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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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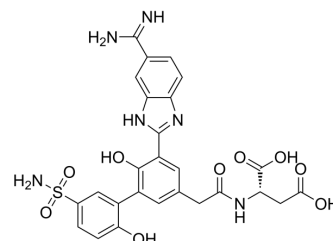
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PCI-27483

Cat. No.:	HY-16382
CAS No.:	871266-63-6
Molecular Formula:	C ₂₆ H ₂₄ N ₆ O ₉ S
Molecular Weight:	596.57
Target:	Others
Pathway:	Others
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 : 62.5 mg/mL (104.77 mM; Need ultrasonic)					
	Preparing Stock Solutions	Solvent Concentration	Mass	1 mg	5 mg	10 mg
		1 mM		1.6762 mL	8.3812 mL	16.7625 mL
		5 mM		0.3352 mL	1.6762 mL	3.3525 mL
		10 mM		0.1676 mL	0.8381 mL	1.6762 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: ≥ 2.08 mg/mL (3.49 mM); Clear solution					
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.08 mg/mL (3.49 mM); Clear solution					
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (3.49 mM); Clear solution					

BIOLOGICAL ACTIVITY

Description	PCI-27483 is a FVIIa/tissue factor inhibitor, with antitumour effects.
In Vitro	PCI-27483 inhibits the TF:FVIIa-complex induced phosphorylation of ERK1/2 and subsequent induction of c-fos in BxPC3 cells, a human pancreatic adenocarcinoma line that highly expresses TF. Furthermore, PCI-27483 blocks the TF:FVIIa induced secretion of IL8 in both BxPC3 cells and MDA-MB-231 breast cancer cells ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

In Vivo

PCI-27483 shows dose-dependent inhibition of thrombus formation, fibrin accumulation and PT. PCI-27483 (4 mg/kg) shows comparable anticoagulation effects as 2 mg/kg enoxaparin^[1]. PCI-27483 (0 and 90 mg/kg, s.c.) results in inhibition of tumor growth in CD1 nu/nu mice implanted with BxPC3 cells^[2].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Gómez-Outes A, et al. New parenteral anticoagulants in development. Ther Adv Cardiovasc Dis. 2011 Feb;5(1):33-59.

[2]. Gómez-Outes A, et al. New parenteral anticoagulants in development. Ther Adv Cardiovasc Dis. 2011 Feb;5(1):33-59.

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

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