PRODUCT INFORMATION



LPA₂ Antagonist 1

Item No. 22051

CAS Registry No.: Formal Name:	1017606-66-4 N-[(1S)-2-[4-[(3,4-dichlorophenyl)sulfonyl]- 1-piperazinyl]-1-methylethyl]-7-methyl- thieno[3,2-d]pyrimidin-4-amine		
MF:	C ₂₀ H ₂₃ CL ₂ N ₅ O ₂ S ₂		· · · · ·
FW:	500.5	ĊI	
Purity:	≥98%		N S
UV/Vis.:	λ _{max} : 242, 302 nm		
Supplied as:	A crystalline solid		N N
Storage:	-20°C		
Stability:	≥4 years		
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Laboratory Procedures

LPA2 antagonist 1 is supplied as a crystalline solid. A stock solution may be made by dissolving the LPA₂ antagonist 1 in the solvent of choice, which should be purged with an inert gas. LPA₂ antagonist 1 is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide. The solubility of LPA₂ antagonist 1 in these solvents is approximately 30 mg/ml.

LPA₂ antagonist 1 is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, LPA₂ antagonist 1 should first be dissolved in ethanol and then diluted with the aqueous buffer of choice. LPA₂ antagonist 1 has a solubility of approximately 0.33 mg/ml in a 1:2 solution of ethanol:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

Description

LPA₂ antagonist 1 is an antagonist of lysophosphatidic acid receptor 2 (LPA₂; IC₅₀ = 17 nM).¹ It is selective for LPA₂ over LPA₁ and LPA₃ (IC₅₀s = >50 μ M). LPA₂ antagonist 1 inhibits HGF-induced phosphorylation of ERK and proliferation of HCT116 colon cancer cells in a concentration-dependent manner.

Reference

1. Beck, H.P., Kohn, T., Rubenstein, S., et al. Discovery of potent LPA₂ (EDG4) antagonists as potential anticancer agents. Bioorg. Med. Chem. Lett. 18(3), 1037-1041 (2008).

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

SAFETY DATA

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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