PRODUCT INFORMATION



Salbutamol-d_o

Item No. 25234

CAS Registry No.:	1173021-73-2		
Formal Name:	α^{1} -[[[1,1-di(methyl-d ₃)ethyl-2,2,2-d ₃]amino]		
	methyl]-4-hydroxy-1,3-benzenedimethanol		
Synonyms:	(±)-Albuterol-d ₉ , (±)-Salbutamol-d ₉ ,	(
	DL-Salbutamol-d ₉		
MF:	$C_{13}H_{12}D_9NO_3$	но	Ň D
FW:	248.4		Ň.
Chemical Purity:	≥95% (Salbutamol)		
Deuterium		но	
Incorporation:	≥99% deuterated forms (d ₁ -d ₉); ≤1% d ₀		
Supplied as:	A solid		
Storage:	-20°C		
Stability:	≥4 years		
Information represents the product specifications. Patch specific analytical results are provided on each cartificate of analysis			

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Laboratory Procedures

Salbutamol-d₉ is intended for use as an internal standard for the quantification of salbutamol (Item No. 21003) by GC- or LC-MS. The accuracy of the sample weight in this vial is between 5% over and 2% under the amount shown on the vial. If better precision is required, the deuterated standard should be quantitated against a more precisely weighed unlabeled standard by constructing a standard curve of peak intensity ratios (deuterated versus unlabeled).

Salbutamol-d_o is supplied as a solid. A stock solution may be made by dissolving the salbutamol-d_o in the solvent of choice, which should be purged with an inert gas. Salbutamol-do is slightly soluble in methanol.

Description

Salbutamol is an agonist of the β_2 -adrenergic receptor (β_2 -AR).¹ It induces bronchodilation in isolated guinea pig trachea rings (EC₅₀ = 19[°] nM). Salbutamol selectively binds to the β_2 -AR over the β_1 -AR and β_3 -AR in CHO cells expressing the human receptors (K_ds = 0.76, 21, and 46 μ M, respectively).² Salbutamol (25 and 50 µg/kg, i.v.) reduces acetylcholine-induced bronchospasm in anesthetized guinea pigs.³ It also reduces the response of bronchial muscle to efferent vagal stimulation in anesthetized cats and dogs. Nebulized salbutamol reduces transpulmonary pressure in recurrent airway obstruction-affected horses $(ED_{50} = 43.6 \ \mu g/animal)$ ⁴ Formulations containing salbutamol have been used in the prevention of exercise-induced asthma and the prevention or treatment of chronic obstructive pulmonary disease (COPD).

References

- 1. Kern, C., Meyer, T., Droux, S., et al. Synthesis and pharmacological characterization of β_2 -adrenergic agonist enantiomers: Zilpaterol. J. Med. Chem. 52(6), 1773-1777 (2009).
- Baker, J.G. The selectivity of β -adrenoceptor antagonists at the human β_1 , β_2 and β_3 adrenoceptors. 2. Br. J. Pharmacol. 144(3), 317-322 (2005).
- 2. Cullum, V.A., Farmer, J.B., Jack, D., et al. Salbutamol: A new, selective β-adrenoceptive receptor stimulant. Br. J. Pharmacol. 35(1), 141-151 (1969).
- 3. Arroyo, M.G., Couëtil, L.L., Nogradi, N., et al. Efficacy of inhaled levalbuterol compared to albuterol in horses with recurrent airway obstruction. J. Vet. Intern. Med. 30(4), 1333-1337 (2016).

WARNING THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

SAFETY DATA

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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