PRODUCT INFORMATION



SB-328437

Item No. 29677

CAS Registry No.:	247580-43-4	
Formal Name:	N-(1-naphthalenylcarbonyl)-4-nitro-	NO ₂
	L-phenylalanine, methyl ester	
MF:	C ₂₁ H ₁₈ N ₂ O ₅	
FW:	378.4	
Purity:	≥98%	
UV/Vis.:	λ _{max} : 221, 278 nm	
Supplied as:	A crystalline solid	
Storage:	-20°C	
Stability:	≥4 years	
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Laboratory Procedures

SB-328437 is supplied as a crystalline solid. A stock solution may be made by dissolving the SB-328437 in the solvent of choice, which should be purged with an inert gas. SB-328437 is soluble in the organic solvent DMSO at a concentration of approximately 100 mM.

Description

SB-328437 is a nonpeptide chemokine (C-C motif) receptor 3 (CCR3) antagonist (IC₅₀ = 4.5 nM in a radioligand binding assay).¹ It is selective for CCR3 over chemokine (C-X-C motif) receptor 1 (CXCR1), CXCR2, CCR7, complement 3a receptor (C3aR), and C5aR (IC₅₀s = >27, >27, >33, >10, and >10 μ M, respectively). SB-328437 (3.3-1,000 nM) inhibits calcium mobilization induced chemokine (C-C motif) ligand 11 (CCL11), CCL24, and CCL13 in RBL-2H3-CCR3 cells and eosinophils. In vivo, SB-328437 reduces pulmonary Ccr3 mRNA expression, the number of eosinophils and monocytes in bronchoalveolar lavage fluid (BALF), and pulmonary IL-4 and TNF- α levels in a mouse model of asthma.²

References

- 1. White, J.R., Lee, J.M., Dede, K., et al. Identification of potent, selective non-peptide CC chemokine receptor-3 antagonist that inhibits eotaxin-, eotaxin-2-, and monocyte chemotactic protein-4-induced eosinophil migration. J. Biol. Chem. 275(47), 36626-36631 (2000).
- 2. Wang, T., Zhang, J.-Y., Wang, X.-D., et al. Effects of CCR3(chemokine receptor-3) on Muc5ac in airway of asthmatic mice. Zhongguo Mianyixue Zazhi 31(8), 1019-1023 (2015).

WARNING THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

SAFETY DATA

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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