Product Data Sheet

PF-9184

Cat. No.: HY-19622 CAS No.: 1221971-47-6

 $\label{eq:molecular-formula:} \textbf{Molecular Formula:} \qquad \textbf{C}_{21}\textbf{H}_{14}\textbf{C}\textbf{I}_{2}\textbf{N}_{2}\textbf{O}_{4}\textbf{S}$

Molecular Weight: 461.32

Target: PGE synthase

Pathway: Immunology/Inflammation

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

Analysis.

BIOLOGICAL ACTIVITY

Description	PF-9184 is a potent and highly selective inhibitor of human microsomal prostaglandin E synthase-1 (mPGES-1), with an IC ₅₀ of 16.5 nM. PF-9184 inhibits IL-1 β -induced PGE ₂ synthesis in vitro ^[1] .
IC ₅₀ & Target	IC50: 16.5 nM (mPGES-1) ^[1]
In Vitro	PF-9184 (0.015-100 μ M; 24 hours) inhibits IL-1 β -stimulated prostaglandin E $_2$ (PGE $_2$) synthesis in rheumatoid arthritis (RASF) in synovial fibroblasts derived from patients and has no apparent cytotoxic effects up to 100 μ M[1]. PF-9184 (0.015-100 μ M) potently blocks mPGES-1 ability to synthesize PGE $_2$ from PGH $_2$, and with no apparent inhibitory effects on COX-2 and prostacyclin synthase in cells[1]. PF-9184 (0.015-100 μ M) inhibits PGE $_2$ weakly but has no effect on TXB2 synthesis except at 100 μ M in human whole blood and modified blood assays[1]. PF-9184 is a poor inhibitor of recombinant rat mPGES-1 (IC $_{50}$ =1080±398 nM)[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	PF-9184 (oral administration or even local delivery) has no effect on PGE_2 synthesis in recombinant rats ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Gabriel M, et, al. Distinction of microsomal prostaglandin E synthase-1 (mPGES-1) inhibition from cyclooxygenase-2 inhibition in cells using a novel, selective mPGES-1 inhibitor. Biochem Pharmacol. 2010 May 15; 79(10): 1445-54.

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

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