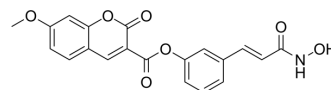


## HDAC-IN-42

Cat. No.:	HY-147892
CAS No.:	2454024-18-9
Molecular Formula:	C <sub>20</sub> H <sub>15</sub> NO <sub>7</sub>
Molecular Weight:	381.34
Target:	HDAC; Apoptosis
Pathway:	Cell Cycle/DNA Damage; Epigenetics; Apoptosis
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	HDAC-IN-42 (compound 14f) is a potent and selective HDAC inhibitor with IC <sub>50</sub> values of 0.19 and 4.98 μM for HDAC1 and HDAC6, respectively. HDAC-IN-42 shows anticancer and anti-proliferative activity. HDAC-IN-42 induces apoptosis and cell cycle arrest at G2/M phase <sup>[1]</sup> .	
<b>IC<sub>50</sub> &amp; Target</b>	HDAC1 0.19 μM (IC <sub>50</sub> )	HDAC6 4.98 μM (IC <sub>50</sub> )
<b>In Vitro</b>	<p>HDAC-IN-42 (compound 14f) shows anti-proliferative activity with C<sub>50</sub>s of 9.56, 13.32, 10.46, 6.91 μM for MCF-7, HCT-116, HepG2, HeLa cells, respectively<sup>[1]</sup>.</p> <p>HDAC-IN-42 (1, 5, 10 μM; 24 h) reduces the colony formation and increases the expression of histone H3 and α-tubulin in HeLa cells<sup>[1]</sup>.</p> <p>HDAC-IN-42 (1, 5, 10 μM; 48 h) induces apoptosis and cell cycle arrest at G2/M phase<sup>[1]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p>	

### REFERENCES

[1]. Ding J, et al. Design, synthesis and biological evaluation of coumarin-based N-hydroxycinnamide derivatives as novel histone deacetylase inhibitors with anticancer activities. *Bioorg Chem.* 2020 Aug;101:104023.

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

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