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ProductInformation

Sulconazole nitrate salt

Product Number **S 9632** Store at Room Temperature

Product Description

Molecular Formula: C₁₈H₁₅Cl₃N₂S • HNO₃

Molecular Weight: 460.8 CAS Number: 61318-91-0 Melting Point: 130.5 - 132 °C¹

Synonym: 1-[2-[[(4-chlorophenyl)methyl]-thio]-2-(2,4-dichlorophyenyl)ethyl]-1H-imidazole nitrate;

 $(\pm)-1-[2,4-dichloro-\beta-[(p-$

chlorobenzyl)thio]phenethyl]imidazole nitrate

Sulconazole is an imidazole derivative with antifungal and antibacterial activity. It has activity against *Malassezia furfur* and *Candida albicans*, as well as dermatophytes and Gram-positive bacteria.² A study of the action of sulconazole and other antifungal agents against various *Candida albicans* and other *Candida* isolates has been reported.³ The comparative activity of sulconazole against *Candida albicans* in logarithmic phase (2 μM) and stationary phase (8 μM) has been investigated.⁴

A study of the formation of aggregates of sulconazole in solution and their inhibitory activity against the enzymes, β -lactamase, chymotrypsin, and malate dehydrogenase, has been described. The inhibition of cytochromes P450 in cDNA-expressing microsomes from human lymphoblast cells or human liver microsomes by sulconazole and other antifungal compounds has been examined.

An HPLC method for the analysis of sulconazole in plasma has been described. HPLC assays for the resolution of the optical isomers of sulconazole that use chiral amylose columns and chiral cellulose columns have been reported. 8,9

Precautions and Disclaimer

For Laboratory Use Only. Not for drug, household or other uses.

Preparation Instructions

This product is soluble in water (50 mg/ml), with heat and sonication as needed, yielding a clear, colorless solution.

References

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- Hernandez Molina, J. M., et al., In vitro activity of cloconazole, sulconazole, butoconazole, isoconazole, fenticonazole, and five other antifungal agents against clinical isolates of Candida albicans and Candida spp. Mycopathologia, 118(1), 15-21 (1992).
- 4. Beggs, W. H., Influence of growth phase on the susceptibility of *Candida albicans* to butoconazole, oxiconazole, and sulconazole. J. Antimicrob. Chemother., **16(3)**, 397-399 (1985).
- 5. Seidler, J., et al., Identification and prediction of promiscuous aggregating inhibitors among known drugs. J. Med. Chem., **46(21)**, 4477-4486 (2003).
- 6. Zhang, W., et al., Inhibition of cytochromes P450 by antifungal imidazole derivatives. Drug Metab. Dispos.. **30(3)**. 314-318 (2002).
- 7. Fass, M., et al., Reversed-phase high-pressure liquid chromatographic analysis of sulconazole in plasma. J. Pharm. Sci., **70(12)**, 1338-1340 (1981).

- 8. Aboul-Enein, H. Y., and Ali, I., Comparison of the chiral resolution of econazole, miconazole, and sulconazole by HPLC using normal-phase amylose CSPs. Fresenius J. Anal. Chem., **370(7)**, 951-955 (2001).
- Aboul-Enein, H. Y., and Ali, I., Comparative study of the enantiomeric resolution of chiral antifungal drugs econazole, miconazole and sulconazole by HPLC on various cellulose chiral columns in normal phase mode. J. Pharm. Biomed. Anal., 27(3-4), 441-446 (2002).

GCY/NSB 1/04