

Catalog # 10-1469 ML297

CAS# 1443246-62-5
N-(3.4-Difluorophenyl)-n'-(3-methyl-1-phenyl-1H-pyrazol-5-yl)urea
VU0456810
Lot # S101107

Selective GIRK $\frac{1}{2}$ (K_{ir}3.1/3.2) channel activator, IC₅₀ = 160, 887 and 914 nM for GIRK1/2, GIRK1/4 and GIRK1/3 respectively. Has no effect on GIRK2, GIRK2/3, K_{ir}2.1 and K_v7.4 channels.^{1,2} Displays antiseizure activity² and decreases anxiety-related behavior without sedative or addictive effects³. Reduces glucose- and IBMX-stimulated GLP-1 secretion with no effect on GIP in murine L and K cells.⁴ Brain penetrant.

- 1) Wen et al. (2014), Discovery of potent and selective GIRK1/2 modulators via "molecular switches' within a series of 1-(3-cyclopropyl-1-phenyl-1H-pyrazol-5-yl)ureas; Bioorg. Med. Chem. Lett., **24** 5102
- 2) Kaufmann et al. (2013), ML-297 (VU0456810), the first potent and selective activator of the GIRK otassium channel, displays antiepileptic properties in mice; ACS Chem. Neurosci., **4** 1278
- 3) Wydeven et al. (2014), Mechanisms underlying the activation of G-protein-gated inwardly rectifying K+ (GIRK) channels by the novel anxiolytic drug, ML297; Proc. Natl. Acad. Sci. USA, **111** 10755
- 4) Psichas et al. (2016), Galanin inhibits GLP-1 and GIP secretion via the GAL1 receptor in enteroendocrine L and K cells; Br. J. Pharmacol., 173 888

PHYSICAL DATA

Molecular Weight: 328.32

NMR: (Conforms)

Solubility: Soluble in DMSO (up to 45 mg/ml) or in Ethanol (up to 20 mg/ml)

Physical Description: 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 ethanol may be stored at -20°C for up to 3 months.

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