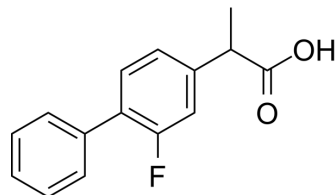


Flurbiprofen

Cat. No.:	HY-10582		
CAS No.:	5104-49-4		
Molecular Formula:	C ₁₅ H ₁₃ FO ₂		
Molecular Weight:	244.26		
Target:	Apoptosis; COX		
Pathway:	Apoptosis; Immunology/Inflammation		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 year



SOLVENT & SOLUBILITY

In Vitro

DMSO : ≥ 100 mg/mL (409.40 mM)
 * "≥" means soluble, but saturation unknown.

	Solvent Concentration	Mass		
		1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	4.0940 mL	20.4700 mL	40.9400 mL
	5 mM	0.8188 mL	4.0940 mL	8.1880 mL
	10 mM	0.4094 mL	2.0470 mL	4.0940 mL

Please refer to the solubility information to select the appropriate solvent.

In Vivo

- Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline
 Solubility: ≥ 2.5 mg/mL (10.23 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)
 Solubility: ≥ 2.5 mg/mL (10.23 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil
 Solubility: ≥ 2.5 mg/mL (10.23 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

Flurbiprofen (dl-Flurbiprofen) is a potent, orally active nonsteroidal anti-inflammatory agent (NSAIA/NSAID), with antipyretic and analgesic activities. Flurbiprofen is commonly used for the research of inflammatory diseases, including osteoarthritis and rheumatoid arthritis. Flurbiprofen is a non-selective cyclooxygenase (COX) inhibitor that can be used for the research of colorectal cancer^{[1][2][3]}.

IC₅₀ & Target

COX	COX-1	COX-2
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In Vitro

Flurbiprofen (2-20 nM; 12-48 hours) significantly decreases SW620 cells proliferation in a concentration- and time-dependent manner^[1].

Flurbiprofen (10 nM; 24 hours) decreases COX-2 expression^[1].

Flurbiprofen (10 nM; 24 hours) inhibits the expression of inflammatory factors by inhibiting COX-2^[1].

Flurbiprofen (10 nM; 24 hours) promotes the apoptosis of colorectal cancer cells by inhibiting COX-2^[1].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Cell Proliferation Assay^[1]

Cell Line:	SW620 cells
Concentration:	2 nM, 4 nM, 10 nM, 20 nM
Incubation Time:	12 hours, 24 hours, 48 hours
Result:	Inhibited colorectal cancer cell proliferation.

Western Blot Analysis^[1]

Cell Line:	SW620 cells
Concentration:	10 nM
Incubation Time:	24 hours
Result:	Significantly decreased the protein and mRNA levels of COX-2.

RT-PCR^[1]

Cell Line:	SW620 cells
Concentration:	10 nM
Incubation Time:	24 hours
Result:	Decreased COX-2 mRNA expression levels

Apoptosis Analysis^[1]

Cell Line:	SW620 cells
Concentration:	10 nM
Incubation Time:	24 hours
Result:	Significantly decreased the expression of Bcl2 and significantly increased the expression of Bax and cleaved-caspase3, with no effect on total caspase-3.

In Vivo

Flurbiprofen (0.3-4.8 mg/kg; p.o.; 4-5 dosages) has acute anti-inflammatory in adrenalectomized rats^[2].

Flurbiprofen (10 mg/kg; i.p.; daily; for 6 days) attenuates high-fat diet-induced obesity in mice^[3].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Rat ^[2]
Dosage:	0.3 mg/kg, 0.6 mg/kg, 1.2 mg/kg, 2.4 mg/kg, 4.8 mg/kg
Administration:	Oral administration, 4-5 dosages
Result:	Inhibited the acute inflammation.

REFERENCES

- [1]. Hosoi, T., et al., Flurbiprofen ameliorated obesity by attenuating leptin resistance induced by endoplasmic reticulum stress. *EMBO Mol Med*, 2014.
- [2]. Xiaobo Wang, et al. Flurbiprofen suppresses the inflammation, proliferation, invasion and migration of colorectal cancer cells via COX2. *Oncol Lett*. 2020 Nov; 20(5): 132.
- [3]. E M Glenn, et al. The pharmacology of 2-(2-fluoro-4-biphenyl)propionic acid (flurbiprofen). A potent non-steroidal anti-inflammatory drug. *Agents Actions*. 1973 Nov;3(4):210-6.
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Caution: Product has not been fully validated for medical applications. For research use only.

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