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Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

Weitere Information auf den folgenden Seiten!
See the following pages for more information!



Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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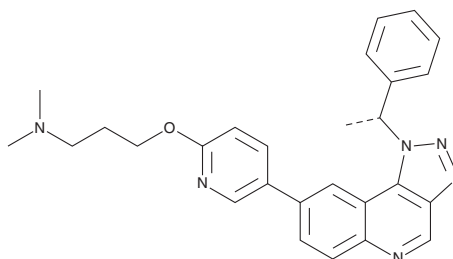
PRODUCT INFORMATION



A011

Item No. 39183

Formal Name: (R)-N,N-dimethyl-3-((5-(1-(1-phenylethyl)-1H-[1,2,3]triazolo[4,5-c]quinolin-8-yl)pyridin-2-yl)oxy)propan-1-amine
MF: C₂₇H₂₈N₆O
FW: 452.6
Purity: ≥95%
UV/Vis.: λ_{max}: 234, 258, 305 nm
Supplied as: A solid
Storage: -20°C
Stability: ≥3 years



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

Laboratory Procedures

A011 is supplied as a solid. A stock solution may be made by dissolving the A011 in the solvent of choice, which should be purged with an inert gas. A011 is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide. The solubility of A011 in these solvents is approximately 10 mg/ml.

Further dilutions of the stock solution into aqueous buffers or isotonic saline should be made prior to performing biological experiments. Ensure that the residual amount of organic solvent is insignificant, since organic solvents may have physiological effects at low concentrations. Organic solvent-free aqueous solutions of A011 can be prepared by directly dissolving the solid in aqueous buffers. A011 is slightly soluble in PBS (pH 7.2). We do not recommend storing the aqueous solution for more than one day.

Description

A011 is an inhibitor of ataxia-telangiectasia mutated (ATM) kinase (IC₅₀ = 1 nM).¹ It is selective for ATM over a panel of 408 additional kinases at 10 μM. A011 inhibits the growth of SW620 and HCT116 colorectal cancer cells (IC₅₀s = 0.14 and 0.91 μM, respectively) and enhances cell death induced by DNA topoisomerase I inhibitor irinotecan (Item No. 14180) in the same cells when used at concentrations of 10, 30, and 100 nM. It also increases irradiation-induced cell death in SW620 cells. *In vivo*, A011 (5 mg/kg) increases irinotecan-induced reductions in tumor growth in an SW620 mouse xenograft model.

Reference

1. Zhang, S., Zhou, P., Liu, J., *et al.* Discovery of [1,2,3]triazolo[4,5- c]quinoline derivatives as a new class of ataxia-telangiectasia mutated kinase inhibitors. *ACS Med. Chem. Lett.* **14**(6), 746-756 (2023).

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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