



# LKT Laboratories, Inc.

## Diaveridine Hydrochloride

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### Product Information

**Product ID** D3302

**CAS No.** 2507-23-5

**Chemical Name** 5-[(3,4-Dimethoxyphenyl)methyl]-2,4-pyrimidine- diamine HCl

**Synonym**

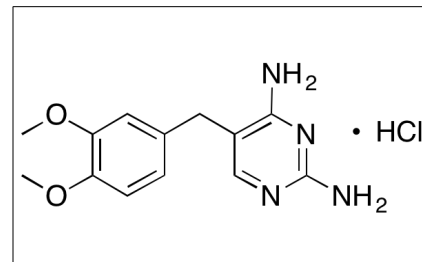
**Formula**  $C_{13}H_{16}N_4O_2 \cdot HCl$

**Formula Wt.** 296.74

**Melting Point**

**Purity**  $\geq 98\%$

**Solubility** Soluble in water.



### Pricing and Availability

**Bulk quantities available upon request**

Product ID	Size	List Price
D3302	1 g	\$81.70
D3302	10 g	\$428.70

**Store Temp** Ambient

**Ship Temp** Ambient

**Description** Diaveridine is a coccidiostat used in veterinary medicine that exhibits anti-parasitic activity. Diaveridine inhibits dihydrofolate reductase, preventing folic acid synthesis in species of *Pneumocystis*. Diaveridine is also genotoxic, inducing structural chromosomal aberrations and inducing DNA damage in vitro.

**References** Ono T, Sekiya T, Takahashi Y, et al. The genotoxicity of diaveridine and trimethoprim. *Environ Toxicol Pharmacol*. 1997 Sep;3(4):297-306. PMID: 21781790.

Cirioni O, Giacometti A, Scalise G. In-vitro activity of atovaquone, sulphamethoxazole and dapsone alone and combined with inhibitors of dihydrofolate reductase and macrolides against *Pneumocystis carinii*. *J Antimicrob Chemother*. 1997 Jan;39(1):45-51. PMID: 9044027.

Walzer PD, Kim CK, Foy JM, et al. Inhibitors of folic acid synthesis in the treatment of experimental *Pneumocystis carinii* pneumonia. *Antimicrob Agents Chemother*. 1988 Jan;32(1):96-103. PMID: 3258144.

**Caution:** This product is intended for laboratory and research use only. It is not for human or drug use.