

Product Name: PF-2545920 Revision Date: 01/10/2021

Product Data Sheet

PF-2545920

Cat. No.:	A4325
CAS No.:	1292799-56-4
Formula:	C25H20N4O
M.Wt:	392.45
Synonyms:	
Target:	Metabolism
Pathway:	PDE
Storage:	Store at -20°C

Solvent & Solubility

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	insoluble in H2O; ≥19.35 mg/mL in DMSO; ≥99.8 mg/mL in EtOH					
In Vitro	Preparing Stock Solutions	Mass Solvent Concentration	1mg	5mg	10mg	
		1 mM	2.5481 mL	12.7405 mL	25.4810 mL	
		5 mM	0.5096 mL	2.5481 mL	5.0962 mL	
		10 mM	0.2548 mL	1.2740 mL	2.5481 mL	

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

Biological Activity

Shortsummary	PDE10A inhibitor, potent and selective		
IC50 & Target	0.37 nM (PDE10A)		
	Cell Viability Assay		
	Cell Line:	Rat striatal cells	
	Preparation method:	The solubility of this compound in DMSO is >19.35 mg/ml. General tips for	
In Vitro		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:	1 μM for 30 min at 30°C	
		1 www.apexbt.com	

	Applications:	Biochemical characterization of PF-2545920 showed that it could modulate		
		both the dopamine D1-direct and D2-indirect striatal pathways and regulate the		
		phosphorylation status of a panel of glutamate receptor subunits in the		
		striatum. It was striking that PDE10A inhibition increased the phosphorylation		
		of the (+/-)-alpha-amino-3-hydroxy-5-methylisoxazole-4-proprionic acid		
	al0	receptor GluR1 subunit at residue serine 845 at the cell surface.		
	Animal experiment	SEL		
In Vivo	Animal models:	Male CF-1 mice or Sprague-Dawley rats		
	Dosage form:	1-30 mg/kg, i.p.		
	Applications:	PF-2545920 was active in a range of antipsychotic models, antagonizing		
		apomorphine-induced climbing in mice, inhibiting conditioned avoidance		
		responding in both rats and mice, and blocking N-methyl-D-aspartate		
		antagonist-induced deficits in prepulse inhibition of acoustic startle response in		
		rats, while improving baseline sensory gating in mice, all of which strengthen		
	810	previously reported observations.		
	Other notes:	Please test the solubility of all compounds indoor, and the actual solubility may		
	Sto Provent	slightly differ with the theoretical value. This is caused by an experimental		
		system error and it is normal.		

Product Citations

See more customer validations on www.apexbt.com.



[1] Grauer SM, et al. Phosphodiesterase 10A inhibitor activity in preclinical models of the positive, cognitive, and negative symptoms of schizophrenia. J Pharmacol Exp Ther, 2009, 331(2), 574-590.

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