DECACHEM DOO, Novi Sad, Serbia



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TECHNICAL DATA SHEET

Ethyl maltol

Cat. No.: HY-W010320
CAS No.: 4940-11-8
Molecular Formula: C₃H₈O₃
Molecular Weight: 140.14

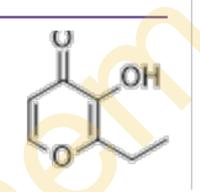
 Target:
 Heme oxygenase (HO)

 Pathway:
 Metabolic Enzyme/Protease

 Storage:
 Powder -20°C 3 years

4°C 2 years In solvent -80°C 6 months

-20°C 1 month



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SOLVENT & SOLUBILITY

In Vitro

DMSO: 100 mg/mL (713.57 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	7.1357 mL	35.6786 mL	71.3572 mL
	5 mM	1.4271 mL	7.1357 mL	14.2714 mL
	10 mM	0.7136 mL	3.5679 mL	7.1357 mL

Please refer to the solubility information to select the appropriate solvent.

In Vivo

- Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.08 mg/mL (14.84 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)
 Solubility: ≥ 2.08 mg/mL (14.84 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (14.84 mM); Clear solution

BIOLOGICAL ACTIVITY

Description	Ethyl maltol (2-Ethyl-3-hydroxy-4H-pyran-4-one) is an orally active and important food additive and flavor enhancer. Ethyl maltol is less toxic to rats and dogs. Ethyl maltol can enhance copper-mediated cytotoxicity and induce apoptosis in lung epithelial cells ^{[1][2][3][4]} .
In Vitro	Ethyl maltol (3 mM; 24 h) induces cell apoptosis and leads to oxidative stress and DNA damage in copper traeated lung epithelial cells ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.



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	Western Blot Analysis ^[1]	Western Blot Analysis ^[1]		
	Cell Line:	Cu treated Calu-6 cells		
	Concentration:	3 mM		
	Incubation Time:	24 h		
	Result:	Increased pATM phosphorylation level.		
	Real Time qPCR[1]			
	Cell Line:	Cu treated Calu-6 cells		
	Concentration:	3 mM		
	Incubation Time:	24 h		
	Result:	Increased the expression of ferritin light chain and hemeoxygenase.		
In Vivo	kidney function impairs	Ethyl maltol (oral administration; 50-1000 mg/kg; 90 days-2 years) is safe in rats and dogs, and the toxicity (such as liver and kidney function impairment, hemolysis) is less than the homolog maltol ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.		

