# Ac-DEVD-AFC

Cat. No.:	HY-P1005			
CAS No.:	201608-14-2	2		
Molecular Formula:	C <sub>30</sub> H <sub>34</sub> F <sub>3</sub> N <sub>5</sub> O	13		0
Molecular Weight:	730			0
Sequence:	N-Acetyl-As	p-Glu-Val	-Asp-7-amido-4-trifluoroMethylcoumarin	ŀ
Sequence Shortening:	Ac-DEVD-7-a	amido-4-	trifluoroMethylcoumarin	
Target:	Fluorescent	: Dye		
Pathway:	Others			
Storage:	Sealed stora	age, away	rfrom moisture and light	
	Powder	-80°C	2 years	
		-20°C	1 year	
	* In solvent	:-80°C,6	months; -20°C, 1 month (sealed storage, away from moisture	
	and light)			

## SOLVENT & SOLUBILITY

In Vitro	H <sub>2</sub> O : < 0.1 mg/mL (in	DMSO : ≥ 50 mg/mL (68.49 mM) H <sub>2</sub> O : < 0.1 mg/mL (insoluble) * "≥" means soluble, but saturation unknown.					
		Solvent Mass Concentration	1 mg	5 mg	10 mg		
	Preparing Stock Solutions	1 mM	1.3699 mL	6.8493 mL	13.6986 mL		
		5 mM	0.2740 mL	1.3699 mL	2.7397 mL		
		10 mM	0.1370 mL	0.6849 mL	1.3699 mL		
	Please refer to the solubility information to select the appropriate solvent.						
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (3.42 mM); Clear solution						
		2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (3.42 mM); Clear solution					
		3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (3.42 mM); Clear solution					

### **BIOLOGICAL ACTIVITY**

Description

Ac-DEVD-AFC is a fluorogenic substrate ( $\lambda_{ex}$ =400 nm,  $\lambda_{em}$ =530 nm).

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# Product Data Sheet

In Vitro	After incubation with Ac-DEVD-AFC for 1 hour, significant increase of caspase-3 activity is observed at 4 hour compare with control. There are no significant increases of caspase-3 activity in Photofrin and LPLI group. The cleavage of Ac-DEVD-AFC in response to caspase-3 activation is remarkably inhibited by shRNA-BimL transfection <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
PROTOCOL	
Cell Assay <sup>[1]</sup>	For the detection of caspase-3 activity, PBS washes cell pellets (derive from either the medium or the adherent cells) which are suspended in extract buffer [25 mM HEPES (pH7.4), 0.1% TritonX-l00, 10% glycerol, 5 mM DTT, 1mM phenylmethylsulfonyl fluoride, 10 mg/mL pepstatin, and 10 mg/mL Leupeptin] and vortexed vigorously. 20µl of extract (corresponding to 10% of the sample) are incubated with the caspase-3 fluorogenic substrates Ac-DEVD-AFC at 100 µM final concentration at room temperature, and caspase-3 activity is measured continuously by monitoring the release of fluorigenic AFC at 37°C <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### **CUSTOMER VALIDATION**

- Cell Rep. 2023 Apr 18;42(5):112414.
- ACS Pharmacol Transl Sci. 2021 Jun 9.
- J Ethnopharmacol. 2020 Nov 15;262:113213.

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#### REFERENCES

[1]. Wang X, et al. Involvement of Bim in Photofrin-mediated photodynamically induced apoptosis. Cell Physiol Biochem. 2015;35(4):1527-36.

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

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