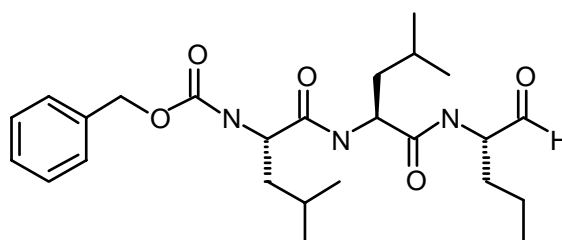


## MATERIAL DATA SHEET

**Z-Leu-Leu-Nva-CHO (MG-115)****Cat. # I-135**

Peptide aldehydes form a covalent hemiacetal adduct with the 20S proteasome, inhibiting the chymotrypsin-like peptidase activity. These inhibitors are reversible and cell-permeable, and also inhibit the activity of calpains and cathepsins.

**Product Information****Quantity:** 5 mg**Formula:** C<sub>25</sub>H<sub>39</sub>N<sub>3</sub>O<sub>5b</sub> **FW:** 461.61**Structure:****Physical/Chemical Characteristics****Solubility:** Soluble in DMSO up to 15 mg/ml. Pellet dry compound prior to reconstitution.**Purity:** > 95% by HPLC. Structure confirmed by NMR.**Activity:** K<sub>i</sub> = 20 nM vs. SDS-activated 20S proteasome.**Use & Storage****Use:** Add from DMSO stock directly to *in vitro* or *in vivo* assay at desired concentration. Pre-incubation with compound prior to assay is recommended for maximal effect.**Storage:** Store stock solution at -20°C. Avoid multiple freeze/thaw cycles.**Literature****References:** Adams J., *et al.* (1998) *Bioorg. Med. Chem. Lett.* **8**:333-338  
Lee D.H. and Goldberg A.L. (1998) *Tren. Cell. Biol.* **8**:397-403  
Lee D.H. and Goldberg A.L. (1996) *J. Biol. Chem.* **271**:27280-27284  
McCormack T.A., *et al.* (1998) *Biochem.* **37**:7792-7800  
Wilk S., *et al.*, (1993) *Enz. Prot.* **47**:306-313

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