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Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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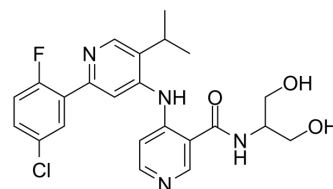
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PF-06952229

Cat. No.:	HY-136244
CAS No.:	1801333-55-0
Molecular Formula:	C ₂₃ H ₂₄ ClFN ₄ O ₃
Molecular Weight:	458.91
Target:	TGF-β Receptor
Pathway:	TGF-beta/Smad
Storage:	Powder -20°C 3 years 4°C 2 years In solvent -80°C 2 years -20°C 1 year



SOLVENT & SOLUBILITY

In Vitro	DMSO : 50 mg/mL (108.95 mM; Need ultrasonic)					
	Preparing Stock Solutions	Solvent Concentration	Mass	1 mg	5 mg	10 mg
		1 mM		2.1791 mL	10.8954 mL	21.7908 mL
		5 mM		0.4358 mL	2.1791 mL	4.3582 mL
		10 mM		0.2179 mL	1.0895 mL	2.1791 mL
Please refer to the solubility information to select the appropriate solvent.						
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (5.45 mM); Clear solution					
	2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (4.53 mM); Clear solution					

BIOLOGICAL ACTIVITY

Description	PF-06952229 is a potent, selective and orally active TGFβR1 inhibitor. PF-06952229 specifically binds to TGFβR1 and prevents TGFβR1-mediated signal transduction. PF-06952229 is a promising antineoplastic agent for the study solid tumors, especially metastatic breast cancer ^[1] .
IC ₅₀ & Target	IC50: transforming growth factor-beta receptor 1 (TGFβR1) ^[1]
In Vivo	PF-06952229 (oral gavage; 30 mg/kg; twice daily; 21 days) combines with Palbociclib 21 days results in an improved inhibition of pSMAD2 in the MCF7 ER ⁺ xenograft breast cancer tumor model. This combination also leads to a significant increase in survival relative to PF-06952229 monotherapy ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	MCF-7 ER ⁺ HER2-xenograft breast cancer tumor model ^[1]
Dosage:	30 mg/kg
Administration:	Oral gavage; twice daily; 44 days
Result:	Resulted in an increase in tumor growth inhibition when combined with Palbociclib.

CUSTOMER VALIDATION

- Cell Discov. 2022 Sep 20;8(1):94.

See more customer validations on www.MedChemExpress.com

REFERENCES

[1]. Flavia Mercer Pernasetti, et al. Combinations of tgfb inhibitors and cdk inhibitors for the treatment of breast cancer. Patent WO2020058820A1.

Caution: Product has not been fully validated for medical applications. For research use only.

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