Revision 1

Store at -20C MG-132	
Stor	Orders: 877-616-CELL (2355) orders@cellsignal.com
st ^{1 mg}	Support: 877-678-TECH (8324)
194	Web: info@cellsignal.com cellsignal.com
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For Research Use Onl	y. Not for Use in Diagnostic Proce	dures.
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Background	MG-132 is a potent proteasome and calpain inhibitor. Research studies have shown that MG-132 prevents degradation of the proteasome substrates SucLLVY-MCA and ZLLL-MCA with an IC ₅₀ of 850 nM and 100 nM, respectively, as well as inhibits casein-degrading activity of m-calpain with an IC ₅₀ of 1.25 μ M (1). Investigators have demonstrated that MG-132 inhibits TNF- α -induced NF- κ B activation and IL-8 release in A549 cells (2). Proteasome inhibition with MG-132 prevents degradation of short-lived proteins, which correlates with increased expression of HSP and ER chaperone proteins (3). MG-132 induces apoptosis in Hep G2 cells in a time- and dose-dependent manner (4). Proteasome inhibitors like MG-132 are important research tools for studying cellular degradation of the ubiquitin-proteasome pathway.
Molecular Formula	$C_{26}H_{41}N_{3}O_{5}$
Molecular Weight	475.6 g/mol
Purity	>98%
CAS	133407-82-6
Solubility	Soluble in DMSO, DMF and EtOH at 45mg/ml.
Storage	Store lyophilized or in solution at -20°C, desiccated. Protect from light. In lyophilized form, the chemical is stable for 24 months. Once in solution, use within 1 month to prevent loss of potency. Aliquot to avoid multiple freeze/thaw cycles.
Directions for Use	MG-132 is supplied as a lyophilized powder. For a 10 mM stock, reconstitute the 1 mg in 210.3 µl DMSO Working concentrations and length of treatments vary depending on the desired effect, but it is typically used at 5-50 µM for 1-24 hours. Soluble in DMSO or EtOH.
Background References	1. Tsubuki, S. et al. (1996) <i>J Biochem</i> 119, 572-6. 2. Fiedler, M.A. et al. (1998) <i>Am J Respir Cell Mol Biol</i> 19, 259-68. 3. Bush, K.T. et al. (1997) <i>J Biol Chem</i> 272, 9086-92. 4. Emanuele, S. et al. (2002) <i>Int J Oncol</i> 21, 857-65.
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