

Produktinformation



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Zellkultur & Verbrauchsmaterial
Diagnostik & molekulare Diagnostik
Laborgeräte & Service

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Zuschläge

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Data Sheet (Cat.No.T5523)



Imidazole ketone erastin

Chemical Propert	ies	
CAS No. :	1801530-11-9	, and the second
Formula:	C35H35ClN6O5	
Molecular Weight:	655.14	
Appearance:	no data available	
Storage:	Powder: -20°C for 3 years In solvent: -80°C for 1 year	

Biological Description Description Imidazole ketone erastin (IKE) is an ferroptosis inducer with inhibitory effects on system Xc-cystine/glutamate transporter proteins. Imidazole ketone erastin has antitumor activity and induces glutathione depletion and lipid peroxidation. Targets(IC50) Ferroptosis In vitro METHODS: Human astrocytoma cells CCF-STTG1 were incubated with Imidazole ketone erastin for 2 h. Glutamate released into the medium was detected using glutamate oxidase, horseradish peroxidase, and Amplex UltraRed fluorimetric assay. RESULTS: The IC50 for system xc- inhibition by Imidazole ketone erastin was 30 nM. [1] METHODS: 18 DLBCL cell lines were treated with Imidazole ketone erastin (0.0001-100 µM) for 24 h. Cell viability was measured using the Cell Titor-Glo luminescent cell viability test. RESULTS: DLBCL cell lines showed different sensitivities to Imidazole ketone erastin inhibition. 7 cell lines with IC50 <100 nM were categorized as sensitive; 5 cell lines with IC50 >10 µM were categorized as resistant; and 6 cell lines with IC50 between 100 nM and 10 µM were categorized as moderately resistant. [2] **METHODS**: To detect anti-tumor activity in vivo, Imidazole ketone erastin (23-40 mg/kg, In vivo 5% DMSO in HBSS at pH 4) was injected intraperitoneally into NCG mice harboring human diffuse histiocytic lymphoma SUDHL6 once daily for thirteen days. **RESULTS**: Imidazole ketone erastin inhibited tumor growth in vivo. [2] METHODS: To test the antitumor activity in vivo, Imidazole ketone erastin (40 mg/kg every two days) and liproxstatin-1 (10 mg/kg once a day) were intraperitoneally injected into C57BL/6 mice with the hepatocellular carcinoma tumor Hepa1-6 for ten days on either a control (Met+) or a methionine-free (Met-) diet. Met-) diet. **RESULTS:** In mice receiving the control diet, Imidazole ketone erastin treatment effectively reduced tumor growth, and this effect was completely blocked by liproxtatin-1. However, in mice receiving a methionine-free diet, Imidazole ketone erastin treatment failed to inhibit tumor growth. [3]

Solubility Information

A DRUG SCREENING EXPERT

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Solu	ubility	

DMSO: 18.33 mg/mL (27.98 mM),Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)

Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	1.5264 mL	7.632 mL	15.2639 mL
5 mM	0.3053 mL	1.5264 mL	3.0528 mL
10 mM	0.1526 mL	0.7632 mL	1.5264 mL
50 mM	0.0305 mL	0.1526 mL	0.3053 mL

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Reference

Liu D, Liang C, Huang B, et al.Tryptophan Metabolism Acts as a New Anti-Ferroptotic Pathway to Mediate Tumor Growth.Advanced Science.2023: 2204006.

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