Indomethacin

Small Molecules

Inhibits COX-1 and COX-2



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TOLL FREE PHONE 1 800 667 0322 • PHONE +1 604 877 0713 INFO@STEMCELL.COM • TECHSUPPORT@STEMCELL.COM FOR GLOBAL CONTACT DETAILS VISIT OUR WEBSITE

Catalog # 73942 1 g

Product Description

Indomethacin is a non-steroidal, anti-inflammatory agent that inhibits cyclooxygenase (COX) activity, thereby blocking the production of prostaglandins (Vane et al.). Indomethacin inhibits both COX-1 and COX-2 ($IC_{50} = 0.08$ and 0.96 μ M for recombinant human COX-1 and COX-2, respectively; Kurumbail et al.).

Molecular Name: Indomethacin
Alternative Names: Indocin
CAS Number: 53-86-1
Chemical Formula: C₁₉H₁₆CINO₄
Molecular Weight: 357.8 g/mol

≥ 99%

Chemical Name: 2-[1-(4-chlorobenzoyl)-5-methoxy-2-methylindol-3-yl]acetic acid

Structure:

Purity:

$$HO_2C$$
 N
 CI

Properties

Physical Appearance: A crystalline solid

Storage: Product stable at room temperature (15 - 25°C) as supplied. Protect product from prolonged exposure to light.

For long-term storage store with a desiccant.

Stable as supplied for 12 months from date of receipt.

Solubility: \cdot DMSO \leq 45 mM

· Absolute ethanol \leq 15 mM

For example, to prepare a 10 mM stock solution in DMSO, resuspend 10 mg in 2.79 mL of DMSO.

Prepare stock solution fresh before use. Information regarding stability of small molecules in solution has rarely been reported, however, as a general guide we recommend storage in DMSO at -20°C. Aliquot into working volumes to avoid repeated freeze-thaw cycles. The effect of storage of stock solution on compound performance should be tested for each application.

Compound has low solubility in aqueous media. For use as a cell culture supplement, stock solution should be diluted into culture medium immediately before use. Avoid final DMSO concentration above 0.1% due to potential cell toxicity.

Small Molecules Indomethacin



Published Applications

DIFFERENTIATION

- Activates peroxisome proliferator-activated receptor-γ (PPARγ), a ligand-activated transcription factor known to play a pivotal role in adipogenesis (Lehmann et al.).
- · Inhibits chondrogenic differentiation in ATDC5 cells and bone marrow stem cells (Caron et al.). CANCER RESEARCH
- · Inhibits growth of mouse mammary tumors (Fulton).
- · Induces apoptosis in prostate and gastric cancer cells (Chiou et al.; Liu et al.).

References

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Kurumbail RG et al. (1996) Structural basis for selective inhibition of cyclooxygenase-2 by anti-inflammatory agents. Nature 384(6610): 644–8.

Lehmann JM et al. (1997) Peroxisome proliferator-activated receptors alpha and gamma are activated by indomethacin and other non-steroidal anti-inflammatory drugs. J Biol Chem 272(6): 3406–10.

Liu C et al. (2017) Inhibition of AKR1C3 activation overcomes resistance to abiraterone in advanced prostate cancer. Mol Cancer Ther 16(1): 35–44.

Vane JR et al. (1998) Cyclooxygenases 1 and 2. Annu Rev Pharmacol Toxicol 38(1): 97-120.

Related Small Molecules

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This product is hazardous. Please refer to the Safety Data Sheet (SDS).

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