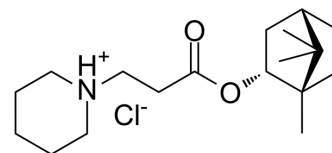


## As-358 hydrochloride

Cat. No.:	HY-146883A
CAS No.:	2374723-26-7
Molecular Formula:	C <sub>18</sub> H <sub>32</sub> ClNO <sub>2</sub>
Molecular Weight:	329.91
Target:	Filovirus
Pathway:	Anti-infection
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	As-358 (hydrochloride) has inhibitory effects against Ebola virus and Marburg virus with IC <sub>50</sub> s of 9.1 μM and 18.1 μM, as well as exhibits good in vivo safety <sup>[1]</sup> .								
<b>IC<sub>50</sub> &amp; Target</b>	IC <sub>50</sub> : 9.1 μM (Ebola virus), 18.1 μM (Marburg virus) <sup>[1]</sup>								
<b>In Vitro</b>	<p>As-358 (hydrochloride) (compound 3b-HCl) (0-500 μM) exhibits inhibitory activities against EBOV and MARV<sup>[1]</sup>. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Proliferation Assay</p> <table border="1"> <tr> <td>Cell Line:</td> <td>Vero (infected with EBOV and MARV)<sup>[1]</sup></td> </tr> <tr> <td>Concentration:</td> <td>0-500 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>10 days</td> </tr> <tr> <td>Result:</td> <td>Exhibits inhibitory activities against EBOV and MARV, with IC<sub>50</sub>s of 18.1 ± 9.3 μM and 9.1 ± 2.1 μM, SI values of 15 and 31, respectively.</td> </tr> </table>	Cell Line:	Vero (infected with EBOV and MARV) <sup>[1]</sup>	Concentration:	0-500 μM	Incubation Time:	10 days	Result:	Exhibits inhibitory activities against EBOV and MARV, with IC <sub>50</sub> s of 18.1 ± 9.3 μM and 9.1 ± 2.1 μM, SI values of 15 and 31, respectively.
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<b>In Vivo</b>	<p>As-358 (hydrochloride) (0.14-1.08 g/kg; IG; single) exhibits no toxicity to adult mice with intragastric administration<sup>[1]</sup>. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1"> <tr> <td>Animal Model:</td> <td>ICR mice<sup>[1]</sup></td> </tr> <tr> <td>Dosage:</td> <td>1.08, 0.44, 0.27, and 0.14 g/kg</td> </tr> <tr> <td>Administration:</td> <td>IG; single (observed for 7 days)</td> </tr> <tr> <td>Result:</td> <td>Exhibited no toxicity to adult mice with intragastric administration, and the LD<sub>50</sub> &gt; 1000 mg/kg.</td> </tr> </table>	Animal Model:	ICR mice <sup>[1]</sup>	Dosage:	1.08, 0.44, 0.27, and 0.14 g/kg	Administration:	IG; single (observed for 7 days)	Result:	Exhibited no toxicity to adult mice with intragastric administration, and the LD <sub>50</sub> > 1000 mg/kg.
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### REFERENCES

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[1]. Sokolova AS, Yarovaya OI, Zybkina AV, et al. Monoterpenoid-based inhibitors of filoviruses targeting the glycoprotein-mediated entry process. Eur J Med Chem. 2020;207:112726.

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

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