PRODUCT INFORMATION

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BAY-3827

Item No. 41448

CAS Registry No.: Formal Name:	2377576-35-5 N-[5-(3,5-dicyano-1,4-dihydro- 1,2,6-trimethyl-4-pyridinyl)-6- fluoro-7-methyl-1H-indazol-3-yl]- 2-ethyl-benzamide	
MF:	C ₂₇ H ₂₅ FN ₆ O	
FW:	468.5	
Purity:	≥98%	
Supplied as:	A 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

BAY-3827 is supplied as a solid. A stock solution may be made by dissolving the BAY-3827 in the solvent of choice, which should be purged with an inert gas. BAY-3827 is sparingly soluble (1-10 mg/ml) in DMSO.

Description

BAY-3827 is an inhibitor of AMP-activated protein kinase (AMPK; IC_{50} = 1.4 nM at 10 μ M ATP).¹ It is selective for AMPK over Aurora A kinase, FMS-related tyrosine kinase 3 (FLT3), and c-Met (IC₅₀s = 1,324, 124, and 788 nM, respectively), p90 ribosomal S6 kinase 2 (RSK2), RSK3, RSK4, p90 ribosomal S6 kinase 1 (MSK1), and MST3 (IC₅₀s = 52, 36, 24, 43, and 94 nM, respectively), and a panel of 331 kinases at 10 μ M but does inhibit RSK1 (IC₅₀ = 9 nM). BAY-3827 selectively inhibits the proliferation of androgendependent LNCaP, VCaP, and 22Rv1 prostate cancer cells (IC₅₀s = 0.28, 1.71, and 5.55 µM, respectively) over androgen receptor-negative C4-2B, PC3, and DU145 prostate cancer cells (IC_{50} = >10 μ M for all).

Reference

1. Lemos, C., Schulze, V.K., Baumgart, S.J., et al. The potent AMPK inhibitor BAY-3827 shows strong efficacy in androgen-dependent prostate cancer models. Cell. Oncol. (Dordr) 44(3), 581-594 (2021).

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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