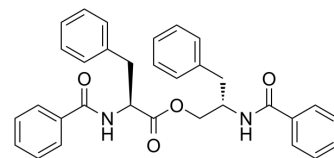


## Asperphenamate

Cat. No.:	HY-129578		
CAS No.:	63631-36-7		
Molecular Formula:	C <sub>32</sub> H <sub>30</sub> N <sub>2</sub> O <sub>4</sub>		
Molecular Weight:	506.59		
Target:	Autophagy; Cathepsin		
Pathway:	Autophagy; Metabolic Enzyme/Protease		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



### SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (197.40 mM; Need ultrasonic)				
		Solvent Concentration	Mass 1 mg	5 mg	10 mg
	Preparing Stock Solutions	1 mM	1.9740 mL	9.8699 mL	19.7398 mL
		5 mM	0.3948 mL	1.9740 mL	3.9480 mL
10 mM		0.1974 mL	0.9870 mL	1.9740 mL	
Please refer to the solubility information to select the appropriate solvent.					
In Vivo	1. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (4.93 mM); Clear solution				

### BIOLOGICAL ACTIVITY

Description	Asperphenamate, a fungal metabolite of <i>Aspergillus flatiipes</i> with anti-cancer effect, exhibits IC <sub>50</sub> values of 92.3 μM, 96.5 μM and 97.9 μM in T47D, MDA-MB-231 and HL-60 cells, respectively <sup>[1][2]</sup> .
In Vitro	Asperphenamate can inhibit cancer cell proliferation by fully inducing autophagy. asperphenamate showed inhibition effects against cathepsin L. At the same time, it also displayed weak inhibitory ability against cathepsin S. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### REFERENCES

[1]. Alice M.Clark, et al. Synthesis of asperphenamate, a novel fungal metabolite. Phytochemistry

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[2]. LeiYuan, et al. Total synthesis and anticancer activity studies of the stereoisomers of asperphenamate and patriscabratine. Chinese Chemical Letters Volume 21, Issue 2, February 2010, Pages 155-158.

[3]. Yuan L, et al. Discovery of novel cathepsin inhibitors with potent anti-metastatic effects in breast cancer cells. Bioorg Chem. 2018 Dec;81:672-680.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: [tech@MedChemExpress.com](mailto:tech@MedChemExpress.com)

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA