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ProductInformation

DL-Isoproterenol hydrochloride

Product Number **I5627** Store at Room Temperature

Product Description

Molecular Formula: C₁₁H₁₇NO₃ • HCl

Molecular Weight: 247.7 CAS Number: 51-30-9 Melting Point: 170-171 °C¹

Synonyms: isoprenaline hydrochloride; 4-[1-hydroxy-

2-(1-methylethyl)-amini]ethyl]-1,2-benzenediol

hydrochloride; 3,4-dihydroxy-

α-[(isopropylamino)methyl]benzyl alcohol

hydrochloride; α-isopropylaminomethyl)protocatechuyl

alcohol hydrochloride

Isoproterenol is a cathecholamine compound and β -adrenergic agonist that is used in cardiovascular research. Isoproterenol can be used to probe the role of β -adrenergic receptors in such processes as N-acetyltransferase activity and the production of melatonin. The effects of isoproterenol and other catecholamines on angiotensinogen gene expression in opossum kidney proximal tubular cells have been studied.

Isoproterenol also causes increases in cytosolic cAMP levels. Isoproterenol (100 $\mu\text{M})$ has been used to study the role of the anti-apoptotic protein Bcl-2 on the growth and death of cultured murine S49 cells. A study of cultured human fetal lung fibroblasts has shown that isoproterenol (10 $\mu\text{M})$ can inhibit fibroblast chemotaxis. The ability of isoproterenol (10 $\mu\text{M})$ to modulate binding of the protein PSD-95 to the μ_1 -adrenergic receptor in COS-7 cells has been investigated.

Precautions and Disclaimer

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

Preparation Instructions

This product is soluble in water at (50 mg/ml), yielding a clear to slightly hazy, colorless to yellow or faint pink solution. The solubility in water has also been reported at 333 mg/ml. It is soluble in ethanol at 20 mg/ml. ¹

Storage/Stability

Solutions of this product become pink to brownish-pink upon exposure to air. Solutions will almost immediately discolor colors when made alkaline. Stock solutions should be protected from light.

References

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- 8. Zhang, L., and Insel, P. A., Bcl-2 protects lymphoma cells from apoptosis but not growth arrest promoted by cAMP and dexamethasone. Am. J. Physiol. Cell. Physiol., **281(5)**, C1642-1647 (2001).
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