Ibuprofen sodium

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Cat. No.: CAS No.: Molecular Formula: Molecular Weight: Target: Pathway: Storage:	HY-78131C 31121-93-4 C ₁₃ H ₁₇ NaO ₂ 228.26 COX; Apoptosis; Parasite Immunology/Inflammation; Apoptosis; Anti-infection 4°C, sealed storage, away from moisture	ONa
Storage:	4°C, sealed storage, away from moisture * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)	

SOLVENT & SOLUBILITY

* "≥" me Prepari	0,	DMSO : 20.83 mg/mL (91.26 mM; ultrasonic and warming and heat to 60°C) * "≥" means soluble, but saturation unknown.					
		Solvent Mass Concentration	1 mg	5 mg	10 mg		
	Preparing Stock Solutions	1 mM	4.3810 mL	21.9048 mL	43.8097 mL		
	Stock Solutions	5 mM	0.8762 mL	4.3810 mL	8.7619 mL		
		10 mM	0.4381 mL	2.1905 mL	4.3810 mL		

BIOLOGICAL ACTIV	ΙΤΥ	
Description	Ibuprofen ((±)-Ibuprofen) sodium is an orally active, selective COX-1 inhibitor with an IC ₅₀ value of 13 μM. Ibuprofen sodium inhibits cell proliferation, angiogenesis, and induces cell apoptosis. Ibuprofen sodium is a nonsteroidal anti-inflammatory agent and a nitric oxide (NO) donor. Ibuprofen sodium can be used in the research of pain, swelling, inflammation, infection, immunology, cancers ^{[1][2][5][8]} .	
IC₅₀ & Target	СОХ-1 13 µМ (IC ₅₀)	COX-2 370 μM (IC ₅₀)
In Vitro	Ibuprofen sodium (24 h) inhibits COX-1 and COX-2 activity with IC ₅₀ values of 13 μM and 370 μM ^[1] . Ibuprofen sodium (500 μM, 48 h) inhibits cell proliferation and angiogenesis, and induces apoptosis in AGS cells (Adenocarcinoma gastric cell line) ^[2] . Ibuprofen sodium (500 μM, 48 h) downregulates transcription of Akt, VEGF-A, PCNA, Bcl2, OCT3/4 and CD44 genes, but upregulates RNA levels of wild type P53 and Bax genes in AGS cell ^[2] . Ibuprofen sodium (500 μM, 24 h) restores microtubule reformation, microtubule-dependent intracellular cholesterol transport, and induces extension of microtubules to the cell periphery in both cystic fibrosis (CF) cell models and primary C	

nasal epithelial cells^[3].

Ibuprofen sodium (500 μ M, 24 h) enhances UV-induced cell death in MCF-7 cells and MDA-MB-231 cells by a photosensitization process^[4].

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

Cell Viability Assay^[2]

Cell Line:	AGS cells
Concentration:	100-1000 μM
ncubation Time:	24 h, 48 h
Result:	Inhibited AGS cell viability with IC $_{50}$ values of 630 μ M (trypan blue staining, 24 h), 456 μ M (neutral red assay, 24 h), 549 μ M (trypan blue staining , 48 h) and 408 μ M (neutral red assay, 48 h).

In Vivo

Ibuprofen sodium (fed in animal feedings, 300 mg/kg, 14 days) reduces overall tumor growth and enhances anti-tumor immune characteristics without adverse autoimmune reactions in a model of postpartum breast cancer^[5].

Ibuprofen sodium (subcutaneous injection, 60 mg/kg, every second day for 15 days) reduces the risk of neuropathy in a rat model of chronic Oxaliplatin@induced peripheral neuropathy^[6].

Ibuprofen sodium (oral administration, 20 mg/kg, every 12 hours, 5 doses total) decreases muscle growth (average muscle fiber cross-sectional area) without affecting regulation of supraspinatus tendon adaptions to exercise^[7].

Ibuprofen sodium (oral administration, 35 mg/kg, twice daily) attenuates the Inflammatory response to pseudomonas aeruginosa in a rat model of chronic pulmonary infection^[8].

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

Animal Model:	Syngeneic (D2A1) orthotopic Balb/c mouse model of PPBC (postpartum) ^[5]	
Dosage:	300 mg/kg, daily for 14 days	
Administration:	Fed in animal feedings (added to pulverized standard chow and mixed dry, then mixed with water, made into chow pellets and dried thoroughly)	
Result:	Suppresed tumor growth, reduced presence of immature monocytes and increased numbers of T cells. Enhanced Th1 associated cytokines as well as promoted tumor border accumulation of T cells.	
Animal Model:	Oxaliplatin⊠induced peripheral neuropathy ^[6]	
Dosage:	60 mg/kg, every second day for 15 days	
Administration:	Subcutaneous injection	
Result:	Lowered sensory nerve conduction velocity (SNCV).	

CUSTOMER VALIDATION

- Cell Rep. 2019 Dec 17;29(12):3847-3858.e5.
- Chemosphere. 2019 Jun;225:378-387.
- Phytomedicine. 1 September 2022, 154427.

- EMBO Rep. 2022 Apr 11;e53932.
- Cells. 2022, 11(12), 1870.

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[2]. Hassan Akrami, et al. Inhibitory effect of ibuprofen on tumor survival and angiogenesis in gastric cancer cell. Tumour Biol. 2015 May;36(5):3237-43.

[3]. Sharon M Rymut, et al. Ibuprofen regulation of microtubule dynamics in cystic fibrosis epithelial cells. Am J Physiol Lung Cell Mol Physiol. 2016 Aug 1;311(2):L317-27.

[4]. Emmanuelle Bignon, et al. Ibuprofen and ketoprofen potentiate UVA-induced cell death by a photosensitization process. Sci Rep. 2017 Aug 21;7(1):8885.

[5]. Nathan D Pennock, et al. Ibuprofen supports macrophage differentiation, T cell recruitment, and tumor suppression in a model of postpartum breast cancer. J Immunother Cancer. 2018 Oct 1;6(1):98.

[6]. Thomas Krøigård, et al. Protective effect of ibuprofen in a rat model of chronic oxaliplatin-induced peripheral neuropathy. Exp Brain Res. 2019 Oct;237(10):2645-2651.

[7]. Sarah Ilkhanipour Rooney, et al. Ibuprofen Differentially Affects Supraspinatus Muscle and Tendon Adaptations to Exercise in a Rat Model. Am J Sports Med. 2016 Sep;44(9):2237-45.

[8]. M W Konstan, et al. Ibuprofen attenuates the inflammatory response to Pseudomonas aeruginosa in a rat model of chronic pulmonary infection. Implications for antiinflammatory therapy in cystic fibrosis. Am Rev Respir Dis. 1990 Jan;141(1):186-92.

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

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