PRODUCT INFORMATION



ML-417

Item No. 36582

CAS Registry No.: 1386162-69-1

Formal Name: 1H-indol-2-yl[4-[2-(4-methoxyphenoxy)

ethyl]-1-piperazinyl]-methanone

MF: $C_{22}H_{25}N_3O_3$ 379.5 FW:

Purity: ≥98%

 λ_{max} : 218, 294 nm A solid UV/Vis.:

Supplied as: Storage: -20°C Stability: ≥4 years

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.

Laboratory Procedures

ML-417 is supplied as a solid. A stock solution may be made by dissolving the ML-417 in the solvent of choice, which should be purged with an inert gas. ML-417 is soluble in ethanol.

Description

ML-417 is a dopamine D_3 receptor agonist.¹ It is selective for the dopamine D_3 receptor over the dopamine D_2 receptor (EC₅₀s = 38 and >10,000 nM, respectively, in β -arrestin recruitment assays using CHO-K1 cells expressing the human receptors) and dopamine D_1 , D_4 , and D_5 receptors, as well as a panel of 45 G protein-coupled receptors (GPCRs), transporters, and ion channels at 10 μM. ML-417 inhibits forskolin-induced cAMP accumulation in HEK293 cells expressing the human dopamine D₃ receptor (IC₅₀ = 86 nM) and induces phosphorylation of ERK in CHO-K1 cells expressing the human dopamine D_3 receptor (EC₅₀ = 21 nM). It protects against cell death induced by the catecholaminergic neurotoxin 6-OHDÅ (Item No. 25330) in human induced pluripotent stem cell-derived (iPSC-derived) cells differentiated into dopaminergic neurons when used at a concentration of 50 nM.

Reference

1. Moritz, A.E., Free, R.B., Weiner, W.S., et al. Discovery, optimization, and characterization of ML417: A novel and highly selective D₃ dopamine receptor agonist. J. Med. Chem. 63(10), 5526-5567 (2020).

WARNING
THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

WARRANTY AND LIMITATION OF REMEDY

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