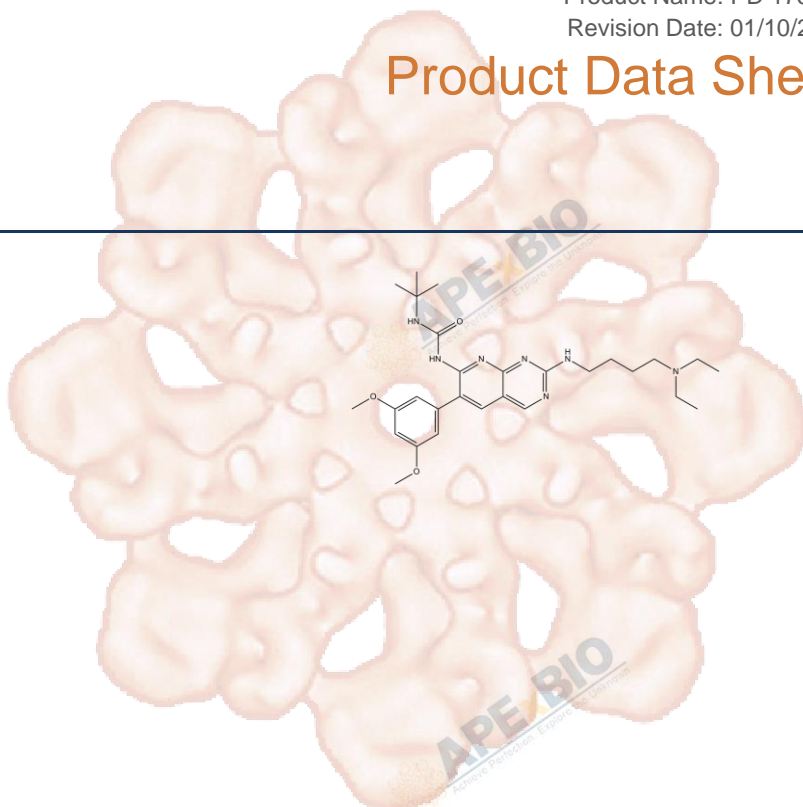


Product Data Sheet

PD 173074

Cat. No.:	A8253
CAS No.:	219580-11-7
Formula:	C ₂₈ H ₄₁ N ₇ O ₃
M.Wt:	523.67
Synonyms:	PD 173074, PD-173074
Target:	Tyrosine Kinase
Pathway:	FGFR
Storage:	Store at 4°C



Solvent & Solubility

≥26.18 mg/mL in DMSO; insoluble in H₂O; ≥108.4 mg/mL in EtOH with ultrasonic

In Vitro

Preparing Stock Solutions	Solvent	Mass		
		1mg	5mg	10mg
	Concentration			
	1 mM	1.9096 mL	9.5480 mL	19.0960 mL
	5 mM	0.3819 mL	1.9096 mL	3.8192 mL
	10 mM	0.1910 mL	0.9548 mL	1.9096 mL

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

Biological Activity

Shortsummary

FGFR inhibitor

IC₅₀ & Target

~25 nM (FGFR1), 100-200 nM (VEGFR2)

In Vitro

Cell Viability Assay

Cell Line: NIH 3T3 cells

Preparation method: The solubility of this compound in DMSO is >10 mM. 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: 0 ~ 1000 nM; 5 mins

	Applications:	PD 173074 dose-dependently inhibited autophosphorylation of FGFR1, with an IC50 value in the range of 1 ~ 5 nM. In addition, PD 173074 inhibited autophosphorylation of VEGFR2 with an IC50 value of 100 ~ 200 nM.
In Vivo	Animal experiment	
	Animal models:	Swiss Webster mice with induced corneal angiogenesis
	Dosage form:	1 or 2 mg/kg/day; i.p.
	Applications:	At the dose of 1 or 2 mg/kg, PD 173074 significantly inhibited angiogenesis induced by either FGF or VEGF in a dose-dependent manner. Besides, it showed no apparent toxicity.
	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

See more customer validations on www.apexbt.com.

References

[1]. Mohammadi M, Froum S, Hamby JM, Schroeder MC, Panek RL, Lu GH, Eliseenkova AV, Green D, Schlessinger J, Hubbard SR. Crystal structure of an angiogenesis inhibitor bound to the FGF receptor tyrosine kinase domain. EMBO J. 1998 Oct 15;17(20):5896-904.

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 APEX BIO products are stable under the recommended conditions. Products are sometimes shipped at a temperature that differs from the recommended storage temperature. Short-term 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.

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