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# **ProductInformation**

## (±)-Metoprolol (+)-tartrate salt

Product Number **M 5391**Store at Room Temperature

## **Product Description**

Molecular Formula: (C<sub>15</sub>H<sub>25</sub>NO<sub>3</sub>)<sub>2</sub> • C<sub>4</sub>H<sub>6</sub>O<sub>6</sub>

Molecular Weight: 684.8 CAS Number: 56392-17-7

 $\lambda_{\text{max}}$ : 223 nm  $(H_2O)^1$ 

Extinction Coefficient: E<sup>mM</sup> = 23.4 <sup>1</sup>

Specific Rotation:  $+6.5^{\circ}$  to  $10.5^{\circ}$  (c = 20 mg/ml,  $H_2O$ ,

20 °C)

Synonyms: 1-[4-(2-methoxyethyl)phenoxy]-3 -[(1-methylethyl)amino]2-propanol tartrate; (±) -1-(isopropylamino)-3-[p-(β-methoxyethyl)phenoxy]

-2-propanol tartrate1

Metoprolol is a cardioselective  $\beta$ -adrenergic receptor antagonist that is used in hypertension and cardiovascular research. Metoprolol can also interact with H1 histamine receptor antagonists such as diphenhydramine. A review of the pharmacodynamics and pharmacokinetics of metoprolol formulations has been reported.

A permeability study of a serum culture system for Caco-2 monolayers has been tested with metoprolol. Metoprolol has also been used to probe the TR146 cell culture model as an *in vitro* model of human buccal epithelium. Pretreatment of cultured human umbilical vein endothelial cells with metoprolol (0.1 - 100  $\mu$ M) has been shown to diminish endothelin-1 production.  $^8$ 

An HPLC method that combines UV and fluorescence detection has been developed for the analysis of metoprolol and other beta blocker compounds. 9

## **Precautions and Disclaimer**

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

## **Preparation Instructions**

The product is soluble in water (50 mg/ml), with heat as needed, yielding a clear, colorless solution. The solubility in water has also been reported at >1000 mg/ml. It is also soluble in methanol (>500 mg/ml) and chloroform (496 mg/ml).<sup>1</sup>

#### References

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- 8. Garlichs, C. D., et al., Beta-blockers reduce the release and synthesis of endothelin-1 in human endothelial cells. Eur. J. Clin. Invest., **29(1)**, 12-16 (1999).
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