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Product Information

Product ID P2133

CAS No. 1403764-72-6

Chemical Name 2-Methoxy-N-(3-Methyl-2-Oxo-1,2,3,4-Tetrahydroquinazolin-6-Yl)

benzenesulfonamide

Synonym PF-06405761

 $\textbf{Formula} \quad C_{16} H_{17} N_3 O_4 S$

Formula Wt. 347.39

Melting Point

Purity ≥98%

Solubility DMSO 69 mg/mL (198.62 mM)

Water Insoluble Ethanol Insoluble

Store Temp -20°C

Ship Temp Ambient

Description PFI-1 is an inhibitor of BET bromodomains (BRDs) that prevents the association of BET with histones; it acts as an acyl-lysine

(Kac) mimetic inhibitor, binding the Kac site of BRD2 and BRD4. PFI-1 exhibits anti-inflammatory and anticancer activities. In H202-treated bronchial epithelial cells, PFI-1 decreases expression of IL-6 and CXCL8. In leukemia cells, PFI-1 downregulates

expression of Myc and AurKB, inducing G1 phase cell cycle arrest and suppressing cell growth.

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Pricing and Availability

Bulk quanitites available upon request

Product ID	Size	List Price
P2133	1 mg	\$55.50
P2133	5 mg	\$98.50
P2133	10 mg	\$167.90
P2133	25 mg	\$318.40

References Khan YM, Kirkham P, Barnes PJ, et al. Brd4 Is Essential for IL-18-Induced Inflammation in Human Airway Epithelial Cells. PLoS One. 2014 Apr 23;9(4):e95051. PMID: 24759736.

Picaud S, Da Costa D, Thanasopoulou A, et al. PFI-1, a highly selective protein interaction inhibitor, targeting BET Bromodomains. Cancer Res. 2013 Jun 1;73(11):3336-46. PMID: 23576556.

Caution: This product is intended for laboratory and research use only. It is not for human or drug use.