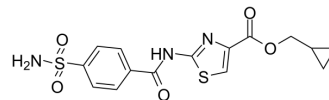


## hCAI/II-IN-4

<b>Cat. No.:</b>	HY-147925
<b>CAS No.:</b>	2480284-01-1
<b>Molecular Formula:</b>	C <sub>15</sub> H <sub>15</sub> N <sub>3</sub> O <sub>5</sub> S <sub>2</sub>
<b>Molecular Weight:</b>	381.43
<b>Target:</b>	Carbonic Anhydrase
<b>Pathway:</b>	Metabolic Enzyme/Protease
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	hCAI/II-IN-4 (compound 6d) is a potent dual hCA I/II inhibitor with K <sub>i</sub> values of 16.95, 15.22 and 27.04 nM for hCA I, hCA II and hCA $\alpha$ , respectively. hCAI/II-IN-4 has anti-hypoxia activities and low toxicity. hCAI/II-IN-4 can be used for acute mountain sickness (AMS) research <sup>[1]</sup> .									
<b>IC<sub>50</sub> &amp; Target</b>	K <sub>i</sub> :16.95 nM (hCA I) 15.22 nM (hCA II) and 27.04 nM (hCA $\alpha$ )									
<b>In Vitro</b>	<p>hCAI/II-IN-4 (compound 6d) (5-200<math>\mu</math>M; 48 hours) has no apparent cytotoxicity in HEK293 cells<sup>[1]</sup>.  MCE has not independently confirmed the accuracy of these methods. They are for reference only.  Cell Cytotoxicity Assay<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>HEK293 cells</td> </tr> <tr> <td>Concentration:</td> <td>5, 25, 50, 100 and 200 <math>\mu</math>M</td> </tr> <tr> <td>Incubation Time:</td> <td>48 hours</td> </tr> <tr> <td>Result:</td> <td>The cell viability rate was higher than 60%.</td> </tr> </table>		Cell Line:	HEK293 cells	Concentration:	5, 25, 50, 100 and 200 $\mu$ M	Incubation Time:	48 hours	Result:	The cell viability rate was higher than 60%.
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Concentration:	5, 25, 50, 100 and 200 $\mu$ M									
Incubation Time:	48 hours									
Result:	The cell viability rate was higher than 60%.									
<b>In Vivo</b>	<p>hCAI/II-IN-4 (compound 6d) (400-2000 mg/kg; p.o.; Male BLAB/c mice of hypoxia) has no apparent toxic effect in vivo<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>Male BLAB/c mice of hypoxia<sup>[1]</sup></td> </tr> <tr> <td>Dosage:</td> <td>400, 500 and 2000 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>Oral administration</td> </tr> <tr> <td>Result:</td> <td>Prolonged the survival time of mice by 29.3% compared with that of the blank control group.</td> </tr> </table>		Animal Model:	Male BLAB/c mice of hypoxia <sup>[1]</sup>	Dosage:	400, 500 and 2000 mg/kg	Administration:	Oral administration	Result:	Prolonged the survival time of mice by 29.3% compared with that of the blank control group.
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### REFERENCES

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[1]. Yang C, et al. N-Quinary heterocycle-4-sulphamoylbenzamides exert anti-hypoxic effects as dual inhibitors of carbonic anhydrases I/II. Bioorg Chem. 2020 Jul;100:103931.

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**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](mailto:tech@MedChemExpress.com)

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