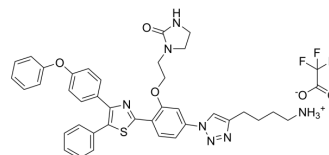


## Trypanothione synthetase-IN-1

Cat. No.:	HY-151148
CAS No.:	2355349-41-4
Molecular Formula:	C <sub>40</sub> H <sub>38</sub> F <sub>3</sub> N <sub>7</sub> O <sub>5</sub> S
Molecular Weight:	785.83
Target:	Parasite
Pathway:	Anti-infection
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

Description	Trypanothione synthetase-IN-1 (Compound 1) is a competitive Leishmania infantum trypanothione synthetase (TryS) inhibitor with an IC <sub>50</sub> of 14.8 μM when triamine spermidine is as polyamine S <sup>[1]</sup> .								
IC <sub>50</sub> & Target	Leishmania								
In Vitro	<p>Trypanothione synthetase-IN-1 (Compound 1) (0-75 μM; 24 or 72 h) shows leishmanicidal activity with cytotoxicity<sup>[1]</sup>. Trypanothione synthetase-IN-1 competes with ATP and the polyamine substrate for binding to LiTryS<sup>[1]</sup>. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Cytotoxicity Assay<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>L. infantum, HepG2</td> </tr> <tr> <td>Concentration:</td> <td>0-75 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h (L. infantum axenic amastigotes) or 72 h</td> </tr> <tr> <td>Result:</td> <td>Showed leishmanicidal activity with EC<sub>50</sub>s of 21.5 ± 2.4 μM and 13.5 ± 0.9 μM against axenic amastigotes and intracellular amastigotes, respectively. Showed cytotoxicity with a CC<sub>50</sub> of 15.9 ± 0.4 μM against HepG2 cells.</td> </tr> </table>	Cell Line:	L. infantum, HepG2	Concentration:	0-75 μM	Incubation Time:	24 h (L. infantum axenic amastigotes) or 72 h	Result:	Showed leishmanicidal activity with EC <sub>50</sub> s of 21.5 ± 2.4 μM and 13.5 ± 0.9 μM against axenic amastigotes and intracellular amastigotes, respectively. Showed cytotoxicity with a CC <sub>50</sub> of 15.9 ± 0.4 μM against HepG2 cells.
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### REFERENCES

[1]. Alcón-Calderón M, et al. Identification of *L. infantum* trypanothione synthetase inhibitors with leishmanicidal activity from a (non-biased) in-house chemical library. European Journal of Medicinal Chemistry, 2022: 114675.

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

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