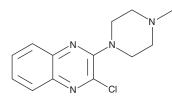
PRODUCT INFORMATION



VUF 10166

Item No. 31423

155584-74-0
2-chloro-3-(4-methyl-1-
piperazinyl)-quinoxaline
$C_{13}H_{15}CIN_4$
262.7
≥98%
λ _{max} : 214, 259 nm
A solid
-20°C
≥4 years
the product specifications. Bat



tch specific analytical results are provided on each certificate of analysis.

Laboratory Procedures

VUF 10166 is supplied as a solid. A stock solution may be made by dissolving the VUF 10166 in the solvent of choice, which should be purged with an inert gas. VUF 10166 is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide (DMF). The solubility of VUF 10166 in ethanol is approximately 10 mg/ml and approximately 30 mg/ml in DMSO and DMF.

VUF 10166 is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, VUF 10166 should first be dissolved in DMSO and then diluted with the aqueous buffer of choice. VUF 10166 has a solubility of approximately 0.25 mg/ml in a 1:3 solution of DMSO:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

Description

VUF 10166 is an antagonist of the serotonin (5-HT) receptor subtype 5-HT_{3A} (K_i = 0.04 nM).¹ It is selective for 5-HT_{3A} over 5-HT_{3AB} receptors in radioligand binding assays (K_i = 2 nM), as well as α 7 nicotinic acetylcholine receptors (nAChRs) at 100 μ M. VUF 10166 also acts as a partial agonist at 5-HT_{3A} receptors $(EC_{50} = 5 \mu M \text{ in a patch-clamp assay using Xenopus oocytes expressing the human receptor).}$

Reference

1. Thompson, A.J., Verheij, M.H.P., de Esch, I.J.P., et al. VUF10166, a novel compound with differing activities at 5-HT_{3A} and 5-HT_{3AB} receptors. J. Pharmacol. Exp. Ther. 341(2), 350-359 (2012).

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

SAFETY DATA

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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