PRODUCT INFORMATION



kb-NB77-78

Item No. 21562

0,	1350622-33-1	Si o
Formal Name:	9-[[(1,1-dimethylethyl)dimethylsilyl]oxy]-1,2,3,4-	
	tetrahydro-5H-[1]benzopyrano[3,4-b]pyridin-5-one	
MF:	C ₁₈ H ₂₅ NO ₃ Si	
FW:	331.5	\sim
Purity:	≥98%	$\int \int $
UV/Vis.:	λ _{max} : 214, 252, 346 nm	
Supplied as:	A crystalline solid	N T
Storage:	-20°C	L Ö
Stability:	≥4 years	п
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Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.

Laboratory Procedures

kb-NB77-78 is supplied as a crystalline solid. A stock solution may be made by dissolving the kb-NB77-78 in the solvent of choice, which should be purged with an inert gas. kb-NB77-78 is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide (DMF). The solubility of kb-NB77-78 in these solvents is approximately 20, 25, and 30 mg/ml, respectively.

kb-NB77-78 is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, kb-NB77-78 should first be dissolved in DMF and then diluted with the aqueous buffer of choice. kb-NB77-78 has a solubility of approximately 0.3 mg/ml in a 1:2 solution of DMF:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

Description

kb-NB77-78 is an inactive analog of the protein kinase D (PKD) inhibitor CID755673 (Item No. 15924).¹ It does not bind PKD1 in a fluorescence polarization assay and has no effect on PKD1 phosphorylation in LNCaP cancer cells.

Reference

1. George, K.M., Frantz, M.C., Bravo-Altamirano, K., et al. Design, synthesis, and biological evaluation of PKD inhibitors. Pharmaceutics 3(2), 186-228 (2011).

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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