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Produktinformation



Forschungsprodukte & Biochemikalien



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Diagnostik & molekulare Diagnostik



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Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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PRODUCT INFORMATION

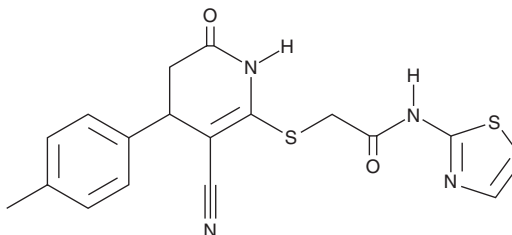


Necrostatin-34

Item No. 37735

CAS Registry No.: 375835-43-1
Formal Name: 2-[[3-cyano-1,4,5,6-tetrahydro-4-(4-methylphenyl)-6-oxo-2-pyridinyl]thio]-N-2-thiazolyl-acetamide

Synonym: Nec-34
MF: C₁₈H₁₆N₄O₂S₂
FW: 384.5
Purity: ≥98%
UV/Vis.: λ_{max}: 280 nm
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

Necrostatin-34 is supplied as a solid. A stock solution may be made by dissolving the necrostatin-34 in the solvent of choice, which should be purged with an inert gas. Necrostatin-34 is soluble in the organic solvent DMSO.

Description

Necrostatin-34 is an inhibitor of receptor-interacting serine/threonine kinase 1 (RIPK1; IC₅₀ = 5.5 μM).¹ It inhibits TNF-α-induced necroptosis in Jurkat cells lacking Fas-associated death domain protein (FADD) and L-929 cells (IC₅₀s = 6.67 and 1.35 μM, respectively). Necrostatin-34 (10 μM) inhibits necroptosis induced by the chemical inducer of dimerization for FK506 binding protein (FKBP) fusion proteins AP20187 (Item No. 36808) and the pan-caspase inhibitor Z-VA-DL-D(OMe)-FMK (zVAD-FMK; Item No. 27421) in L-929 cells expressing an FKBP and RIPK1 fusion protein.

Reference

1. Meng, H., Wu, G., Zhao, X., *et al.* Discovery of a cooperative mode of inhibiting RIPK1 kinase. *Cell Discov.* **7**(1), 41 (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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