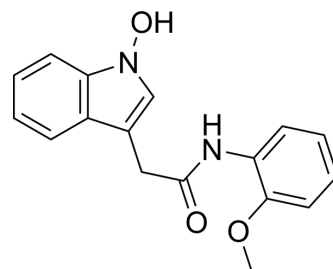


## ZT55

Cat. No.:	HY-124727
CAS No.:	2138488-38-5
Molecular Formula:	C <sub>17</sub> H <sub>16</sub> N <sub>2</sub> O <sub>3</sub>
Molecular Weight:	296.32
Target:	JAK; Apoptosis
Pathway:	Epigenetics; JAK/STAT Signaling; Protein Tyrosine Kinase/RTK; Stem Cell/Wnt; Apoptosis
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	ZT55 is an orally active and highly-selective JAK2 inhibitor with an IC <sub>50</sub> value of 0.031 μM. ZT55 inhibits the proliferation of JAK2 <sup>V617F</sup> -expressing HEL cell lines and induces apoptosis and cycle arrest. ZT-55 also effectively inhibits the growth of HEL xenograft tumours in a mice model. ZT-55 can be used in studies of myeloproliferative neoplasms, polycythemia vera and primary thrombocythemia <sup>[1]</sup> .																
<b>IC<sub>50</sub> &amp; Target</b>	JAK2 0.031 μM (IC <sub>50</sub> )																
<b>In Vitro</b>	<p>ZT55 (0-100 μM; 48 h) inhibits the proliferation of JAK2<sup>V617F</sup> (+) HEL cells in a concentration-dependent manner (IC<sub>50</sub>=18.05 μM)<sup>[1]</sup>.</p> <p>ZT55 (12.5, 25, 50 μM; 24, 48, 72 h) inhibits the cell viability and induces apoptosis of HEL cells in a concentration- and time-dependent manner<sup>[1]</sup>.</p> <p>ZT55 (12.5, 25, 50 μM; 24 h) arrests HEL cells in the G2/M phase in a concentration-dependent manner<sup>[1]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Proliferation Assay<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>HEL cells (JAK2<sup>V617F</sup>(+))</td> </tr> <tr> <td>Concentration:</td> <td>0-100 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>48 h</td> </tr> <tr> <td>Result:</td> <td>Significantly inhibited proliferation in a concentration-dependent manner (IC<sub>50</sub>=18.05 μM).</td> </tr> </table> <p>Cell Viability Assay<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>HEL cells</td> </tr> <tr> <td>Concentration:</td> <td>12.5, 25, 50 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24, 48, 72 h</td> </tr> <tr> <td>Result:</td> <td>Inhibited viability of HEL cells in a concentration- and time-dependent manner.</td> </tr> </table> <p>Apoptosis Analysis<sup>[1]</sup></p>	Cell Line:	HEL cells (JAK2 <sup>V617F</sup> (+))	Concentration:	0-100 μM	Incubation Time:	48 h	Result:	Significantly inhibited proliferation in a concentration-dependent manner (IC <sub>50</sub> =18.05 μM).	Cell Line:	HEL cells	Concentration:	12.5, 25, 50 μM	Incubation Time:	24, 48, 72 h	Result:	Inhibited viability of HEL cells in a concentration- and time-dependent manner.
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Cell Line:	HEL cells (JAK2 <sup>V617F</sup> (+))
Concentration:	0-100 μM
Incubation Time:	24, 48, 72 h
Result:	Induced cell apoptosis in a concentration- and time-dependent manner.

#### Cell Cycle Analysis<sup>[1]</sup>

Cell Line:	HEL cells (JAK2 <sup>V617F</sup> (+))
Concentration:	0-100 μM
Incubation Time:	24, 48, 72 h
Result:	Significantly increased the number of HEL cells in the G2/M phase in a concentration-dependent manner.

#### In Vivo

ZT55 (100 mg/kg; p.o; once a day for 2 weeks) drastically attenuates the growth of subcutaneous HEL tumors in nude mice<sup>[1]</sup>

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Female, athymic BALB/c nude mice (6 to 8-week-old; JAK2 <sup>V617F</sup> xenograft model) <sup>[1]</sup> .
Dosage:	100 mg/kg
Administration:	Oral administration; once a day for 2 weeks
Result:	Induced marked reductions in tumor volume.

## REFERENCES

[1]. Hu M, et al. Discovery and evaluation of ZT55, a novel highly-selective tyrosine kinase inhibitor of JAK2<sup>V617F</sup> against myeloproliferative neoplasms. J Exp Clin Cancer Res. 2019 Feb 4;38(1):49.

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

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