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Data Sheet (Cat.No.T0984)



5-Fluorouracil

Chemical Properties

CAS No.: 51-21-8

Formula: C4H3FN2O2

Molecular Weight: 130.08

Appearance: no data available

Storage: Powder: -20°C for 3 years | In solvent: -80°C for 1 year

Biological Description

Biological Descript	lon
Description	5-Fluorouracil (5-FU) is a uracil analog, an inhibitor of DNA synthesis. 5-Fluorouracil has antitumor activity and affects pyrimidine synthesis through inhibition of thymidylate synthase. 5-Fluorouracil causes apoptosis and autophagy.
Targets(IC50)	Apoptosis, Nucleoside Antimetabolite/Analog, HIV Protease, DNA/RNA Synthesis, Endogenous Metabolite
In vitro	METHODS: Human cardiomyocytes HCM, human umbilical vein endothelial cells HUVE, and human colon cancer cells HCT116 and HT29 were treated with 5-Fluorouracil (0.01-1000 μM) for 24-96 h, and the growth inhibition of the cells was detected by MTT. RESULTS: The EC50 of 5-Fluorouracil on HCM, HUVE, PHCT116 and HT29 cells were 4.866 μM, 3.832 μM, 13.72 μM and 106.8 μM, respectively, at 72 h. [1] METHODS: Smooth muscle cells were treated with 5-Fluorouracil (0.05-10 mM) for 24 h. Apoptosis was detected using Flow Cytometry. RESULTS: 5-Fluorouracil at concentrations of 0.1, 1, and 10 mM induced apoptosis in cultured smooth muscle cells after 24 h of treatment, and the apoptotic cells were detached from the culture dishes. [2] METHODS: Human colon cancer cells SW620 were incubated with 5-Fluorouracil (13 μg/mL) for 24-48 h. ALP activity was detected using an ALP detection kit. RESULTS: Alkaline phosphatase (ALP) has been used to monitor the differentiation effects of certain anticancer compounds. Untreated SW620 cells showed relatively low ALP activity, while in cells treated with 13 μg/ml 5-FU, ALP activity reached high levels in a time-dependent manner. [3]
In vivo	METHODS: To detect the antitumor activity in vivo, 5-Fluorouracil (10-40 mg/kg) was injected intraperitoneally once a day for ten days into mice bearing ascites hepatocellular carcinoma tumor H22. RESULTS: 5-Fluorouracil at 10 mg/kg inhibited tumor growth while maintaining immune function in mice. 5-Fluorouracil exerts low-dose, low-toxicity antitumor effects and stimulates the host immune system. [4] METHODS: To study 5-Fluorouracil-induced intestinal injury, 5-Fluorouracil (100-200 mg/kg) was administered as a single intraperitoneal injection to BALB/c mice. RESULTS: The body weight and diarrhea symptoms of 5-Fluorouracil-treated animals were significantly reduced in a dose-dependent manner. Occludin and claudin-1 protein expression was significantly reduced in the 5-Fluorouracil-treated group. 5-Fluorouracil-treated group showed significantly higher NF-κBp65 protein and TNF-α mRNA expression than the control group. NF-κBp65 protein and TNF-α mRNA expression were significantly higher in the 5-Fluorouracil treated group than in the control group.

[5]

Cell Research

After a 7-day habituation period, the mice were divided into three groups (vehicle group, dextrin group, and ED group; n = 6 mice per group) that had the same mean body weight (time of grouping was designated as day 0). Then, mice were treated by tail vein injections from day 0 to 4; mice in the dextrin and ED groups received 40 mg/kg/day of 5- fluorouracil (5-FU) injection 250 mg, while mice in the vehicle group received 10 mL/kg/day physiological saline, which was equivalent to the dose of 5-FU. Additionally, twice a day from day 0 to 6, ED group mice received 1.6 kcal/0.8 mL/day ED administered orally, while mice in the vehicle and dextrin groups received dextrin containing the same amount of calories. Body weight and food consumption were measured before administration of 5-FU and ED on days 0 and 7. Food consumption was measured with respect to each group. Mice were accommodated individually in specially prepared polycarbonate cages, and two or more fresh stools per mouse were scored as follows: 0, the stools were not crushed when pushed by human fingers; 1, the stools were crushed, but the core remained when pushed by human fingers; 2, the stools were crushed and the core did not remain when pushed by human fingers; 3, the stools were crushed and stuck to the fingers when pushed by human fingers; 4, the stools lost their shape just from being touched. Autopsies were conducted on day 7. After bleeding, the large intestine (the colon and rectum) was taken, and the length was measured. The lumen was washed with physiological saline, the excess water was wiped off, and specimens were weighed. Section 3 cm distal from the center of the large intestine was fixed with formalin for histological evaluation. Salivary glands (the submandibular gland and sublingual gland) were collected and weighed. The collected salivary glands were fixed with formalin, and after the tissue sections were prepared, they were stained with hematoxylin and eosin [3].

Solubility Information

Solubility

Ethanol: 1.3 mg/mL (10 mM),

H2O: 10 mg/mL (76.8 mM), Heating at 50℃ is recommended.

DMSO: 18.33 mg/mL (140.94 mM),

(< 1 mg/ml refers to the product slightly soluble or insoluble)

Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	7.6876 mL	38.4379 mL	76.8758 mL
5 mM	1.5375 mL	7.6876 mL	15.3752 mL
10 mM	0.7688 mL	3.8438 mL	7.6876 mL
50 mM	0.1538 mL	0.7688 mL	1.5375 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

Focaccetti C, et al. Effects of 5-fluorouracil on morphology, cell cycle, proliferation, apoptosis, autophagy and ROS production in endothelial cells and cardiomyocytes. PLoS One.

br/>Zhao X, Lian X, Xie J, et al. Accumulated

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