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## Product Data Sheet

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Product Name: Rivanicline hemioxalate

Cat. No.: GC37544

### Chemical Properties

Cas. No.

SMILES CNCC/C=C/C1=CC=CN=C1.OC(C(O)=O)=O.[0.5]

Formula C11H15N2O2 M.Wt 207.23

Solubility DMSO: 50 mg/mL (241.28 mM) Storage Store at -20°C

General tips For obtaining a higher solubility , please warm the tube at 37 °C and shake it in the ultrasonic bath for a while. Stock solution can be stored below -20°C for several months.

Shipping Condition Evaluation sample solution : ship with blue ice All other available size: ship with RT , or blue ice upon request.

Structure

### Background

Rivanicline hemioxalate (RJR-2403 hemioxalate) is a neuronal nicotinic receptor agonist, showing high selectivity for the  $\alpha 4\beta 2$  subtype ( $K_i=26$  nM); > 1,000 fold selectivity than  $\alpha 7$  receptors ( $K_i= 36000$  nM). IC<sub>50</sub> value: 26 nM [1] Target:  $\alpha 4\beta 2$  nAChR in vitro: At concentrations up to 1 mM, Rivanicline does not significantly activate nAChRs in PC12 cells, muscle type nAChRs or muscarinic receptors. Dose-response curves for agonist-induced ileum contraction indicate that Rivanicline is less than one-tenth as potent as nicotine with greatly reduced efficacy. Rivanicline does not antagonize nicotine-stimulated muscle or ganglionic nAChR function (IC<sub>50</sub> > 1 mM). Chronic exposure of M10 cells to Rivanicline (10 microM) results in an up-regulation of high-affinity nAChRs phenomenologically similar to that seen with nicotine [1]. in vivo: Rivanicline significantly improved passive avoidance retention after scopolamine-induced amnesia and enhanced both working and reference memory in rats with ibotenic acid lesions of the forebrain cholinergic projection system in an 8-arm radial maze paradigm. By comparison, Rivanicline was 15 to 30-fold less potent than nicotine in decreasing body temperature, respiration, Y-maze rears and crosses and acoustic startle response [2]. Metanicotine was about 5-fold less potent than nicotine in the tail-flick test after s.c

**Caution: Product has not been fully validated for medical applications. For research use only.**

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administration, but slightly more potent after central administration [3].

[1]. Bencherif M, et al. RJR-2403: a nicotinic agonist with CNS selectivity I. In vitro characterization. J Pharmacol Exp Ther. 1996 Dec;279(3):1413-21. [2]. Lippiello PM, et al. RJR-2403: a nicotinic agonist with CNS selectivity II. In vivo characterization. J Pharmacol Exp Ther. 1996 Dec;279(3):1422-9. [3]. Damaj MI, et al. Antinociceptive and pharmacological effects of metanicotine, a selective nicotinic agonist. J Pharmacol Exp Ther. 1999 Oct;291(1):390-8.

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