

Product Name: Atractyloside Dipotassium Salt Revision Date: 09/16/2021

Product Data Sheet

Atractyloside Dipotassium Salt

Cat. No.:	A8188	S=0 K* K*
CAS No.:	102130-43-8	
Formula:	C30H44K2O16S2	
M.Wt:	802.99	
Synonyms:		ОН ОН
Target:	Apoptosis	ОН
Pathway:	Apoptosis Inducers	
Storage:	Store at -20°C	
	E Burnen	ОН
Solvent	& Solubility	SS Protocol
	Contraction of the second seco	

	≥37.05 mg/mL in DM	≥37.05 mg/mL in DMSO; ≥51 mg/mL in EtOH; ≥110.4 mg/mL in H2O					
In Vitro		Mass					
	Preparing Stock Solutions	Solvent	1mg	5mg	10mg		
		Concentration					
		1 mM	1.2453 mL	6.2267 mL	12.4535 mL		
		5 mM	0.2491 mL	1.2453 mL	2.4907 mL		
		10 mM	0.1245 mL	0.6227 mL	1.2453 mL		

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

Biological Activity

Shortsummary	AAT inhibitor			
IC50 & Target		al Quer		
	Cell Viability Assay	O E Constant		
	Cell Line:	Ehrlich ascites tumor cells, J2-3T3 cells and cervical carcinoma cells		
In Vitro	Preparation method:	The solubility of this compound in DMSO is > 37.05 mg/mL. 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:	3 mM, 24 h		
1 www.apexbt.com				

	Applications:	In cultured Ehrlich ascites tumor cells, atractyloside (3 mM, 24 h) inhibited cell
	PERSONAL PROPERTY AND	growth by 70% with not severe influence on cell viability. Atractyloside retarded
		cell cycle progression. After transfer of atractyloside treated cells to normal
		medium, proliferation and macromolecular synthesis normalized within 3 to 6 h.
		Atractyloside increased glucose consumption and lactate production.
		Lactate/glucose ratio was 1.9 after 24 h. Atractyloside reduced oxygen uptake.
		Atractyloside increased the ATP/ADP concentration ratio in the mitochondrial
		and extramitochondrial compartment. Atractyloside dose-dependently induced
		apoptosis in normal J2-3T3 cells and cervical carcinoma cells by loss of cell
		viability, nuclear fragmentation and DNA laddering. The sensitivity of cells to
		atractyloside-induced apoptosis was found to be: HPV 16 E6-J2-3T3 > CaSki >
		normal-J2-3T3 cells approximately ts p53-J2-3T3 approximately vector-J2-3T3
		cells > Hela > SiHa > C33A approximately C33A 16 E6. Atractyloside can
		induce opening of the mitochondrial permeability transition pore (mPTP) in
	Bloom	arteriolar smooth muscle cells (ASMCs). Treatment with 7.5, 10, and 15 μM
	Espaceme C	atractyloside for 10 min significantly reduced the relative ATP content in
		ASMCs by 48%, 63% and 66% of control, and ASMCs were hyperpolarized.
	Animal experiment	
	Applications:	
In Vivo	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. Zhu H, Ding Y, et al."Prostaglandin E1 protects coronary microvascular function via the glycogensynthase kinase 3β-mitochondrial permeability transition pore pathway in rathearts subjected to sodium laurate-induced coronary microembolization." Am JTransl Res. 2017 May 15;9(5):2520-2534.PMID:28560002

See more customer validations on www.apexbt.com.

References

Pick-Kober K H, Schneider F. Proliferation, macromolecular synthesis and energy metabolism of in vitro grown Ehrlich ascites tumor cells after inhibition of ATP-ADP translocation by atractyloside[J]. European journal of cell biology, 1984, 34(2): 323-329.
Brown J, Higo H, Mckalip A, et al. Human papillomavirus (HPV) 16 E6 sensitizes cells to atractyloside - induced apoptosis: Role of p53, ICE - like proteases and the mitochondrial permeability transition[J]. Journal of cellular biochemistry, 1997, 66(2): 245-255.
Song R, Bian H, Huang X, et al. Atractyloside induces low contractile reaction of arteriolar smooth muscle through mitochondrial damage[J]. Journal of Applied Toxicology, 2012, 32(6): 402-408.

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 guality of the reagents. Upon receipt of the product, follow the storage recommendations on the product data sheet.



APExBIO Technology www.apexbt.com

7505 Fannin street, Suite 410, Houston, TX 77054. Tel: +1-832-696-8203 | Fax: +1-832-641-3177 | Email: info@apexbt.com







