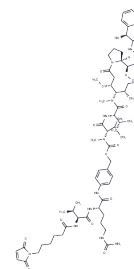


VCMMAE

Chemical Properties

CAS No. :	646502-53-6
Formula:	C ₆₈ H ₁₀₅ N ₁₁ O ₁₅
Molecular Weight:	1316.63
Appearance:	no data available
Storage:	Powder: -20°C for 3 years In solvent: -80°C for 1 year



Biological Description

Description	VCMMAE (mc-vc-PAB-MMAE) is a drug-linker conjugate for ADC with potent antitumor activity.
Targets(IC50)	Microtubule Associated
In vitro	MMAE sensitized colorectal and pancreatic cancer cells to IR in a schedule and dose dependent manner correlating with mitotic arrest. Monomethyl auristatin E (MMAE) is efficiently released from SGN-35 within CD30+ cancer cells and, owing to its membrane permeability, it has the ability to exert cytotoxic activity on bystander cells. Radiosensitization is evidenced by decreased clonogenic survival and increased DNA double strand breaks in irradiated cells.
In vivo	Monomethyl auristatin E (MMAE) in combination with IR results in increased DNA damage signaling and activation of CHK1 that delay the tumor growth. Tumor-targeted ACPD-cRGD-MMAE with IR produces a more robust and significantly prolonged tumor regression in xenograft models.
Cell Research	Monomethyl auristatin E is reconstituted in DMSO at a concentration of 5 nM. Monomethyl auristatin E (MMAE, 5 nM) and ionizing radiation (IR) treated cells are harvested and lysed in RIPA buffer with protease and phosphatase inhibitors. 30 µg of lysate undergo electrophoresis using 4-12% Bis-Tris gels, transferred to PVDF membranes and incubated with indicated primary antibodies.
Animal Research	6-8 week old female athymic nu/nu mice are injected subcutaneously into thighs with 5×10 ⁶ HCT-116 or PANC-1 cells in a 1:1 Matrigel and PBS solution. Mice are treated with IR or intravenous (IV) injection of ACPD-cRGD-MMAE (6 nmoles/day, 18 nmoles total, i.v.), tumor tissue is harvested, formalin fixed and paraffin embedded followed by staining with indicated antibodies. The primary antibody is used at a 1:250 dilution and is visualized using DAB as a chromagen with the UltraMap system.

Solubility Information

Solubility	DMSO: 54 mg/mL (< 1 mg/ml refers to the product slightly soluble or insoluble)
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A DRUG SCREENING EXPERT

Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	0.7595 mL	3.7976 mL	7.5951 mL
5 mM	0.1519 mL	0.7595 mL	1.519 mL
10 mM	0.076 mL	0.3798 mL	0.7595 mL
50 mM	0.0152 mL	0.076 mL	0.1519 mL

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Reference

Okeley, et al. Intracellular Activation of SGN-35, a Potent Anti-CD30 Antibody-Drug Conjugate. Clinical Cancer Research (2010), 16(3), 888-897.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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