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Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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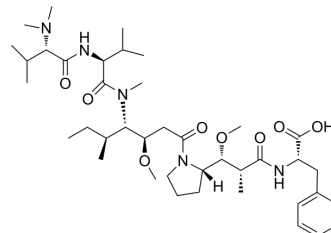
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Auristatin F

Cat. No.:	HY-15583
CAS No.:	163768-50-1
Molecular Formula:	C ₄₀ H ₆₇ N ₅ O ₈
Molecular Weight:	745.99
Target:	Microtubule/Tubulin; ADC Cytotoxin
Pathway:	Cell Cycle/DNA Damage; Cytoskeleton; Antibody-drug Conjugate/ADC Related
Storage:	Powder -20°C 3 years 4°C 2 years



* The compound is unstable in solutions, freshly prepared is recommended.

SOLVENT & SOLUBILITY

In Vitro	DMSO : 200 mg/mL (268.10 mM; Need ultrasonic)					
	Preparing Stock Solutions	Solvent	Mass	1 mg	5 mg	10 mg
		Concentration				
		1 mM		1.3405 mL	6.7025 mL	13.4050 mL
		5 mM		0.2681 mL	1.3405 mL	2.6810 mL
	10 mM		0.1341 mL	0.6703 mL	1.3405 mL	
Please refer to the solubility information to select the appropriate solvent.						
In Vivo	<ol style="list-style-type: none"> Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 5 mg/mL (6.70 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 5 mg/mL (6.70 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 5 mg/mL (6.70 mM); Clear solution 					

BIOLOGICAL ACTIVITY

Description	Auristatin F is a potent cytotoxin in antitumor-conjugated agents and an analogue of MMAF. Auristatin F is a potent microtubule inhibitor and vascular damaging agent (VDA). Auristatin F inhibits cell division by preventing tubulin aggregation. Auristatin F can be used in antibody-drug conjugates (ADC) ^{[1][2]} .
IC₅₀ & Target	Microtubule ^[1]
In Vitro	Auristatin F and Monomethyl Auristatin F (MMAF) are potent ADC cytotoxin used in antibody-drug conjugates ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

In Vivo

Auristatin F (5 mg/kg; i.v.; male Sprague-Dawley rats) has C_{max} of 8276.76 ng/mL. The AUC_{last} is 65661.30 min*ng/mL, and the clearance (CL) is 77.33 mL/min/kg, which is above the hepatic blood flow in the rat^[2].

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

Animal Model: Male Sprague-Dawley rats^[2]

Dosage: 5 mg/kg

Administration: Intravenous injection

Result:

Parameter	
Dose (i.v.) mg/kg	5
C_{max} (ng/mL)	8276.76
AUC_{last} (min*ng/mL)	62661.30
CL (mL/min/kg)	77.33
Vss (mL/Kg)	1057.13

REFERENCES

[1]. Park MH, et al. Pharmacokinetic and Metabolism Studies of Monomethyl Auristatin F via Liquid Chromatography-Quadrupole-Time-of-Flight Mass Spectrometry. *Molecules*. 2019 Jul 29;24(15):2754.

[2]. Roy S, et al. SMI-Ribosome inactivating protein conjugates selectively inhibit tumor cell growth. *Chem Commun (Camb)*. 2017 Apr 11;53(30):4234-4237.

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

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