

**Catalog # 10-1309**

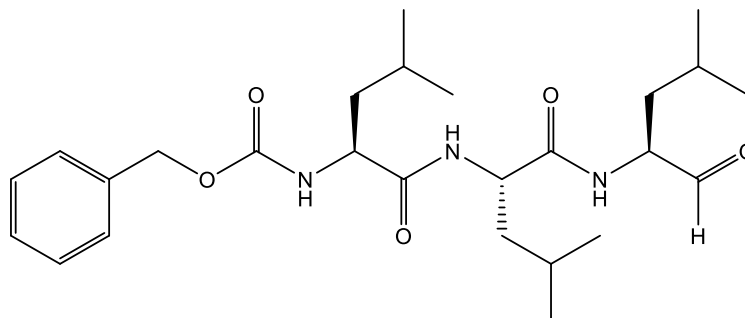
**MG-132**

CAS# 133407-82-6

N-(Benzyloxycarbonyl)leucinylleucinylleucinal

Z-Leu-Leu-Leu-CHO

Lot # X104137



Specific inhibitor of the chymotrypsin-like activity of the 20S proteasome ( $IC_{50}=100$  nM with Z-LLL-AMC as substrate).<sup>1</sup> Also inhibits calpain ( $IC_{50}=1.25$   $\mu$ M)<sup>1</sup> Suppresses gastric cancer cell proliferation and induces macro-autophagy<sup>2</sup>. Activates stress kinases and induces Hsp72.<sup>3</sup> Induces neurite outgrowth.<sup>1</sup> Blocks NF $\kappa$ B activation by blocking I $\kappa$ B proteolysis ( $IC_{50}=3$   $\mu$ M).<sup>4</sup> Cell permeable.

- 1) Tsubuki *et al.* (1996), *Differential inhibition of calpain and proteasome activities by peptidyl aldehydes of di-leucine and tri-leucine*; J. Biochem., **119** 572
- 2) Wu *et al.* (2010), *Macroautophagy and ERK phosphorylation counteract the antiproliferative effect of proteasome inhibitor in gastric cancer cells*; Autophagy, **6** 228
- 3) Meriin *et al.* (1998), *Proteasome inhibitors activate stress kinases and induce Hsp72. Diverse effects on apoptosis*; J. Biol. Chem., **273** 6373
- 4) Fiedler *et al.* (1998), *Inhibition of TNF-alpha-induced NF-kappaB activation and IL-8 release in A549 cells with the proteasome inhibitor MG-132*; Am. J. Respir. Cell Mol. Biol., **19** 259

**PHYSICAL DATA**

Molecular Weight: 475.63  
Molecular Formula: C<sub>26</sub>H<sub>41</sub>N<sub>3</sub>O<sub>5</sub>  
Purity: 98% by HPLC  
Solubility: DMSO (up to 45 mg/ml), DMF (up to 45 mg/ml) or ethanol (up to 45 mg/ml)  
Physical Description: White solid  
Storage and Stability: Store as supplied at -20°C for up to 2 years from the date of purchase. Protect from exposure to moisture. Solutions in DMSO, DMF, or ethanol may be stored at -20°C for up to 1 week

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