

D8-MMAF

Chemical Properties

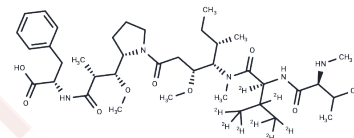
CAS No. :

Formula: C₃₉H₅₇D₈N₅O₈

Molecular Weight: 740.01

Storage: Powder: -20°C for 3 years | In solvent: -80°C for 1 year

Actual storage temperature shall be subject to the COA.



Biological Description

Description	D8-MMAF is a microtubule destroyer, deuterated form of MMAF (T3256-GMP).
In vitro	MMAF showed cytotoxicity in vitro against a group of cell lines. Karpas 299, H3396, 786-O and Caki-1 have IC ₅₀ values of 119, 105, 257 and 200 nM, respectively. Targeted MMAF is more effective than free drug, and the cAC10 conjugate of MMAF shows significant activity. On a molar basis, the average potency of cAC10-L1-MMAF4 is more than 2200 times higher than free MMAF, and it is active in all tested CD30 positive cell
In vivo	The maximum tolerated dose of MMAF (> 16 mg / kg) mice is much higher than that of MMAE (1 mg / kg). The MTD of cAC10-L1-MMAF4 is 50 mg / kg in mice and 15 mg / kg in rats. The corresponding cAC10-L4-MMAF4 ADC is much less toxic, with MTD greater than 150 mg / kg and 90 mg / kg in mice and rats, respectively.

Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	1.3513 mL	6.7567 mL	13.5133 mL
5 mM	0.2703 mL	1.3513 mL	2.7027 mL
10 mM	0.1351 mL	0.6757 mL	1.3513 mL
50 mM	0.027 mL	0.1351 mL	0.2703 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.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

Reference

Doronina SO, et al. Enhanced activity of monomethylauristatin F through monoclonal antibody delivery: effects of linker technology on efficacy and toxicity. *Bioconjug Chem.* 2006 Jan-Feb;17(1):114-24.

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