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Lieferung & Zahlungsart

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Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
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- Expressversand

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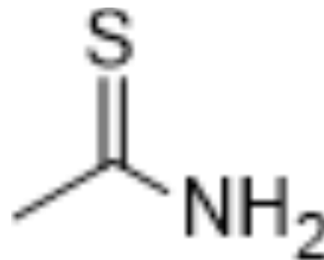
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Thioacetamide

Cat. No.:	HY-Y0698
CAS No.:	62-55-5
Molecular Formula:	C ₂ H ₅ NS
Molecular Weight:	75.13
Target:	Necroptosis
Pathway:	Apoptosis
Storage:	4°C, stored under nitrogen * In solvent : -80°C, 6 months; -20°C, 1 month (stored under nitrogen)



SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (1331.03 mM; Need ultrasonic)					
	H ₂ O : 50 mg/mL (665.51 mM; Need ultrasonic)					
	Preparing Stock Solutions	Solvent	Mass	1 mg	5 mg	10 mg
		Concentration				
		1 mM		13.3103 mL	66.5513 mL	133.1026 mL
5 mM			2.6621 mL	13.3103 mL	26.6205 mL	
10 mM		1.3310 mL	6.6551 mL	13.3103 mL		
Please refer to the solubility information to select the appropriate solvent.						
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (33.28 mM); Clear solution					
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (33.28 mM); Clear solution					
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (33.28 mM); Clear solution					

BIOLOGICAL ACTIVITY

Description	Thioacetamide (TAA) is an indirect hepatotoxin and causes parenchymal cell necrosis. Thioacetamide requires metabolic activation by microsomal CYP2E1 to thioacetamide-S-oxide initially and then to thioacetamide-S-dioxide, which is a highly reactive metabolite, and its reactive metabolites covalently bind to proteins and lipids thereby causing oxidative stress and centrilobular necrosis. Thioacetamide can induce chronic liver fibrosis, encephalopathy and other events model ^{[1][2][3][4]} .
In Vitro	Thioacetamide (TAA; 0-10000 μM; 24 h; WB-F344 cells) has cytotoxicity in a concentration-dependent manner ^[4] . Thioacetamide (TAA; 1000 and 10000 μM; 0-24 h; WB-F344 cells) has differentially-expressed genes in the early phases at low (1000 μM) and high (10000 μM) concentrations ^[4] .

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Cell Viability Assay^[4]

Cell Line:	WB-F344 cells
Concentration:	0-10000 μ M
Incubation Time:	24 hours
Result:	Had 20% and 50% cell death at the 1000 and 10000 μ M concentrations, respectively.

In Vivo

Thioacetamide (TAA; 100 mg/kg; i.p.; three times weekly) can induce chronic liver fibrosis in male ICR mice^[2].

Thioacetamide (200-1200 mg/kg; i.p.; once) induces hepatic encephalopathy model in C57BL/6 mice^[3].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Male ICR mice ^[2]
Dosage:	100 mg/kg
Administration:	Intraperitoneal injection; three times weekly for eight weeks
Result:	Induced chronic liver fibrosis in male ICR mice and resulted in lower body weight, serum cholesterol and triglycerides as well as increased liver size, ALT, AST and LDH values.

Animal Model:	Male C57BL/6 mice (20-25g, aged 8-12 weeks) ^[3]
Dosage:	200, 600, and 1,200 mg/kg
Administration:	Intraperitoneal injection; once
Result:	Altered the neuropsychiatric state, motor behavior and reflex and sensory functions. Increased in the glutamate release in the cerebral cortex of Hepatic encephalopathy (HE) mice.

REFERENCES

- [1]. Wallace MC, et, al. Standard operating procedures in experimental liver research: thioacetamide model in mice and rats. *Lab Anim.* 2015 Apr;49(1 Suppl):21-9.
- [2]. Chen IS, et, al. Hepatoprotection of silymarin against thioacetamide-induced chronic liver fibrosis. *J Sci Food Agric.* 2012 May;92(7):1441-7.
- [3]. Miranda AS, et, al. A thioacetamide-induced hepatic encephalopathy model in C57BL/6 mice: a behavioral and neurochemical study. *Arq Neuropsiquiatr.* 2010 Aug;68(4):597-602.
- [4]. Yeom HJ, et, al. Expression analysis of early response-related genes in rat liver epithelial cells exposed to thioacetamide in vitro. *J Vet Med Sci.* 2009 Jun;71(6):719-27.

Caution: Product has not been fully validated for medical applications. For research use only.

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