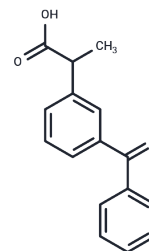


Ketoprofen

Chemical Properties

CAS No. :	22071-15-4
Formula:	C ₁₆ H ₁₄ O ₃
Molecular Weight:	254.28
Appearance:	no data available
Storage:	Powder: -20°C for 3 years In solvent: -80°C for 1 year



Biological Description

Description	Ketoprofen (RP-19583) is a propionic acid derivative and nonsteroidal anti-inflammatory drug (NSAID) with anti-inflammatory, analgesic, and antipyretic effects. It inhibits cyclo-oxygenase I and II, decreasing the formation of prostaglandin and thromboxane precursors. This reduction in prostaglandin synthesis, mediated by prostaglandin synthase, is responsible for its therapeutic effects. Additionally, Ketoprofen decreases thromboxane A ₂ formation via thromboxane synthase, inhibiting platelet aggregation.
Targets(IC50)	Apoptosis, COX, MRP
In vitro	Ketoprofen combined with UVB irradiation induces the cytotoxicity and suppresses DNA synthesis in HaCaT cells in a concentration-dependent manner. Ketoprofen combined with UVB irradiation inhibits the cell growth and induces G2/M cell cycle arrest by modulating the levels of cdc2, cyclin B1, Chk1, Tyr15-phosphorylated cdc2 and p21. Ketoprofen combined with UVB irradiation also provokes a striking accumulation of cyclin B1-cdc2-p21 complexes, concomitantly with an increase in the levels of Tyr15-phosphorylated cdc2 and p21 protein. Ketoprofen combined with UVB irradiation accentuates the apoptotic response to UVB radiation in HaCaT cells as evidenced by DAPI staining. [1]
In vivo	Ketoprofen at 1% level in suitable topical vehicles can effectively inhibit GCF-LTB ₄ and GCF-PGE ₂ and positively alter alveolar bone activity in the ligature-induced model of periodontitis in the monkey. [2] Ketoprofen (3.63 mg/kg bwt) reduces hoof pain and lameness to a greater extent than the 2.2 mg/kg dose and phenylbutazone. [3] Ketoprofen is more effective than local anesthesia (LA), or caudal epidural anesthesia (EPI) in decreasing cortisol and partially reverses the reduction in average daily gain (ADG) following castration. [4] Ketoprofen (40 and 80 mg/kg diet) reduces the incidence of transitional cell carcinoma of the urinary bladder by >70% from that seen in dietary mice. [5]

Solubility Information

Solubility	Ethanol: 48 mg/mL (188.77 mM), Sonication is recommended. DMSO: 55 mg/mL (216.3 mM), Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
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Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	3.9327 mL	19.6634 mL	39.3267 mL
5 mM	0.7865 mL	3.9327 mL	7.8653 mL
10 mM	0.3933 mL	1.9663 mL	3.9327 mL
50 mM	0.0787 mL	0.3933 mL	0.7865 mL

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Reference

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Ting ST, et al. J Anim Sci, 2003, 81(5), 1281-1293.

Rao KV, et al. Carcinogenesis, 1996, 17(7), 1435-1438.

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