

## Saccharopine hydrochloride

Cat. No.: HY-W040307B Molecular Formula:  $C_{11}H_{21}CIN_2O_6$ 

Molecular Weight: 313

Target: **Endogenous Metabolite** Pathway: Metabolic Enzyme/Protease

Storage: -20°C, sealed storage, away from moisture and light

\* In solvent: -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture

and light)

$$HO$$
 $NH_2$ 
 $OOH$ 
 $OOH$ 
 $OOH$ 

**Product** Data Sheet

## **SOLVENT & SOLUBILITY**

In Vitro

H<sub>2</sub>O: 250 mg/mL (798.72 mM; Need ultrasonic)

| Preparing<br>Stock Solutions | Solvent Mass<br>Concentration | 1 mg      | 5 mg       | 10 mg      |
|------------------------------|-------------------------------|-----------|------------|------------|
|                              | 1 mM                          | 3.1949 mL | 15.9744 mL | 31.9489 mL |
|                              | 5 mM                          | 0.6390 mL | 3.1949 mL  | 6.3898 mL  |
|                              | 10 mM                         | 0.3195 mL | 1.5974 mL  | 3.1949 mL  |

Please refer to the solubility information to select the appropriate solvent.

| DI | $\cap$ | <b>OCI</b> | CAI | ۸CT    | IVITY     |
|----|--------|------------|-----|--------|-----------|
| DI | UL     | UUI        | CAL | _ AC I | 1 1 1 1 1 |

| Description               | Saccharopine (L-Saccharopine) hydrochloride, a lysine degradation intermediate, is a mitochondrial toxin. Lysine and $\alpha$ -ketoglutarate are converted into Saccharopine hydrochloride by the lysine-ketoglutarate reductase. Saccharopine hydrochloride is then oxidized to $\alpha$ -aminoapidate semialdehyde and glutamate by the saccharopine dehydrogenase. Saccharopine hydrochloride impairs development by disrupting mitochondrial homeostasis [1][2][3]. |
|---------------------------|---|
| IC <sub>50</sub> & Target | Human Endogenous Metabolite   |
| In Vitro                  | Saccharopine accumulation leads to mitochondrial damage and functional loss which is induced by saccharopine dehydrogenase (SDH) mutations of $\alpha$ -aminoadipate semialdehyde synthase (AASS)-1 in C. elegans <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.  |
| In Vivo                   | Saccharopine accumulation induces mitochondrial damage and progressive postnatal growth retardation in Aass mutant $mice^{[1]}$ .  MCE has not independently confirmed the accuracy of these methods. They are for reference only.  |

## **REFERENCES**

- [1]. Zhou J, et, al. The lysine catabolite saccharopine impairs development by disrupting mitochondrial homeostasis. J Cell Biol. 2019 Feb 4;218(2):580-597.
- [2]. Leandro J, et, al. Saccharopine, a lysine degradation intermediate, is a mitochondrial toxin. J Cell Biol. 2019 Feb 4;218(2):391-392.
- [3]. Papes F, et, al. Lysine degradation through the saccharopine pathway in mammals: involvement of both bifunctional and monofunctional lysine-degrading enzymes in mouse. Biochem J. 1999 Dec 1;344 Pt 2(Pt 2):555-63.

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

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