

Catalog # 10-1384 Epoxomicin CAS# 134381-21-8 Lot # FBM2094



Epoxomicin is a potent, selective and cell permeable irreversible inhibitor of the 20S proteasome.¹ It does not inhibit non-proteasomal proteases such as papain, chymotrypsin, trypsin, calpain and cathepsin B at concentrations up to $50 \mu M.^{1}$ Epoxomicin was isolated from Actinomycete strain Q996-17 and displayed in vivo antitumor activity against B16 melanoma cells.² Epoxomicin caused a progressive model of Parkinson's disease in various systems.^{3,4,5} This model has been disputed.^{6,7}

- 1) Meng *et al.* (1999), *Epoxomicin, a potent and selective proteasome inhibitor, exhibits in vivo anti-inflammatory activity;* Proc. Natl. Acad. Sci. USA, **96** 10403
- 2) Hanada et al. (1992), Epoxomicin, a new antitumor agent of microbial origin; J. Antibiot. (Tokyo), 45 1746
- 3) McNaught *et al.* (2004), *Systemic exposure to proteasome inhibitors causes a progressive model of Parkinson's disease;* Ann. Neurol., **56** 149
- 4) Matsui *et al.* (2010), *Protesasome inhibition in medaka brain induces the features of Parkinson's disease;* J. Neurochem., **115** 178
- 5) Metcalfe *et al.* (2012), *Coordination between proteasome impairment and caspase activation leading to TAU pathology:neuroprotection by cAMP;* Cell Death Diff., **3** e326
- 6) Kordower *et al.* (2006), *Failure of proteasome inhibitor administration to provide a model of Parkinson's disease in rats and monkeys;* Ann. Neurol., **60** 264
- 7) Bove et al. (2006), Proteasome inhibition and Parkinson's disease modeling; Ann. Neurol., 60 260

PHYSICAL DATA

Molecular Weight:	554.73
Molecular Formula:	C ₂₈ H ₅₀ N ₄ O ₇
Purity:	>97% by HPLC
	NMR: (Conforms)
Solubility:	DMSO (up to 10 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. Solutions in DMSO may be stored at -20°C for up to 2 months.

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