

## Catalog # 10-4530 Tubastatin A

CAS# 1310693-92-5

*N*-Hydroxy-4-[(2-methyl-3,4-dihydro-1*H*-pyrido[4,3-*b*]indol-5-yl)methyl]benzamide hydrochloride Lot # FBM2202

Tubastatin A is a potent and highly selective inhibitor of HDAC6 (IC $_{50}$  = 15 nM, >1000-fold selectivity against other HDAC isoforms, approx. 60-fold versus HDAC8).¹ It displayed anti-inflammatory and anti-rheumatic effects in Freund's competitive adjuvant induced animal model of inflammation.² Tubastatin A inhibited TNF- $\alpha$  (IC $_{50}$  = 272 nM) and IL-6 (IC $_{50}$  = 712 nM) in LPS stimulated human THP-1 macrophages.² Tubastatin A has also been shown to have therapeutic potential for treatment of Alzheimers disease via altered amyloid- $\beta$  load and reduced tau hyperphosphorylation.³,4,5

- 1) Butler et al. (2010), Rational design and simple chemistry yield a superior, neuroprotective HDAC6 inhibitor, tubastatin A; J.Am.Chem.Soc. **132** 10842
- 2) Vishwakarma et al. (2013), *Tubastatin, a selective histone deacetylase 6 inhibitor shows anti-inflammatory and anti-rheumatic effects*; Int.Immunopharmacol, **16** 72
- 3) Noack and Richter-Landsberg (2014), HDAC6 inhibition results in tau acetylation and modulates tau phosphorylation and degradation in oligodendrocytes; Glia, **62** 535
- 4) Selenica et al. (2014), Histone deacetylase 6 inhibition improves memory and reduces total tau levels in a mouse model of tau deposition; Alzheimers Res.Ther. **6** 12
- 5) Zhang et al. (2014) Tubastatin A/ACY-1215 improves cognition in Alzheimers disease transgenic mice; J.Alzheimers Dis. **41** 1193

## PHYSICAL DATA

Molecular Weight: 371.86

Molecular Formula:  $C_{20}H_{22}N_3O_2 \cdot HCI$ Purity: 99% by HPLC

NMR: (Conforms)

Solubility: DMSO or water Physical Description: White solid

Storage and Stability: Store as supplied at room -20°C for up to 2 years from the date of purchase. Solutions in

DMSO or water may be stored at -20°C for up to 3 months.

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