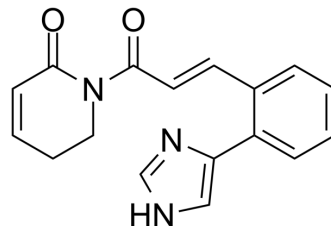


## ZC0101

<b>Cat. No.:</b>	HY-147772
<b>CAS No.:</b>	2541604-52-6
<b>Molecular Formula:</b>	C <sub>17</sub> H <sub>15</sub> N <sub>3</sub> O <sub>2</sub>
<b>Molecular Weight:</b>	293.32
<b>Target:</b>	Indoleamine 2,3-Dioxygenase (IDO); Apoptosis; Reactive Oxygen Species
<b>Pathway:</b>	Metabolic Enzyme/Protease; Apoptosis; Immunology/Inflammation; NF-κB
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	ZC0101 is a potent, orally active IDO1 and TrxR dual inhibitor with IC <sub>50</sub> values of 0.084 μM and 7.98 μM, respectively. ZC0101 effectively induces apoptosis and ROS accumulation in cancer cells <sup>[1]</sup> .	
<b>IC<sub>50</sub> &amp; Target</b>	IDO1 0.084 μM (IC <sub>50</sub> )	TrxR 7.98 μM (IC <sub>50</sub> )
<b>In Vivo</b>	ZC0101 (60 mg/kg; p.o.; once) effectively suppresses IDO1 activity in vivo <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	

### REFERENCES

[1]. Fan QZ, et al. Design, synthesis, and biological evaluation of a novel indoleamine 2,3-dioxygenase 1 (IDO1) and thioredoxin reductase (TrxR) dual inhibitor. *Bioorg Chem.* 2020 Dec;105:104401.

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

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