## **Product** Data Sheet

## Reproterol

 $\begin{array}{lll} \textbf{Cat. No.:} & \text{HY-135490} \\ \textbf{CAS No.:} & 54063\text{-}54\text{-}6 \\ \textbf{Molecular Formula:} & \text{$C_{18}$H}_{23}$N_{5}$O_{5} \\ \end{array}$ 

Molecular Weight: 389.41

Target: Adrenergic Receptor; Phosphodiesterase (PDE)

Pathway: GPCR/G Protein; Neuronal Signaling; Metabolic Enzyme/Protease

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

Analysis.

## **BIOLOGICAL ACTIVITY**

Description	Reproterol is a dual acting $\beta$ 2-adrenoceptor agonist and PDE inhibitor. The theophylline constituent of Reproterol inhibits phosphodiesterase activity induced by adenylyl cyclase. Reproterol has the potential for asthma research <sup>[1][2]</sup> .
IC <sub>50</sub> & Target	β2 adrenoceptor

## **REFERENCES**

[1]. Uwe R Juergens, et al. Different mechanisms of action of beta2-adrenergic receptor agonists: a comparison of reproterol, fenoterol and salbutamol on monocyte cyclic-AMP and leukotriene B4 production in vitro. Eur J Med Res. 2004 Jul 30;9(7):365-70.

[2]. C G Georgakopoulos, et al. Excretion study of the beta2-agonist reproterol in human urine. J Chromatogr B Biomed Sci Appl. 1999 Apr 16;726(1-2):141-8.

Caution: Product has not been fully validated for medical applications. For research use only.

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