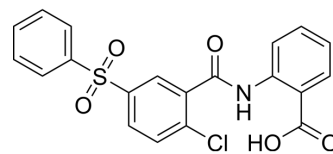


## BAY-8002

Cat. No.:	HY-122312
CAS No.:	724440-27-1
Molecular Formula:	C <sub>20</sub> H <sub>14</sub> ClNO <sub>5</sub> S
Molecular Weight:	415.85
Target:	Monocarboxylate Transporter
Pathway:	Membrane Transporter/Ion Channel
Storage:	Powder    -20°C    3 years 4°C        2 years In solvent   -80°C    6 months -20°C    1 month



### SOLVENT & SOLUBILITY

In Vitro	DMSO : 125 mg/mL (300.59 mM; Need ultrasonic)																					
	<table border="1"> <thead> <tr> <th rowspan="2">Solvent</th> <th rowspan="2">Mass</th> <th colspan="3">Concentration</th> </tr> <tr> <th>1 mg</th> <th>5 mg</th> <th>10 mg</th> </tr> </thead> <tbody> <tr> <td rowspan="3">Preparing Stock Solutions</td> <td>1 mM</td> <td>2.4047 mL</td> <td>12.0236 mL</td> <td>24.0471 mL</td> </tr> <tr> <td>5 mM</td> <td>0.4809 mL</td> <td>2.4047 mL</td> <td>4.8094 mL</td> </tr> <tr> <td>10 mM</td> <td>0.2405 mL</td> <td>1.2024 mL</td> <td>2.4047 mL</td> </tr> </tbody> </table>	Solvent	Mass	Concentration			1 mg	5 mg	10 mg	Preparing Stock Solutions	1 mM	2.4047 mL	12.0236 mL	24.0471 mL	5 mM	0.4809 mL	2.4047 mL	4.8094 mL	10 mM	0.2405 mL	1.2024 mL	2.4047 mL
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Please refer to the solubility information to select the appropriate solvent.																						
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.08 mg/mL (5.00 mM); Clear solution																					
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: 2.08 mg/mL (5.00 mM); Suspended solution; Need ultrasonic																					
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (5.00 mM); Clear solution																					

### BIOLOGICAL ACTIVITY

Description	BAY-8002 is a potent, selective, orally active inhibitor of monocarboxylate transporter 1 (MCT1), with an IC <sub>50</sub> of 85 nM in the MCT1-expressing DLD-1 cells, displays excellent selectivity against MCT4. Anti-tumor activity <sup>[1]</sup> .
IC <sub>50</sub> & Target	MCT1
In Vitro	BAY-8002 is an inhibitor of MCT1, with an IC <sub>50</sub> of 85 nM in the MCT1-expressing DLD-1 cells, displays excellent selectivity against MCT4 (IC <sub>50</sub> >50 μM in EVSA-T cells) <sup>[1]</sup> .

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

#### In Vivo

BAY-8002 (80 and 160 mg/kg, p.o., twice daily, for more than 26 days) significantly inhibits tumor growth in Raji tumor-bearing mice<sup>[1]</sup>.

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

Animal Model:	Female NOD SCID mice bearing Raji cells (7-10 weeks old) <sup>[1]</sup>
Dosage:	80 and 160 mg/kg
Administration:	P.O. twice daily for more than 26 days
Result:	Inhibited tumor growth with no obvious body weight loss, but showed no effect on tumor regression.

## CUSTOMER VALIDATION

- Cell. 2021 Jan 21;184(2):370-383.e13.
- Blood. 2021 Oct 5;blood.2021011563.
- Crit Rev Anal Chem. 2021 Mar 10;1-15.

See more customer validations on [www.MedChemExpress.com](http://www.MedChemExpress.com)

## REFERENCES

[1]. Quanz M, et al. Preclinical Efficacy of the Novel Monocarboxylate Transporter 1 Inhibitor BAY-8002 and Associated Markers of Resistance. Mol Cancer Ther. 2018 Nov;17(11):2285-2296.

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

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