

(S,R,S)-AHPC-PEG1-NH2 Biochemical

(S,R,S)-AHPC-PEG1-NH2

CATALOG #
AAA228240

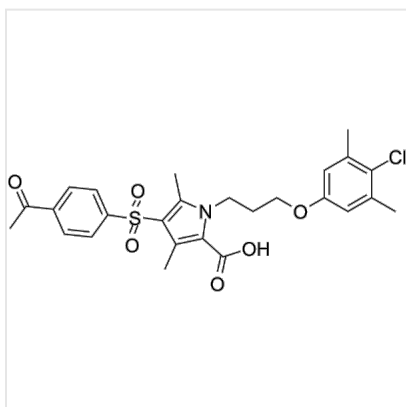
PRODUCT SYNONYMS

(S; R; S)-AHPC-PEG1-NH2; S)-AHPC-PEG1-NH2 biochemical

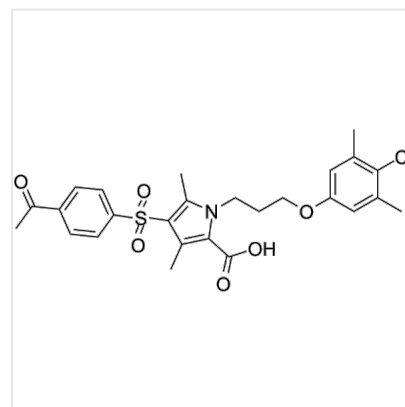
PRODUCT SPECIFICATIONS

Purity/Purification	98%
CAS	2241641-57-4
Solubility	<1mg/ml refers to the product slightly soluble or insoluble
In vitro	PROTACs contain two different ligands connected by a linker; one is a ligand for an E3 ubiquitin ligase and the other is for the target protein. PROTACs exploit the intracellular ubiquitin-proteasome system to selectively degrade target proteins[2].
Preparation and Storage	Powder: -20 degree C for 3 years In solvent: -80 degree C for 2 years

IMAGES



Structure
Mcl-1 inhibitor 6, CAS 2598978-56-2



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ADDITIONAL INFORMATION

Related Product Information for (S,R,S)-AHPC-PEG1-NH2 biochemical	(S,R,S)-AHPC-PEG1-NH2 (VH032-PEG1-NH2) is a synthesized E3 ligase ligand-linker conjugate that incorporates the VH032 based VHL ligand and a linker used in PROTAC technology.
References	1. Peng-Ju Zhu, et al. Discovery of 3,5-Dimethyl-4-Sulfonyl-1 H-Pyrrole-Based Myeloid Cell Leukemia 1 Inhibitors with High Affinity, Selectivity, and Oral Bioavailability. J Med Chem. 2021 Aug 12;64(15):11330-11353.

For Research Use Only. Not for use in diagnostic or therapeutic procedures. Small volumes of product may become entrapped in the seal of the vial during shipment and storage; if necessary, briefly centrifuge the vial on a tabletop centrifuge to dislodge any liquid in the cap.