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Product Data Sheet

Trifluridine/tipiracil hydrochloride mixture

Cat. No.:	HY-16478	
CAS No.:	733030-01-8	
Molecular Formula:	C ₁₀ H ₁₁ F ₃ N ₂ O ₅ -1/2C ₉ H ₁₁ ClN ₄ O ₂ -1/2HCl	F
Molecular Weight:	435.76	
Target:	Nucleoside Antimetabolite/Analog; Thymidylate Synthase	-
Pathway:	Cell Cycle/DNA Damage; Apoptosis	F. F
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

In Vitro H ₂ O DMF DMS Prep Stoc	H ₂ O : 100 mg/mL (229.48 mM; Need ultrasonic) DMF : 20 mg/mL (45.90 mM; Need ultrasonic) DMSO : 2.34 mg/mL (5.37 mM; Need ultrasonic and warming)					
		Solvent Mass Concentration	1 mg	5 mg	10 mg	
	Preparing Stock Solutions	1 mM	2.2948 mL	11.4742 mL	22.9484 mL	
		5 mM	0.4590 mL	2.2948 mL	4.5897 mL	
		10 mM	0.2295 mL	1.1474 mL	2.2948 mL	
	Please refer to the solubility information to select the appropriate solvent.					
In Vivo	1. Add each solvent one by one: Saline Solubility: 100 mg/mL (229.48 mM); Clear solution; Need ultrasonic					
	2. Add each solvent one by one: PBS Solubility: 50 mg/mL (114.74 mM); Clear solution; Need ultrasonic					

DIOLOGICALACITY	
Description	Trifluridine/tipiracil hydrochloride mixture (TAS-102) is a potent and orally active nucleoside antitumor agent. The composition of Trifluridine/tipiracil hydrochloride mixture (TAS-102) is a 1:0.5 mixture (on a molar basis) of alpha,alpha,alpha-tri-fluorothymidine (FTD) and thymidine phosphorylase inhibitor (TPI). Trifluridine/tipiracil hydrochloride mixture (TAS-102) is a 1:0.5 mixture (TAS-102) and thymidine phosphorylase inhibitor (TPI). Trifluridine/tipiracil hydrochloride mixture (TAS-102) is a 1:0.5 mixture (TAS-102). Trifluridine/tipiracil hydrochloride mixture (TAS-102) shows the antitumor activity mainly via the inhibition of thymidylate synthase (TS) and incorporation into DNA ^{[1][2]} .
In Vivo	Trifluridine/tipiracil hydrochloride mixture (150 mg/kg/day; p.o.; twice a day for 14 days) prevents body weight loss and reduces the relative tumor volume in colorectal cancer and gastric cancer mice models ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Male nude mice bearing KM12C, KM12C/5-FU, DLD-1, DLD-1/5-FU, and SC-2 ${\rm cells}^{[2]}$
Dosage:	150 mg/kg/day
Administration:	p.o.; twice a day for 14 days
Result:	Prevented body weight loss and reduced the relative tumor volume in colorectal cance and gastric cancer mice models.

CUSTOMER VALIDATION

• Am J Cancer Res. 2020 Nov 1;10(11):3752-3764.

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REFERENCES

[1]. Limagne E, Thibaudin M, Nuttin L, et al. Trifluridine/Tipiracil plus Oxaliplatin Improves PD-1 Blockade in Colorectal Cancer by Inducing Immunogenic Cell Death and Depleting Macrophages. Cancer Immunol Res. 2019;7(12):1958-1969.

[2]. Suzuki N, Nakagawa F, Takechi T. Trifluridine/tipiracil increases survival rates in peritoneal dissemination mouse models of human colorectal and gastric cancer [published correction appears in Oncol Lett. 2021 Jul;22(1):511]. Oncol Lett. 2017;14(1):639-646.

[3]. Emura T, et al. A novel antimetabolite, TAS-102 retains its effect on FU-related resistant cancer cells. Int J Mol Med. 2004;13(4):545-549.

[4]. Nukatsuka M, et al. Efficacy of combination chemotherapy using a novel oral chemotherapeutic agent, TAS-102, with irinotecan hydrochloride on human colorectal and gastric cancer xenografts. Anticancer Res. 2015;35(3):1437-1445.

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

Tel: 609-228-6898 Fax: 609-228-5909 E-mail: tech@MedChemExpress.com Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA