

Product Data Sheet

Mabuterol

 Cat. No.:
 HY-13338

 CAS No.:
 56341-08-3

 Molecular Formula:
 C₁₃H₁₈ClF₃N₂O

Molecular Weight: 310.74

Target: Adrenergic Receptor

Pathway: GPCR/G Protein; Neuronal Signaling

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.

BIOLOGICAL ACTIVITY

Description	Mabuterol is a selective and orally active beta-2 adrenergic receptor (ADRB2) agonist. Mabuterol inhibits the proliferation and suppresses the increase of intracellular Ca2+ induced by PDGF-BB. Mabuterol suppresses the protein expressions of Drp-1, cyclinD1 and PCNA and enhanced the expression of Mfn-2 induced by PDGF-BB ^{[1][2]} .	
IC ₅₀ & Target	Beta-2 adrenergic receptor	
In Vitro	Mabuterol (10-60 μ M; 48 h) inhibits the proliferation of ASMCs induced by PDGF-BB ^[2] . Mabuterol (50 μ M; 1h) suppresses the increase of intracellular Ca2+ induced by PDGF-BB (25 ng/ml) ^[2] . Mabuterol (50 μ M; 48 h) significantly suppresses the protein expressions of Drp-1, cyclinD1 and PCNA and enhanced the expression of Mfn-2 induced by PDGF-BB ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Proliferation Assay ^[2]	
	Cell Line:	rat ASMCs ^[2]
	Concentration:	10, 20, 30, 40, 50, 60 μΜ
	Incubation Time:	24, 48 h
	Result:	Inhibited the proliferation of ASMCs induced by PDGF-BB (25 ng/ml) in a dose-dependent maner at 48 h.

REFERENCES

 $[1]. Yamamoto\ I, et\ al.\ Enzyme\ immunoassay.\ 1985; 6(3): 261-76.$

[2]. Yaru Gu, et al. Inhibitory effect of mabuterol on proliferation of rat ASMCs induced by PDGF-BB via regulating [Ca2+]i and mitochondrial fission/fusion. Interactions, 2019, 307: 63-72.

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$

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