Product Data Sheet

COX-2-IN-1

Cat. No.: HY-U00275
CAS No.: 787623-48-7

 $\label{eq:molecular-formula:} \textbf{Molecular Formula:} \qquad \textbf{C}_{18}\textbf{H}_{14}\textbf{ClF}_{3}\textbf{N}_{4}\textbf{O}_{2}\textbf{S}$

Molecular Weight: 442.84

Target: COX

Pathway: Immunology/Inflammation

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

Analysis.

BIOLOGICAL ACTIVITY

Description	COX-2-IN-1 is potent and slective COX-2 inhibitor with an IC $_{50}$ of 3.9 $\mu\text{M}.$
IC ₅₀ & Target	COX-2 3.9 μM (IC ₅₀)
In Vitro	COX-2-IN-1(compound 5f) is an inhibitor of COX-1 and COX-2 with IC $_{50}$ s of >100 and 3.9 μ M, respectively. Comparison of 5b and COX-2-IN-1 shows that smaller halogen atom at 5th position on the indole ring enhances COX-2 enzyme inhibition activity, whereas the presence of a larger bromine atom at 6th position of the ring improves the inhibitory activity (5d, 5e and 5h) of the compound ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Reddy MV, et al. Design, synthesis, and biological evaluation of 1-(4-sulfamylphenyl)-3-trifluoromethyl-5-indolyl pyrazolines as cyclooxygenase-2 (COX-2) and lipoxygenase (LOX) inhibitors. Bioorg Med Chem. 2008 Apr 1;16(7):3907-16.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA