

Product Name: PF-573228 Revision Date: 01/10/2021

Product Data Sheet

PF-573228

Cat. No.: B1523

CAS No.: 869288-64-2

Formula: C22H20F3N5O3S

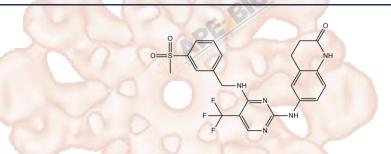
M.Wt: 491.49

Synonyms:

Target: Tyrosine Kinase

Pathway: FAK

Storage: Store at -20°C



Solvent & Solubility

insoluble in EtOH; insoluble in H2O; \geq 166.6 mg/mL in DMSO

In Vitro

Preparing Stock Solutions	Solvent Concentration	1mg	5mg	10mg
	1 mM	2.0346 mL	10.1731 mL	20.3463 mL
	5 mM	0.4069 mL	2.0346 mL	4.0693 mL
	10 mM	0.2035 mL	1.0173 mL	2.0346 mL

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

Biological Activity

Reacting conditions:

Shortsummary	ATP-competitive FAK inhibitor		
IC ₅₀ & Target	4 nM (FAK)		
	Cell Viability Assay		
	Cell Line:	REF52, PC3 or MDCK cells, REF52 cells	
	Preparation method:	The solubility of this compound in DMSO is >166.6mg/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.	

1-10 µM, 24 h

	Applications:	PF-228 inhibited FAK phosphorylation in A431 cells with IC50 of 11 nM. PF-228
		blocked FAK Tyr397 phosphorylation in PC3 (prostate carcinoma), SKOV-3
		(ovarian carcinoma), L3.6p1 and F-G (pancreatic carcinomas), and MDCK
		cells with IC50 of 30-500 nM. In REF52 cells, treatment with 1-3 µM PF-228
		reduced FN-stimulated FAK Tyr397 phosphorylation by ~65–85%. Treatment
	310	with 10 μM PF-228 blocked random migration and efficiently blocked serum
	SE STREET	and FN-stimulated migration. Treatment of cultures with 1 µM PF-228
	Sales La Collection	significantly reduced the rate of movement of individual cells into the wound.
	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. Kath C, Goni-Oliver P, et al. "PTEN suppresses axon outgrowth by down-regulating the level of detyrosinated microtubules." PLoS One. 2018 Apr 4;13(4):e0193257.PMID:29617365

See more customer validations on www.apexbt.com.

References

[1]. Slack-Davis J K, Martin K H, Tilghman R W, et al. Cellular characterization of a novel focal adhesion kinase inhibitor[J]. Journal of Biological Chemistry, 2007, 282(20): 14845-14852.

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.

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













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