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Product Information

Product ID D1995

CAS No. 149003-01-0

Chemical Name

Synonym

Formula C₁₁H₁₆N₄O₄ • HCl

Formula Wt. 304.73 Melting Point 191-197°C

Purity ≥98%

Solubility Sparingly soluble in water and 0.1 N HCl, slightly soluble in ethanol and

methanol, and practically insoluble in nonpolar organic solvents.

Pricing and Availability

Bulk quanitites available upon request

Product ID	Size	List Price
D1995	5 mg	\$78.80
D1995	25 mg	\$225.80
D1995	100 mg	\$614.30

Store Temp Ambient Ship Temp Ambient

Description Dexrazoxane is an iron chelator that exhibits cardioprotective, antioxidative, pro-angiogenic, and anti-parasitic activities.

Dexrazoxane decreases superoxide formation and prevents anthracycline-induced cardiotoxicity. In animal models of myocardial infarction, dexrazoxane decreases infarct size, increases capillary density, and improves cardiac function.

Additionally, this compound displays antimalarial benefit, inhibiting growth of Plasmodium.

References Doroshow JH. Dexrazoxane for the prevention of cardiac toxicity and treatment of extravasation injury from the anthracycline antibiotics. Curr Pharm Biotechnol. 2012 Aug;13(10):1949-56. PMID: 22352729.

> Zhou L, Sung RY, Li K, et al. Cardioprotective effect of dexrazoxane in a rat model of myocardial infarction: anti-apoptosis and promoting angiogenesis. Int J Cardiol. 2011 Oct 20;152(2):196-201. PMID: 20692056.

Jones RL. Utility of dexrazoxane for the reduction of anthracycline-induced cardiotoxicity. Expert Rev Cardiovasc Ther. 2008 Nov;6(10):1311-7. PMID: 19018683.

Lovevsky M. Sacci JB Jr, Boehme P, et al. Plasmodium falciparum and Plasmodium yoelii: effect of the iron chelation prodrug dexrazoxane on in vitro cultures. Exp Parasitol. 1999 Feb;91(2):105-14. PMID: 9990337.

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