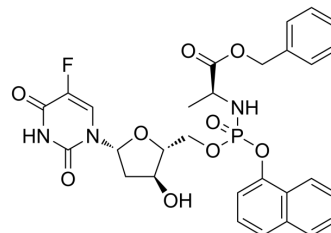


## Fosifloxuridine nafalbenamide

Cat. No.:	HY-109115		
CAS No.:	1332837-31-6		
Molecular Formula:	C <sub>29</sub> H <sub>29</sub> FN <sub>3</sub> O <sub>9</sub> P		
Molecular Weight:	613.53		
Target:	Thymidylate Synthase		
Pathway:	Apoptosis		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 year



### SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (162.99 mM; Need ultrasonic)			
		Solvent Concentration	Mass	
			1 mg	5 mg
			10 mg	
Preparing Stock Solutions	1 mM	1.6299 mL	8.1496 mL	16.2991 mL
	5 mM	0.3260 mL	1.6299 mL	3.2598 mL
	10 mM	0.1630 mL	0.8150 mL	1.6299 mL
Please refer to the solubility information to select the appropriate solvent.				
In Vivo	<ol style="list-style-type: none"> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 40% PEG300 &gt;&gt; 5% Tween-80 &gt;&gt; 45% saline Solubility: ≥ 2.5 mg/mL (4.07 mM); Clear solution</li> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (4.07 mM); Clear solution</li> <li>Add each solvent one by one: 10% DMSO &gt;&gt; 90% corn oil Solubility: ≥ 2.5 mg/mL (4.07 mM); Clear solution</li> </ol>			

### BIOLOGICAL ACTIVITY

Description	Fosifloxuridine nafalbenamide (NUC-3373), a pyrimidine nucleotide analogue, is a Thymidylate synthase inhibitor. Fosifloxuridine nafalbenamide has anticancer activity. Fosifloxuridine nafalbenamide has the potential to evoke a host immune response and enhance immunoresearch <sup>[1][2]</sup> .
In Vitro	Fosifloxuridine nafalbenamide induces the release of damage-associated molecular patterns (DAMPs), with increased expression of calreticulin (CRT) at the cell surface and concomitant loss of nuclear nuclear high mobility group box protein 1 (HMGB1) <sup>[1]</sup> .

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	MCE has not independently confirmed the accuracy of these methods. They are for reference only.
<b>In Vivo</b>	Fosifloxuridine nafalbenamide exhibits anti-cancer activity in HT-29 nude mouse xenograft models <sup>[2]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

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## REFERENCES

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[1]. Fiona G. McKissock, et al. Abstract 1848: NUC-3373 induces ER stress and the release of damage associated molecular patterns in colorectal cancer cells. Cancer Res August 15 2020 (80) (16 Supplement) 1848.

[2]. Essam A. Ghazaly, et al. Abstract B46: NUC-3373: A novel pyrimidine nucleotide analogue that overcomes key cancer drug resistance limiting patient survival. Mol Cancer Ther December 1 2015 (14) (12 Supplement 2) B46.

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**Caution: Product has not been fully validated for medical applications. For research use only.**

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