

Product Name: Rupatadine Fumarate Revision Date: 01/10/2021

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Product Data Sheet

Rupatadine Fumarate

Cat. No.:	B1566	
CAS No.:	182349-12-8	
Formula:	C26H26CIN3·C4H4O4	
M.Wt:	532.03	
Synonyms:		
Target:	Neuroscience	
Pathway:	Histamine Receptor	
Storage:	Store at -20°C	

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Solvent & Solubility

	insoluble in H2O; \geq	insoluble in H2O; \geq 11.75 mg/mL in DMSO; \geq 12.55 mg/mL in EtOH with gentle warming			
Preparing In Vitro Stock Solu	Preparing	Mass Solvent Concentration	1mg	5mg	10mg
	Slock Solutions	1 mM	1.8796 mL	9.3980 mL	18.7959 mL
	018	5 mM	0.3759 mL	1.8796 mL	3.7592 mL
	PERF	10 mM	0.1880 mL	0.9398 mL	1.8796 mL

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

Biological Activity

Shortsummary

Inhibitor of PAF and histamine (H1) receptor

IC₅₀ & Target

In Vitro

Cell Viability Assay	and the second se
Cell Line:	Platelets
Preparation method:	The solubility of this compound in warm ethanol is > 12.6 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:	Up to 100 μM; 5 mins

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	Applications:	Rupatadine Fumarate competitively inhibited platelet-activating factor				
		(PAF)-induced platelet aggregation in washed rabbit platelets (pA2 = 6.68 \pm				
		0.08) and in human platelet-rich plasma (IC50 = 0.68 μ M). However,				
		Rupatadine Fumarate did not affect ADP- or arachidonic acid (AA)-induced				
		platelet aggregation.				
	Animal experiment	619				
In Vivo	Animal models:	Mice and rats				
	Dosage form:	i.v. or p.o				
	Applications:	Rupatadine Fumarate (i.v.) blocked histamine- and PAF-induced hypotension				
		in rats with the ID50 values of 1.4 and 0.44 mg/kg, respectively. Moreover,				
		Rupatadine Fumarate potently inhibited PAF-induced mortality in mice (ID50 =				
		0.31 and 3.0 mg/kg, respectively, for the i.v. and p.o. administrations) and				
		endotoxin-induced mortality in mice and rats (ID50 = 1.6 and 0.66 mg/kg,				
		respectively. i.v.).				
	Other notes:	Please test the solubility of all compounds indoor, and the actual solubility may				
	PERMIT	slightly differ with the theoretical value. This is caused by an experimental				
	P. P. Contraction	system error and it is normal.				

Product Citations

1. Bravo DD, Chernov-Rogan T, et al. "An impedance-based cell contractionassay using human primary smooth muscle cells and fibroblasts." J PharmacolToxicol Methods. 2017 Oct 19;89:47-53.PMID:29056519

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References

[1]. Merlos M, Giral M, Balsa D, Ferrando R, Queralt M, Puigdemont A, García-Rafanell J, Forn J. Rupatadine, a new potent, orally active dual antagonist of histamine and platelet-activating factor (PAF). J Pharmacol Exp Ther. 1997 Jan;280(1):114-21.

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.













