



## **Pafuramidine**

Catalog No: tcsc1470

Available Sizes
Size: 5mg
Size: 10mg
Size: 50mg
Size: 100mg
Specifications
<b>CAS No:</b> 186953-56-0
<b>Formula:</b> $C_{20}^{H}_{20}^{N}_{4}^{O}_{3}$
Pathway: Anti-infection
<b>Target:</b> Parasite
Purity / Grade: >98%
Solubility: H2O:
Alternative Names: DB289
Observed Molecular Weight: 364.4





## **Product Description**

Pafuramidine (DB289) is an orally bioavailable prodrug of furamidine, which has clinical activity against Pneumocystis pneumonia.

IC50 Value: 4.5 nM (In vitro inhibitory activity against Trypanosoma brucei rhodesiense) [4]

Target: Antiparasitic

DB289 (pafuramidine maleate; 2,5-bis[4-(N-methoxyamidino)phenyl]furan monomaleate) is a prodrug of DB75 (furamidine dihydrochloride; 2,5-bis(4-guanylphenyl)furan dihydrochloride), an aromatic dication related to pentamidine that has demonstrated good efficacy against African trypanosomiasis, Pneumocystis carinii pneumonia, and malaria, but lacks adequate oral availability.

in vitro: The results of this investigation suggest that DB75 inhibits mitochondrial function. Yeast cells relying upon mitochondrial metabolism for energy production are especially sensitive to DB75 [1].

in vivo: Clearance of DB289 approximated the liver plasma flow and its large volume of distribution was consistent with extensive tissue binding. Plasma protein binding of DB289 was 97 to 99% in four animal species and humans, but that of DB75 was noticeably less and more species- and concentration-dependent [2]. Despite excellent oral activity against early-stage sleeping sickness, oral administration of DB289 exhibited limited efficacy in mouse models of late-stage disease [3].

Clinical trial: DB289, a novel orally active prodrug of DB75, is undergoing phase IIb clinical trials for early-stage human African trypanosomiasis, Pneumocystis jiroveci carinii pneumonia, and malaria [1].

All products are for RESEARCH USE ONLY. Not for diagnostic & therapeutic purposes!