

Product Data Sheet

Inhibitors • Screening Libraries • Proteins

Proteasome-activating peptide 1 TFA

Cat. No.:	HY-P3414A			
Molecular Formula:	C ₆₅ H ₁₁₇ F ₃ N ₂₂	0 ₁₅ S ₄		
Molecular Weight:	1632.02			
Sequence Shortening:	IPRCRKMPG	VKMC-NF	12	IPRCRKMPGVKMC-NH ₂ (TFA)
Target:	Proteasome	1		2 ()
Pathway:	Metabolic E	nzyme/Pr	otease	
Storage:	Sealed stora	age, 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

$H_2O: \geq 50 \text{ mg/mL}$	(30.64	mM)
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* "≥" means soluble, but saturation unknown.

	Solvent Mass Concentration	1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	0.6127 mL	3.0637 mL	6.1274 mL
	5 mM	0.1225 mL	0.6127 mL	1.2255 mL
	10 mM	0.0613 mL	0.3064 mL	0.6127 mL

Please refer	to the soli	lbility inforr	nation to se	elect the ap	propriate solver	it.

DIOEOGICALACITY					
Description	Proteasome-activating peptide 1 TFA is a peptide and a potent proteasome activator. Proteasome-activating peptide 1 TFA increases the chymotrypsin-like proteasomal catalytic activity and, consequently, proteolytic rates both in vitro and in culture. Proteasome-activating peptide 1 TFA prevents protein aggregation in a cellular model of amyotrophic lateral sclerosis ^[1] .				
In Vitro	 Proteasome-activating peptide 1 (TFA) (0-100 μM, 2 h) increases ChT-L catalytic activity of the 20S proteasome in a dose-dependent manner^[1]. Proteasome-activating peptide 1 (TFA) (50 μM, 24 h) induces the degradation of SOD1, promotes a decrease of oxidized proteins^[1]. Proteasome-activating peptide 1 (TFA) (0-100 μM) protects fibroblast and SH-SY5Y cells from oxidative stress^[1]. Proteasome-activating peptide 1 (TFA) only affects the free 20S proteasomal pool^[1]. Proteasome-activating peptide 1 (TFA) promotes proteasomal gate opening^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Viability Assay 				

Cell Line:	Human neuroblastoma (SH-SY5Y) and fibroblast ${\sf cells}^{[1]}$		
Concentration:	0, 50, 100 μM (in DMEM without fetal bovine serum followed by the addition of H2O2 (500 or 750 $\mu M)$ or water (0.4%))		
Incubation Time:	30 min or 2 h		
Result:	Promoted long-term proteasomal activation through interaction with the 20S core particle, increased ChT-L activity of the 20S proteasome in a dose-dependent manner, and did not reduce cell fibroblast viability.		
Western Blot Analysis			
Cell Line:	SH-SY5Y ^{G93A} cells ^[1]		
Concentration:	50 μΜ		
Incubation Time:	24 h		
Result:	Increased the degradation of soluble SOD1, prevented the formation of SOD1-insolub aggregates, and decreased the oxidized proteins in the total intracellular content.		

REFERENCES

[1]. Dal Vechio FH, Cerqueira F, Augusto O, Lopes R, Demasi M. Peptides that activate the 20S proteasome by gate opening increased oxidized protein removal and reduced protein aggregation. Free Radic Biol Med. 2014 Feb;67:304-13.

[2]. Dal Vechio FH, et al. Peptides that activate the 20S proteasome by gate opening increased oxidized protein removal and reduced protein aggregation. Free Radic Biol Med. 2014;67:304-313.

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

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