

## Catalog # 10-5112 Xanomeline tartrate

CAS# 152854-19-8

3-[4-(Hexyloxy)-1,2,5-thiadiazol-3-yl]-1,2,5,6-tetrahydro-1-methylpyridine tartrate;

LY246708 tartrate

Lot # X109238

A functionally-selective M1 muscarinic receptor agonist, EC<sub>50</sub>=0.3, 92.5, 5, 52 and 42 nM for M1, M2, M3, M4 and M5 respectively. Displays positive cognitive and behavioral effects in schizophrenia and Alzheimer's disease. Suppresses proinflammatory cytokine responses and improves survival in sepsis. Displays potent analgesic activity in rodent models of chronic inflammatory and neuropathic pain.

- 1) Heinrich et al. (2009), Pharmacological Comparison of Muscarininc Ligands: Historical Versus more Recent Muscarinic M1-preferring Receptor Agonists; Eur. J. Pharmacol., **605** 53
- 2) Jakubik et al. (2008), Importance and prospects for design of selective muscarinic agonists; Physiol. Res., 57 Suppl. 3 S39
- 3) Bender et al. (2017), Classics in Chemical Neuroscience: Xanomeline; ACS Chem. Neurosci, 8 435
- 4) Rosas-Ballina et al. (2015), Xanomeline Suppresses Excessive Pro-Inflammatory Cytokine responses Through Neural Signal-Mediated Pathways and Improves Survival in Lethal Inflammation; Brain Behav. Immun., 44 19
- 5) Martino et al. (2011), The M1/M4 Preferring Agonist Xanomeline Is Analgesic in Rodent Models of Chronic Inflammatory and Neuropathic Pain via Central Site of Action; Pain, **152** 2852

## PHYSICAL DATA

Molecular Weight: 431.50

Molecular Formula: C<sub>14</sub>H<sub>23</sub>N<sub>3</sub>OS Purity: 98% by HPLC

NMR: (Conforms)

Solubility: DMSO (up to 45 mg/ml), Water (up to 70 mg/ml)

Physical Description: Off-white solid

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

DMSO or distilled water may be stored at -20°C for up to 2 months.

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