## Stilbamidine

| Cat. No.: | HY-U00007 |
| :--- | :--- |
| CAS No.: | $122-06-5$ |
| Molecular Formula: | $\mathrm{C}_{16} \mathrm{H}_{16} \mathrm{~N}_{4}$ |
| Molecular Weight: | 264.33 |
| Target: | Fungal |
| Pathway: | Anti-infection |
| Storage: | Please store the product under the recommended conditions in the Certificate of |
|  | Analysis. |



## BIOLOGICAL ACTIVITY

## Description

$I_{50}$ \& Target fungal

In Vitro
fungal

Stilbamidine is a diamidine compound derived from Stilbene and used chiefly in the form of its crystalline isethionate salt in treating various fungal infections.

The high-affinity pentamidine transporter (HAPT1) is inhibited by Propamidine but displays only low affinity to Stilbamidine. adenosine-sensitive pentamidine transporter (ASPT1), in contrast, is strongly inhibited by Stilbamidine, and Propamidine. [3 H ]pentamidine uptake is determined in the presence of various concentrations of adenosine ( $\mathrm{IC}_{50}=1.2 \mu \mathrm{M}$ ) or melarsen oxide ( $\mathrm{IC}_{50}=0.7 \mu \mathrm{M}$ ), as well as in the presence of $250 \mu \mathrm{M}$ adenosine and increasing concentrations of hypoxanthine, Propamidine $\left(\mathrm{IC}_{50}=6.1 \mu \mathrm{M}\right)$ and Stilbamidine $\left(\mathrm{IC}_{50}=110 \mu \mathrm{M}\right)^{[1]}$. The two diamidine compounds, Stilbamidine and Pentamidine are used to treat in multiple myeloma, a disease in which increase of the globulin content of the serum is of frequent occurrence ${ }^{[2]}$
MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## REFERENCES

[1]. De Koning HP. Uptake of pentamidine in Trypanosoma brucei brucei is mediated by three distinct transporters: implications for cross-resistance with arsenicals. Mol Pharmacol. 2001 Mar;59(3):586-92.
[2]. BREWER AE.et al. Multiple myeloma treated with stilbamidine and pentamidine. Br Med J. 1948 Dec 4;2(4587):978-82.

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