

## Catalog # 10-5386 Z-Leu-Arg-AMC · HCI

CAS# 156192-32-4 Z-LR-AMC; CBZ-Leu-Arg-AMC, hydrochloride Lot # X105623



Sensitive Cathepsin K fluorogenic substrate ( $K_m$ =8µM,  $k_{cat}/K_m$ =4 x 10<sup>5</sup> M<sup>-1</sup>s<sup>-1</sup>)<sup>1-3</sup>. It is cleaved more slowly by the following cathepsins ( $k_{cat}/K_m$ : B (10<sup>5</sup>)<sup>4</sup>, F (10<sup>6</sup>)<sup>4</sup>, L (10<sup>6</sup>)<sup>4,5</sup>, L2/V (10<sup>4</sup>)<sup>5</sup>, S (10<sup>5</sup>)<sup>2,4,5</sup>. Also cleaved by malaria parasite proteases berghepain<sup>6</sup>, vivapain-2 and -3<sup>7</sup>, and falcipain-1, -2, and -3<sup>6,8,9</sup>. Excitation: 365nm, Emission: 440nm.

- 1) Bossard et al. (1996), Proteolytic activity of human osteoclast cathepsin K. Expression, purification, activation, and substrate identification; J. Biol. Chem. 271 12517
- Brömme et al. (1996), Human cathepsin O2, a matrix protein-degrading cysteine protease expressed in osteoclasts. Functional expression of human cathepsin O2 in Spodoptera frugiperda and characterization of the enzyme; J. Biol. Chem. 271 2126
- 3) Linnevers et al. (1997), Expression of human cathepsin K in Pichia pastoris and preliminary crystallographic studies of an inhibitor complex; Protein Sci. 6 919
- 4) Wang et al. (1998), Human cathepsin F. Molecular cloning, functional expression, tissue localization, and enzymatic characterization; J. Biol. Chem., 273 32000
- 5) Brömme et al. (1999), Human cathepsin V functional expression, tissue distribution, electrostatic surface potential, enzymatic characterization and chromosomal localization; Biochemistry, **38** 2377
- 6) Singh et al. (2007), A chimeric cysteine protease of Plasmodium berghei engineered to resemble the Plasmodium falciparum protease falcipain-2; Protein Eng. Des. Sel. 20 171
- 7) Na et al. (2004), Identification and biochemical characterization of vivapains, cysteine proteases of the malaria parasite Plasmodium vivax; Biochem. J. 378(Pt2) 529
- 8) Goh et al. (2005), Cysteine protease falcipain 1 in Plasmodium falciparum is biochemically distinct from its isozymes; Parasitol. Res. 97 295
- 9) Pandey et al. (2004), Independent intramolecular mediators of folding, activity, and inhibition for the Plasmodium falciparum cysteine protease falcipain-2; J. Biol. Chem. 279 3484

## PHYSICAL DATA

Molecular Weight:	615.13
Molecular Formula:	C <sub>30</sub> H <sub>38</sub> N <sub>6</sub> O <sub>6</sub> · HCI
Purity:	>97% by HPLC
	Amino acid analysis and identity: Confirmed
	Peptide Content: 93%
Solubility:	DMSO (up to at least 20 mg/ml)
Physical Description:	White solid
Storage and Stability:	Store as supplied desiccated at -20°C for up to 2 years from the date of purchase.
	Solutions in DMSO may be stored at -20°C for up to 3 months. Protect from light.

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