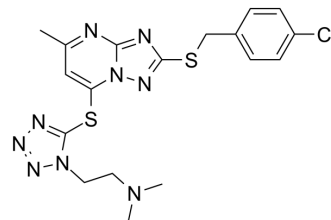


WS-383 free base

Cat. No.:	HY-126075
CAS No.:	2247543-65-1
Molecular Formula:	C ₁₈ H ₂₀ ClN ₉ S ₂
Molecular Weight:	461.99
Target:	E1/E2/E3 Enzyme
Pathway:	Metabolic Enzyme/Protease
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	WS-383 free base is a potent, selective and reversible inhibitor of DCN1-UBC12 interaction, with an IC ₅₀ of 11 nM. WS-383 free base inhibits Cul3/1 neddylation, induces accumulation of p21, p27 and NRF2 ^[1] .																
IC₅₀ & Target	IC ₅₀ : 11 nM (DCN1-UBC12 interaction) ^[1]																
In Vitro	<p>WS-383 (10 μM) is against a panel of kinases such as BTK, CDKs, and EGFR [L858R] using staurosporine and afatinib as the positive controls. WS-383 showed weak inhibitory activity at 10.0 μM, it is selective to the DCN1-UBC12 interaction over the selected kinases^[1].</p> <p>WS-383 (0.03-3 μM; 24 hours) blocks Cul3 neddylation at 3 μM and also has certain inhibition of Cul1 neddylation at 10 μM but was not effective in inhibiting neddylation of other cullin members^[1].</p> <p>WS-383 (0.03-3 μM; 24 hours) increases Cul1, Skp1 (adaptor protein), F-box protein, and RBX1/RBX2 RING protein form SCF E3 complex. Cyclin dependent kinase inhibitor 1A (p21) and cyclin dependent kinase inhibitor 1B (p27) expression in a dose-dependent manner in MGC-803 and KYSE70 manner^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Western Blot Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>MGC-803 cells</td> </tr> <tr> <td>Concentration:</td> <td>0.03 μM; 0.3 μM; 3 μM; 10 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 hours</td> </tr> <tr> <td>Result:</td> <td>Decreased N8-Cul1 and N8-Cul2 protein expression.</td> </tr> </table> <p>Western Blot Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>MGC-803 and KYSE70 cells</td> </tr> <tr> <td>Concentration:</td> <td>0.03 μM; 0.3 μM; 3 μM; 10 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 hours</td> </tr> <tr> <td>Result:</td> <td>Induced accumulation of p21, p27, and NRF2 in MGC-803 cells.</td> </tr> </table>	Cell Line:	MGC-803 cells	Concentration:	0.03 μM; 0.3 μM; 3 μM; 10 μM	Incubation Time:	24 hours	Result:	Decreased N8-Cul1 and N8-Cul2 protein expression.	Cell Line:	MGC-803 and KYSE70 cells	Concentration:	0.03 μM; 0.3 μM; 3 μM; 10 μM	Incubation Time:	24 hours	Result:	Induced accumulation of p21, p27, and NRF2 in MGC-803 cells.
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REFERENCES

[1]. Wang S, et al. Development of Highly Potent, Selective, and Cellular Active Triazolo[1,5- a]pyrimidine-Based Inhibitors Targeting the DCN1-UBC12 Protein-Protein Interaction. J Med Chem. 2019 Mar 14;62(5):2772-2797.

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

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