

Product Name: Marimastat Revision Date: 01/10/2021

Product Data Sheet

Marimastat

Cat. No.: A4049

CAS No.: 154039-60-8
Formula: C15H29N3O5

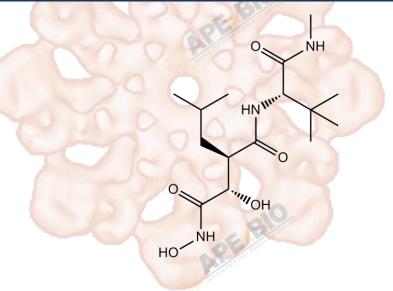
M.Wt: 331.41

Synonyms:

In Vitro

Target: Proteases
Pathway: MMP

Storage: Store at -20°C



Solvent & Solubility

≥2.8 mg/mL in H2O with gentle warming and ultrasonic; ≥20.43 mg/mL in EtOH; ≥80.1 mg/mL in DMSO

Preparing Stock Solutions	Solvent Concentration	1mg	5mg	10mg
	1 mM	3.0174 mL	15.0871 mL	30.1741 mL
	5 mM	0.6035 mL	3.0174 mL	6.0348 mL
	10 mM	0.3017 mL	1.5087 mL	3.0174 mL

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

Biological Activity

Shortsummary	MMPs inhibitor,board spe	MMPs inhibitor,board spectrum		
IC ₅₀ & Target	3 nM (MMP-9), 5 nM (MMP-1), 6 nM (MMP-2), 9 nM (MMP-14), 13 nM (MMP-7)			
	Cell Viability Assay			
In Vitro	Cell Line:	Human glioma cell lines U251 and GaMG.		
	Preparation method:	Soluble in DMSO > 10 mM. General tips for obtaining a higher concentration:		
		Please warm the tube at 37°C for 10 minutes and/or shake it in the ultrasonic		
		bath for a while. Stock solution can be stored below -20°C for several months.		

	Reacting conditions: 0.3 μM, 0.6 μM, 1.0 μM, 10 μM and 50 μM; 48 h or 6 days.			
	Applications:	In co-cultures of tumor spheroids derived from human glioma cell lines U251		
		and GaMG with RBA, marimastat strongly inhibits tumor invasion at		
		concentrations of 10 µM. Marimastat (10 µM) significantly reduces cell		
		proliferation by 54% and completely inhibits cell growth at 50 µM over 6 days.		
	610	Also, marimastat (10 μM) reduces U251 spheroid growth by 65%.		
	Animal experiment			
In Vivo	Animal models:	Male 6-week-old C57BL6/J mice with liver fibrosis.		
	Dosage form:	100 mg/kg; twice daily via orogastric gavage; one week.		
	Applications:	Marimastat significantly reduces liver injury and inflammation but induces a		
		25% increase in collagen deposition following acute CCl4-administration.		
		Marimastat inhibits MMP activities and reduces fibrolysis. In CCl4-induced		
		chronic hepatic injury, marimastat significantly reduces serum alanine		
	~1O.	aminotransferase (ALT) levels by 14-fold and downregulates the mRNA levles		
		of major pro-fibrogenic genes.		
	Other notes:	Please test the solubility of all compounds indoor, and the actual solubility may		
	The state of the s	slightly differ with the theoretical value. This is caused by an experimental		
	13-0	system error and it is normal.		

Product Citations

1. Sapoznikov A, Gal Y, et al. "Early disruption of the alveolar-capillary barrier in a ricin-induced ARDS mouse model: neutrophil-dependent and -independent impairment of junction proteins." Am J Physiol Lung Cell Mol Physiol. 2019 Jan 1;316(1):L255-L268.PMID:30382767

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References

- [1]. Tonn JC, Kerkau S, Hanke A, et al. Effect of synthetic matrix-metalloproteinase inhibitors on invasive capacity and proliferation of human malignant gliomas in vitro. Int J Cancer, 1999, 80(5): 764-772.
- [2]. de Meijer VE, Sverdlov DY, Popov Y, et al. Broad-spectrum matrix metalloproteinase inhibition curbs inflammation and liver injury but aggravates experimental liver fibrosis in mice. PLoS One, 2010, 5(6): e11256.

Caution

FOR RESEARCH PURPOSES ONLY.

NOT FOR HUMAN, VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

Specific storage and handling information for each product is indicated on the product datasheet. Most APExBIO products are stable under the recommended conditions. Products are sometimes shipped at a temperature that differs from the recommended storage temperature. Shortterm storage of many products are stable in the short-term at temperatures that differ from that required for

long-term storage. We ensure that the product is shipped under conditions that will maintain the quality of the reagents. Upon receipt of the product, follow the storage recommendations on the product data sheet.



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