## MLKL-IN-3

**MedChemExpress** 

Cat. No.: CAS No.:	HY-151541 3031406-27-3	
Molecular Formula:	C <sub>31</sub> H <sub>29</sub> ClN <sub>4</sub> O <sub>6</sub>	
Molecular Weight:	589.04	
Target:	Mixed Lineage Kinase; Necroptosis	
Pathway:	MAPK/ERK Pathway; Apoptosis	
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.	

BIOLOGICAL ACTIVITY		
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Description	MLKL-IN-3 (compound 66) is a potent MLKL (Mixed lineage kinase domain-like protein) inhibitor. MLKL-IN-3 inhibits necroptosis in HT-29 cells and acts downstream of MLKL phosphorylation, with EC <sub>50</sub> of 31 nM <sup>[1]</sup> . MLKL-IN-3 is a click chemistry reagent, it contains an Alkyne group and can undergo copper-catalyzed azide-alkyne cycloaddition (CuAAc) with molecules containing Azide groups.	
In Vitro	MLKL-IN-3 (compound 66) (1 μM, 24 h) inhibits the translocation of MLKL to cell membranes in HT-29 cells <sup>[1]</sup> . MLKL-IN-3 does not affect the phosphorylation status of RIPK1 and MLKL <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	
In Vivo	MLKL-IN-3 (compound 66) shows T1/2 over 48 h and more than 150-fold lower reaction rates with glutathione (GSH), which potentially decreased their off-target effects and cell toxicity <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	

## REFERENCES

[1]. Cui B, et al. Discovery of a New Class of Uracil Derivatives as Potential Mixed Lineage Kinase Domain-like Protein (MLKL) Inhibitors. J Med Chem. 2022 Oct 13;65(19):12747-12780.

Caution: Product has not been fully validated for medical applications. For research use only.

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