

Catalog # 10-4210 BD1047 dihydrobromide

N-[2-(3,4Dichlorophenyl)ethyl]-N-methyl-2-(dimethylamino)ethylamine dihydrobromide 138356-21-5
Lot # JKM1193

BD1047 is a potent and selective sigma-1 antagonist (IC₅₀'s: σ 1 = 0.93nM, σ 2 = 47nM). ¹ Pretreatment of mice with BD1047 reduced convulsions, lethality, and locomotor activity induced by cocaine. ² BD1047 attenuated high fat diet-induced peripheral neuropathy in mice. ³ BD1047 displayed antinociceptive effects in several rodent pain models. ⁴⁻⁶ It also inhibits the β -adrenoreceptor (IC₅₀ = 145 nM). ¹

- 1) Matsumoto et al. (1995), Characterisation of two novel σ receptor ligands: antidystonic effects in rats suggest σ receptor antagonism; Eur.J.Pharmacol. **280** 301
- 2) McCracken et al. (1999), Two novel sigma receptor ligands, BD1047 and LR172, attenuate cocaine-induced toxicity and locomotor activity; Eur.J.Pharmacol. **370** 225
- 3) Song et al. (2017), Role of sigma 1 receptor in high fat diet-induced peripheral neuropathy; Biol.Chem. 398 1141
- 4) Jeong et al. (2005), The spinal antinociceptive mechanism determined by systemic administration of BD1047 in zymosan-induced hyperalgesia in rats; Brain Res. Bull. **119(Pt.A)** 93
- 5) Roh and Yoon (2014), Sigma-1 receptor antagonist BD1047 reduces nociceptive response and phosphorylation of p38 MAPK in mice orofacial formalin model; Biol.Pharm.Bull. **37** 145
- 6) Zhu et al. (2015), Sigma-1 Receptor Antagonist BD1047 Reduces Mechanical Allodynia in a Rat Model of Bone Cancer Pain through the Inhibition of Spinal NR1 Phosphorylation and Microglia Activation; Mediators Inflamm. **2015** 265056

PHYSICAL DATA

Molecular Weight: 437.04

Molecular Formula: $C_{13}H_{20}Cl_2N_2 \cdot 2HBr$ Purity: >98% by HPLC NMR: Conforms

Solubility: DMSO (20 mg/mL) and water (20 mg/mL)

Physical Description: Off-white solid

Storage and Stability: Store as supplied desiccated at -20°C for up to 1 year 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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