Pharmacol. 2010 Oct;78(4):617-630.</p>				