

Product Name: Rocilinostat (ACY-1215) Revision Date: 01/10/2021

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## **Product Data Sheet**

# Rocilinostat (ACY-1215)

Cat. No.:	A4083
CAS No.:	1316214-52-4
Formula:	C24H27N5O3
M.Wt:	433.5
Synonyms:	
Target:	DNA Damage/DNA Repair
Pathway:	HDAC
Storage:	Store at -20°C
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## Solvent & Solubility

	$\geq$ 21.675 mg/mL in DMSO; insoluble in H2O; insoluble in EtOH				
In Vitro	Preparing Stock Solutions	Mass Solvent Concentration	1mg	5mg	10mg
		1 mM	2.3068 mL	11.5340 mL	23.0681 mL
		5 mM	0.4614 mL	2.3068 mL	4.6136 mL
		10 mM	0.2307 mL	1.1534 mL	2.3068 mL

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

## **Biological Activity**

Shortsummary	Selective HDAC6 inhibitor	
IC <sub>50</sub> & Target	5 nM (HDAC6)	
In Vitro	Cell Viability Assay	
	Cell Line:	MM cells; PBMCs.
	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:	0.16, 0.8, 4, 20 nM; MM cells: 24 h, PBMCs: 48 h.

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	Applications:	ACY-1215 induces less cytotoxicity in PHA-stimulated or unstimulated PBMCs.		
		In purified CD4+ T cells, ACY-1215 induces toxicity with IC50 value of 2.5 $\mu\text{M}.$		
		In MM cell lines, ACY-1215 dose-dependently decreases viability with IC50		
		value of 2-8 $\mu\text{M}$ and induces significant cytotoxicity. In MM.1S cells, ACY-1215		
		dose-dependently reduces DNA synthesis of MM cells adherent to BMSCs and		
	29	also inhibits growth induced by IL-6 and IGF-1.		
	Animal experiment	DE		
In Vivo	Animal models:	Male SCID mice inoculated subcutaneously with MM.1S cells.		
	Dosage form:	50 mg/kg; 5 days a week for 3 weeks; administrated orally.		
	Applications:	In human MM xenograft mouse model, ACY-1215 (50 mg/kg) significa		
		delays tumor growth. Treatment mice with ACY-1215 plus bortezomib		
		significantly prolonged overall survival (OS) and induces a significant		
		accumulation of polyubiquitinated proteins. Treatment with ACY-1215 plus		
		bortezomib is well tolerated and have no significant influence on the body		
	PEBIO	weight. In female SCID-beige mice inoculated intravenously with		
		MM.1S-LucNeo cells, treatment with ACY-1215 (75 mg/kg) and bortezomib		
	State Constant	(1.5 mg/kg) significantly inhibits tumor growth and prolongs OS.		
	Preparation method:	Dissolved in 10% DMSO in 5% dextrose in water		
	Other notes:	Please test the solubility of all compounds indoor, and the actual solubility may		
		slightly differ with the theoretical value. This is caused by an experimental		
		system error and it is normal.		

### **Product Citations**



1.Sun, Yuefeng, et al. "Fe65 Suppresses Breast Cancer Cell Migration and Invasion through Tip60 Mediated Cortactin Acetylation." Scientific reports 5 (2015).PMID:26166158

See more customer validations on www.apexbt.com.

#### References

[1]. Santo L, Hideshima T, Kung AL, et al. Preclinical activity, pharmacodynamic, and pharmacokinetic properties of a selective HDAC6 inhibitor, ACY-1215, in combination with bortezomib in multiple myeloma. Blood, 2012, 119(11): 2579-2589.

#### 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

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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.



### APExBIO Technology

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