

Cilengitide free base

产品编号: D51601

CAS: 188968-51-6

分子式: C₂₇H₄₀N₈O₇

纯度: ≥98%

InChi: InChI=1S/C27H40N8O7/c1-15(2)22-25(41)33-17(10-7-11-30-27(28)29)23(39)31-14-20(36)32-18(13-21(37)38)24(40)34-19(26(42)35(22)3)12-16-8-5-4-6-9-16/h4-6,8-9,15,17-19,22H,7,10-14H,2,1-3H3,(H,31,39)(H,32,36)(H,33,41)(H,34,40)(H,37,38)(H4,28,29,30)/t17-,18-,19+,22-/m0/s1

InChi Key: AMLYAMJWYAIXIA-VWNVYAMZSA-N

Smiles: CC(C)[C@H]1C(=O)N[C@@H](CCCN=C(N)N)C(=O)NCC(=O)N[C@@H](CC(O)=O)C(=O)N[C@H](CC2C=CC=CC=2)C(=O)N1C

外观: 固体粉末

作用通路: Autophagy

溶解性: Soluble in DMSO

保存条件: Store in dry, dark place for one year.

产品介绍: Cilengitide is a cyclic Arg-Gly-Asp peptide with potential antineoplastic activity. Cilengitide binds to and inhibits the activities of the $\alpha(v)\beta(3)$ and $\alpha(v)\beta(5)$ integrins, thereby inhibiting endothelial cell-cell interactions, endothelial cell-matrix interactions, and angiogenesis.

基本信息:

CAS: 188968-51-6

分子式: C₂₇H₄₀N₈O₇

分子量: 588.66

MDL: MFCD04307743

纯度: ≥99%

溶解性: 溶于DMSO, 水。

产品简介:

作用靶点: Integrin; Autophagy;

作用通路: Cytoskeleton; Autophagy;

描述: 西仑吉肽 Cilengitide (EMD 121974) 是一种有效的选择性整合素 $\alpha v\beta 3$ 和 $\alpha v\beta 5$ 抑制剂。Cilengitide 抑制 $\alpha v\beta 3$ 和 $\alpha v\beta 5$ 与玻连蛋白结合, IC₅₀ 值分别为 4 和 79 nM。