

## **Product** Data Sheet

## N3-PEG3-vc-PAB-MMAE

Cat. No.: HY-100874

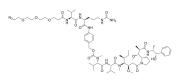
Molecular Formula:  $C_{67}H_{109}N_{13}O_{16}$ Molecular Weight: 1352.66

Target: Drug-Linker Conjugates for ADC

Pathway: Antibody-drug Conjugate/ADC Related

Storage: Powder  $-20^{\circ}\text{C}$  3 years  $4^{\circ}\text{C}$  2 years

\* The compound is unstable in solutions, freshly prepared is recommended.



## **BIOLOGICAL ACTIVITY**

Description	N3-PEG3-vc-PAB-MMAE is a synthesized agent-linker conjugate for ADC that incorporates the MMAE (a tubulin inhibitor) and 3-unit PEG linker. N3-PEG3-vc-PAB-MMAE shows potent antitumor activity. N3-PEG3-vc-PAB-MMAE is a click chemistry reagent, it contains an Azide group and can undergo copper-catalyzed azide-alkyne cycloaddition reaction (CuAAc) with molecules containing Alkyne groups. Strain-promoted alkyne-azide cycloaddition (SPAAC) can also occur with molecules containing DBCO or BCN groups.
IC <sub>50</sub> & Target	Auristatin
In Vitro	Monomethyl auristatin E (MMAE; SGD-1010) is a synthetic derivative of Dolastatin 10 and functions as a potent mitotic inhibitor by inhibiting tubulin polymerization. MMAE is widely used as a cytotoxic component of antibody-drug conjugates (ADCs) to treat several different cancer types.  MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## **CUSTOMER VALIDATION**

- Angew Chem Int Ed Engl. 2020 Jul 27;59(31):12885-12893.
- Chembiochem. 2019 Sep 16;20(18):2411-2419.

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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