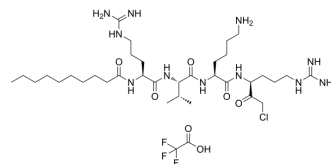


## Decanoyl-RVKR-CMK TFA

Cat. No.:	HY-107760A
CAS No.:	2098497-25-5
Molecular Formula:	C <sub>36</sub> H <sub>67</sub> ClF <sub>3</sub> N <sub>11</sub> O <sub>7</sub>
Molecular Weight:	858.43
Target:	HIV
Pathway:	Anti-infection
Storage:	Sealed storage, away from moisture
	Powder    -80°C    2 years
	-20°C    1 year

\* In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : 50 mg/mL (58.25 mM; Need ultrasonic)  
H<sub>2</sub>O : 4 mg/mL (4.66 mM; Need ultrasonic)

Concentration	Solvent	Mass	1 mg	5 mg	10 mg
			1 mM	1.1649 mL	5.8246 mL
5 mM	0.2330 mL	1.1649 mL	2.3298 mL		
10 mM	0.1165 mL	0.5825 mL	1.1649 mL		

Please refer to the solubility information to select the appropriate solvent.

#### In Vivo

- Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline  
Solubility: 2.5 mg/mL (2.91 mM); Clear solution; Need ultrasonic
- Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)  
Solubility: 2.5 mg/mL (2.91 mM); Clear solution; Need ultrasonic
- Add each solvent one by one: 10% DMSO >> 90% corn oil  
Solubility: 2.5 mg/mL (2.91 mM); Clear solution; Need ultrasonic

### BIOLOGICAL ACTIVITY

#### Description

Decanoyl-RVKR-CMK (DecRVKRcmk) TFA inhibits over-expressed gp160 processing and HIV-1 replication<sup>[1]</sup>.

#### IC<sub>50</sub> & Target

HIV-1

HIV-2

#### In Vitro

Decanoyl-RVKR-CMK (DecRVKRcmk) TFA inhibits HIV-2<sub>ROD</sub> replication by blocking envelope glycoprotein precursor processing in the Jurkat lymphocyte cell<sup>[1]</sup>.

?Decanoyl-RVKR-CMK (DecRVKRcmk) TFA blocks regulated secretion of VGF<sup>[2]</sup>.

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

#### Cell Viability Assay<sup>[1]</sup>

Cell Line:	HeLaCD4 cells infected with recombinant vaccinia viruses at a multiplicity of infection (MOI) of 5 PFU/mL
Concentration:	35 and 70 $\mu$ M
Incubation Time:	7 days
Result:	Peptide at 35 $\mu$ M significantly inhibited ex vivo HIV-1 and HIV-2 replications (70-80% inhibition).

## CUSTOMER VALIDATION

- PLoS Pathog. 2023 Nov 10;19(11):e1011789.

See more customer validations on [www.MedChemExpress.com](http://www.MedChemExpress.com)

## REFERENCES

[1]. B Bahbouhi, et al. Inhibition of HIV-2(ROD) replication in a lymphoblastoid cell line by the alpha1-antitrypsin Portland variant (alpha1-PDX) and the decRVKRcmk peptide: comparison with HIV-1(LAI). *Microbes Infect.* 2001 Nov;3(13):1073-84.

[2]. Angelo L Garcia, et al. A prohormone convertase cleavage site within a predicted alpha-helix mediates sorting of the neuronal and endocrine polypeptide VGF into the regulated secretory pathway. *J Biol Chem.* 2005 Dec 16;280(50):41595-608.

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

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