Catalog \# 10-1469<br>ML297<br>CAS\# 1443246-62-5<br>N -(3.4-Difluorophenyl)-n'-(3-methyl-1-phenyl-1H-pyrazol-5-yl)urea<br>VU0456810<br>Lot \# S101107



Selective GIRK $1 / 2\left(\mathrm{~K}_{\mathrm{ir}} 3.1 / 3.2\right)$ channel activator, $\mathrm{IC}_{50}=160$, 887 and 914 nM for GIRK1/2, GIRK1/4 and GIRK1/3 respectively. Has no effect on GIRK2, GIRK2/3, $\mathrm{K}_{\mathrm{i} r} 2.1$ and $\mathrm{K}_{\mathrm{v}} 7.4$ channels. ${ }^{1,2}$ Displays antiseizure activity $^{2}$ and decreases anxiety-related behavior without sedative or addictive effects ${ }^{3}$. 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., 245102
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., 41278
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, 11110755
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., 173888

## PHYSICAL DATA

| Molecular Weight: | 328.32 |
| :---: | :---: |
| Molecular Formula: | $\mathrm{C}_{17} \mathrm{H}_{14} \mathrm{~F}_{2} \mathrm{~N}_{4} \mathrm{O}$ |
| Purity: | 98\% by TLC |
|  | NMR: (Conforms) |
| Solubility: | Soluble in DMSO (up to $45 \mathrm{mg} / \mathrm{ml}$ ) or in Ethanol (up to $20 \mathrm{mg} / \mathrm{ml}$ ) |
| Physical Description: | White solid |
| Storage and Stability: | Store as supplied desiccated at $-20^{\circ} \mathrm{C}$ for up to 1 year from the date of purchase. |

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