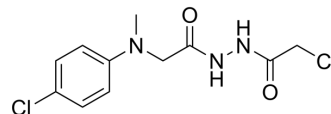


UCHL1-IN-1

Cat. No.:	HY-161378
Molecular Formula:	C ₁₁ H ₁₃ Cl ₂ N ₃ O ₂
Molecular Weight:	290.15
Target:	Deubiquitinase
Pathway:	Cell Cycle/DNA Damage
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	UCHL1-IN-1 (Compound 46) is an inhibitor for Ubiquitin C-terminal Hydrolase L1 (UCHL1), with an IC ₅₀ of 5.1 μM. UCHL1-IN-1 can be used for cancer research ^[1] .
IC ₅₀ & Target	UCHL1
In Vitro	UCHL1-IN-1 (0-50 μM, 4 h) inhibits UCHL1 in BT549 cell ^[1] . UCHL1-IN-1 (0-50 μM, 24 h) inhibits cell migration and cell viability in BT549 cell ^[1] . UCHL1-IN-1 is relatively metabolically stable in human liver microsomes assay, with a t _{1/2} > 120 min, and low intrinsic hepatic clearance ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Imhoff RD, et al. Covalent Fragment Screening and Optimization Identifies the Chloroacetohydrazide Scaffold as Inhibitors for Ubiquitin C-terminal Hydrolase L1. J Med Chem. 2024 Mar 15.

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

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