# PRODUCT INFORMATION



## NIDA-41020

Item No. 21020

CAS Registry No.: 502486-89-7

Formal Name: 1-(2,4-dichlorophenyl)-5-(4-

methoxyphenyl)-4-methyl-N-1-

piperidinyl-1H-pyrazole-3-carboxamide

MF:  $C_{23}H_{24}CI_2N_4O_2$ 

459.4 FW: **Purity:** ≥95%

Supplied as: A crystalline solid

Storage: -20°C Stability: ≥4 years

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

## **Laboratory Procedures**

NIDA-41020 is supplied as a crystalline solid. A stock solution may be made by dissolving the NIDA-41020 in the solvent of choice, which should be purged with an inert gas. NIDA-41020 is soluble in organic solvents such as ethanol and DMSO. The solubility of NIDA-41020 in these solvents is approximately 100 mM.

## Description

NIDA-41020 is a cannabinoid (CB) receptor 1 antagonist ( $K_i = 4.1 \text{ nM}$ ).<sup>1</sup> It binds selectively to CB<sub>1</sub> over CB<sub>2</sub> (K<sub>b</sub>s = 26 and 831 nM, respectively).<sup>2</sup> NIDA-41020 was designed as a potential radioligand for use in positron emission tomography (PET).1

### References

- 1. Katoch-Rouse, R., Pavlova, O.A., Caulder, T., et al. Synthesis, structure-activity relationship, and evaluation of SR141716 analogues: Development of central cannabinoid receptor ligands with lower lipophilicity. J. Med Chem. 46(4), 642-645 (2003).
- 2. Donohue, S.R., Halldin, C., and Pike, V.W. Synthesis and structure-activity relationships (SARs) of 1,5-diarylpyrazole cannabinoid type-1 (CB<sub>1</sub>) receptor ligands for potential use in molecular imaging. Bioorg. Med. Chem. 14(11), 3712-3720 (2006).

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.

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