For Research Use Only Bardoxolone Methyl



Catalog Number: CM04717

产品信息	Catalog Number: CM04717 CAS号: 218600-53-4 分子式: C ₃₂ H ₄₃ NO ₄ 主要靶点: Nrf2 Autophagy I × B/IKK Apoptos	分子量: 505.69 溶解度: H2O:<1 mg/mL,DMSO:20 mg/mL (39.5 mM),Ethanol:<1 mg/mL	
	主要通路: 凋亡 NF-κB信号通路 免疫与炎症 自噬	ISFEITOPLOSIS	
体外活性	Bardoxolone Methyl exhibits potent inhibit macrophages with IC50 of 0.1 nM. [1] Bardo IC50 of 0.4, 0.4, and 0.27 μ M, respectively. it blocks Bcl-2 phosphorylation, which cont constitutive and inducible NF-kappaB activ lipopolysaccharide, and cigarette smoke. [20]	xolone Methyl decreases the viability of l CDDO-Me induces pro-apoptotic Bax prote ributes to the induction of apoptosis. [2] Ba rated by TNF, interleukin (IL)-1beta, phorbo	eukemic HL-60, KG-1, and NB4 cells with in, inhibits the activation of ERK1/2, and ardoxolone Methyl potently inhibits both
体内活性	Bardoxolone Methyl (60 mg/kg) reduces th significantly reduces the in vivo inflamma the spleen, and protects mice against letha	tory cytokine response following LPS chall	s in vivo. [4] Bardoxolone Methyl enge, induces HO-1 protein expression in
细胞实验	Leukemic cell lines are cultured at a c mononuclear cells are cultured at 56 indicated concentrations of CDDO-M are included as control. For cytotoxici viable cells are counted with the tryp Reference)	thinsp;× 105 cells/m e. Appropriate amounts of DMSO (fina ity studies, 1 μM ara-C is added to	L in the presence or absence of al concentration less than 0.05%) o the cultures. After 24 to 72 hours,
描述	Bardoxolone Methyl (TP-155), an IKK inhibi	tor, exhibits effective proapoptotic and an	ti-inflammatory activities.
储存	Powder: -20°C for 3 years In solvent:	-80°C for 2 years	