

# (S) - (+) -Rolipram

## Catalog No: tcsc2478

Available Sizes

Size: 10mg

Size: 50mg

Size: 100mg

Specifications

CAS No:

85416-73-5

Formula:

C<sub>16</sub>H<sub>21</sub>NO<sub>3</sub>

**Pathway:** Metabolic Enzyme/Protease

**Target:** Phosphodiesterase (PDE)

**Purity / Grade:** 

## Solubility:

10 mM in DMSO

#### **Alternative Names:**

(+)-Rolipram;(S)-Rolipram

#### **Observed Molecular Weight:**

275.34

### **Product Description**

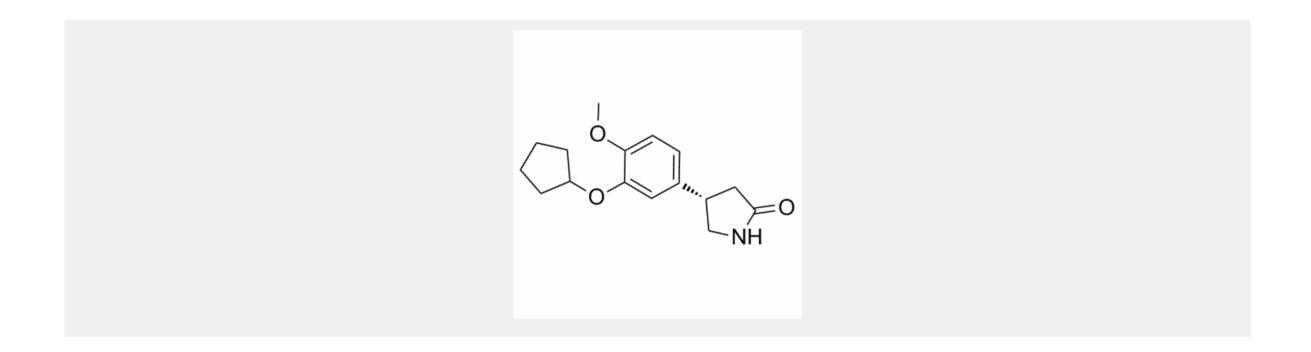
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(S)-(+)-Rolipram is a PDE4-inhibitor and an anti-inflammatory agent, less potent than its R enantiomer.

#### Target: PDE4B; PDE4D

Rolipram, a selective inhibitor of the cyclic AMP-specific phosphodiesterase (PDE IV). Rolipram did not inhibit 5-lipoxygenase activity but did inhibit human monocyte production of leukotriene B4 (LTB4, IC50 3.5 microM). Rolipram inhibited arachidonic acid-induced inflammation in the mouse, while the low Km-cyclic-GMP PDE inhibitor. Rolipram had a modest effect on LTB4 production in the mouse, but markedly reduced LTB4-induced PMN infiltration [1]. In humans and animals rolipram produces thereby a variety of biological effects. These effects include attenuation of endogenous depression and inflammation in the central nervous system (CNS), both effects are of potential clinical relevance [2].



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