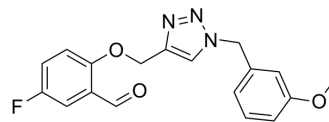


## Xanthine oxidase-IN-5

Cat. No.:	HY-144456
CAS No.:	2276711-87-4
Molecular Formula:	C <sub>18</sub> H <sub>16</sub> FN <sub>3</sub> O <sub>3</sub>
Molecular Weight:	341.34
Target:	Xanthine Oxidase
Pathway:	Metabolic Enzyme/Protease
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Xanthine oxidase-IN-5 is an effective and orally active xanthine oxidase (XO) inhibitor with IC <sub>50</sub> value of 0.70 μM. Xanthine oxidase-IN-5 displays favorable agent-like properties with ligand efficiency (LE) and lipophilic ligand efficiency (LLE) values of 0.33 and 3.41, respectively. Xanthine oxidase-IN-5 shows potent hypouricemic effects in hyperuricemic rat model <sup>[1]</sup> .								
<b>IC<sub>50</sub> &amp; Target</b>	IC <sub>50</sub> : 0.70 μM (XO) <sup>[1]</sup>								
<b>In Vivo</b>	<p>Xanthine oxidase-IN-5 (compound 9m) shows potent hypouricemic effects at an oral dose of 20 mg/kg in a rat hyperuricemia model induced by potassium oxonate<sup>[1]</sup>.</p> <p>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>Potassium oxonate-induced hyperuricemic rat model<sup>[1]</sup></td> </tr> <tr> <td>Dosage:</td> <td>20 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>PO; single (measured serum uric acid after 0.5 - 8 hours)</td> </tr> <tr> <td>Result:</td> <td>Showed potent hypouricemic effects at an oral dose of 20 mg/kg in a rat hyperuricemia model induced by potassium oxonate.</td> </tr> </table>	Animal Model:	Potassium oxonate-induced hyperuricemic rat model <sup>[1]</sup>	Dosage:	20 mg/kg	Administration:	PO; single (measured serum uric acid after 0.5 - 8 hours)	Result:	Showed potent hypouricemic effects at an oral dose of 20 mg/kg in a rat hyperuricemia model induced by potassium oxonate.
Animal Model:	Potassium oxonate-induced hyperuricemic rat model <sup>[1]</sup>								
Dosage:	20 mg/kg								
Administration:	PO; single (measured serum uric acid after 0.5 - 8 hours)								
Result:	Showed potent hypouricemic effects at an oral dose of 20 mg/kg in a rat hyperuricemia model induced by potassium oxonate.								

### REFERENCES

[1]. Zhang TJ, Zhang Y, Zhang ZH, et al. Discovery of 4-(phenoxyethyl)-1H-1,2,3-triazole derivatives as novel xanthine oxidase inhibitors. *Bioorg Med Chem Lett*. 2022;60:128582.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA