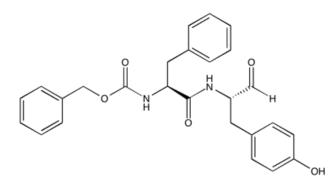


Catalog # 10-3079 Z-Phe-Tyr-CHO

CAS# 167498-29-5 Z-FY-CHO; Z-Phe-Tyr-Aldehyde; SB-412515 Lot # X106290



A potent and selective inhibitor of cathepsin L, IC_{50} =0.85 nM¹ selective over cathepsin B and calpain II (IC_{50} s=85.1 and 184 nM respectively). Suppresses osteoclastic pit formation at 1.5 nM and markedly inhibited parathyroid hormone-stimulated osteoclastic bone resorption.² Also inhibits cathepsin K (K_i s=0.052 and 1.57 nM for cat L and cat K respectively) and partially provides the basis for the finding that cathepsin K is the protease responsible for osteoclastic bone resorption.³ Provides partial protection against serum and potassium deprivation-induced neuronal death.⁴ Active *in vivo*.²

- 1) Woo et al. (1995), Peptidyl aldehyde derivatives as potent and selective inhibitors of cathepsin L; Bioorg. Med. Chem. Lett., 5 1501
- 2) Woo et al. (1996), Suppressive effect of N-(benzyloxycarbonyl)-L-phenylalanyl-L-tyrosinal on bone resorption in vitro and in vivo; Eur. J. Pharmacol., **300** 131
- 3) James et al. (2001) Potent and selective cathepsin L inhibitors do not inhibit human osteoclast resoption in vitro; J. Biol. Chem., 276 11507
- 4) Kaasik et al. (2005), Up regulation if lysosomal cathepsin L and autophagy during neuronal death induced by reduced serum and potassium; Eur. J. Neurosci., **22** 1023

PHYSICAL DATA

Molecular Weight:	446.50
Molecular Formula:	$C_{26}H_{26}N_2O_5$
Purity:	98% by TLC
-	NMR: (Conforms)
Solubility:	DMSO (up to 30 mg/ml) or Ethanol (up to 50 mg/ml)
Physical Description:	Off-white or pale yellow solid
Storage and Stability:	Store as supplied desiccated at -20°C for up to 1 year from the date of purchase. Solutions in
v	DMSO or ethanol may be stored at -20°C for up to 1 month.

Materials provided by Focus Biomolecules are for laboratory research use only and are not intended for human or veterinary applications.