BRM/BRG1 ATP Inhibitor-3

Cat. No.:	HY-148373	
CAS No.:	2368901-31-7	NH ₂
Molecular Formula:	$C_{26}H_{25}N_{5}O_{2}S_{2}$	
Molecular Weight:	503.64	
Target:	Epigenetic Reader Domain	O NH H
Pathway:	Epigenetics	`s N
Storage:	Please store the product under the recommended conditions in the Certificate of	0 3-7
	Analysis.	

BIOLOGICAL ACTIVITY				
Description	BRM/BRG1 ATP Inhibitor-3 is a BRG1/BRM inhibitor. BRM/BRG1 ATP Inhibitor-3 inhibits BRM and BRG1 with IC ₅₀ values of 10.4 nM and 19.3 nM, respectively. BRM/BRG1 ATP Inhibitor-3 can be used for the research of cancers and BAF complex-related disorders ^[1] .			
IC ₅₀ & Target	IC50: 10.4 nM (BRM); 19.3 nM (BRG1) ^[1] .			
In Vitro	BRM/BRG1 ATP Inhibitor-3 has inhibitory activity for BRM and BRG1 with IC ₅₀ values of 10.4 nM and 19.3 nM, respectively ^[1] . BRM/BRG1 ATP Inhibitor-3 (0.01-10 μM) has inhibition for cell proliferation in several cancer cell lines ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Proliferation Assay ^[1]			
	Cell Line:	Uveal melanoma cell lines (92-1 , MP41 , MP38, MP46), prostate cancer cell lines (LNCAP), lung cancer cell lines (NCI-H1299), and immortalized embryonic kidney lines (HEK293T)		
	Concentration:	0-10 μΜ		
	Incubation Time:	3 days		
	Result:	Inhibited cell proliferation of uveal melanoma cell line 92-1 and MP41.		

REFERENCES

[1]. Kevin J. Wilson, et al. Compounds and uses thereof. Patent. WO2021155320A1.

Product Data Sheet



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

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