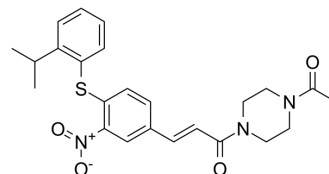


A-286982

Cat. No.:	HY-107587		
CAS No.:	280749-17-9		
Molecular Formula:	C ₂₄ H ₂₇ N ₃ O ₄ S		
Molecular Weight:	453.55		
Target:	Integrin		
Pathway:	Cytoskeleton		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



SOLVENT & SOLUBILITY

In Vitro	DMSO : 50 mg/mL (110.24 mM; ultrasonic and warming and heat to 60°C)				
		Solvent Concentration	Mass 1 mg	5 mg	10 mg
	Preparing Stock Solutions	1 mM	2.2048 mL	11.0241 mL	22.0483 mL
		5 mM	0.4410 mL	2.2048 mL	4.4097 mL
10 mM		0.2205 mL	1.1024 mL	2.2048 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: ≥ 1 mg/mL (2.20 mM); Clear solution 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 1 mg/mL (2.20 mM); Clear solution				

BIOLOGICAL ACTIVITY

Description	A-286982 is a potent and allosteric LFA-1/ICAM-1 interaction inhibitor with IC ₅₀ s of 44 nM and 35 nM in an LFA-1/ICAM-1 binding and LFA-1-mediated cellular adhesion assay, respectively ^{[1][2]} .
IC₅₀ & Target	IC ₅₀ : 44 nM (LFA-1/ICAM-1 interaction in an LFA-1/ICAM-1 binding assay) and 35 nM (LFA-1/ICAM-1 interaction in LFA-1-mediated cellular adhesion assay) ^[1]
In Vitro	A-286982 binds to the I domain allosteric site (IDAS). The allosteric ICAM inhibition such as this would be expected to exhibit the unsurmountable competition we have observed for A-286982 as a result of the passage of this allostery through the A-286982 binding site in its transmission from the β subunit I-like domain to the α subunit ICAM binding site ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

CUSTOMER VALIDATION

- Hematol Oncol. 2023 Jul 26.

See more customer validations on www.MedChemExpress.com

REFERENCES

- [1]. G Liu, et al. Discovery of novel p-arylthio cinnamides as antagonists of leukocyte function-associated antigen-1/intracellular adhesion molecule-1 interaction. 1. Identification of an additional binding pocket based on an anilino diaryl sulfide lead. J Me
- [2]. Susan M Keating, et al. Competition between intercellular adhesion molecule-1 and a small-molecule antagonist for a common binding site on the alpha subunit of lymphocyte function-associated antigen-1. Protein Sci. 2006 Feb;15(2):290-303.
-

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA