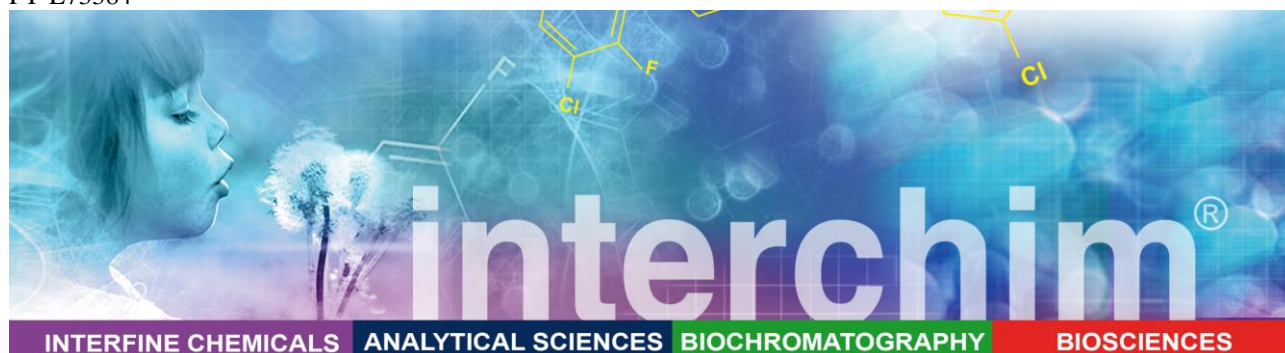


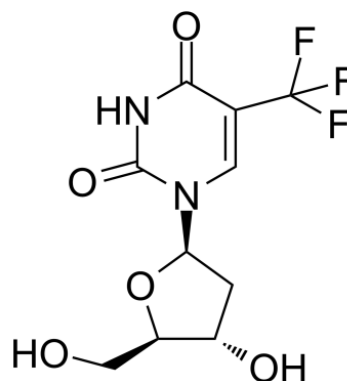
FT-E73384



Trifluridine

Product Description

Catalog #:	E73384, 100mg E73385, 500mg AX7MN0, 1ml 10mM in DMSO.
Name:	Trifluridine Syn: Trifluorothymidine; 5-Trifluorothymidine; TFT CAS : 70-00-8
MW :	296.20
Formula :	C ₁₀ H ₁₁ F ₃ N ₅ O ₅
Properties :	Soluble : in DMSO : ≥ 100 mg/mL (337.61 mM) Purity: 99.96%
Storage:	Powder: -20°C (long term; possible at +4°C (2 years) (M) In solvent: -80°C (6 months) -20°C (1 month)



For Research Use Only

Introduction

Trifluridine (Trifluorothymidine; 5-Trifluorothymidine; TFT) is an irreversible thymidylate synthase inhibitor, and thereby suppresses DNA synthesis. Trifluridine is an antiviral drug for herpes simplex virus (HSV) infection. IC₅₀ & Targets : DNA Synthesis (thymidylate synthase)

In Vitro

Solubility : DMSO : >100 mg/mL (337.61 mM)
Preparation : 1mM = 1mg in 3.3761 mL

In Vivo

Preparation :

1. Add each solvent one by one: 10% DMSO 40% PEG300 5% Tween-80 45% saline
Solubility: ≥ 2.5 mg/mL (8.44 mM); Clear solution
2. Add each solvent one by one: 10% DMSO 90% (20% SBE-β-CD in saline)
Solubility: ≥ 2.5 mg/mL (8.44 mM); Clear solution
3. Add each solvent one by one: 10% DMSO 90% corn oil
Solubility: ≥ 2.5 mg/mL (8.44 mM); Clear solution

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References

- [1]. Suzuki N, et al. Mode of action of trifluorothymidine (TFT) against DNA replication and repair enzymes. Int J Oncol. 2011 Jul;39(1):263-70.
- [2]. Suzuki N, et al. Trifluorothymidine exhibits potent antitumor activity via the induction of DNA double-strand breaks. Exp Ther Med. 2011 May;2(3):393-397.
- [3]. Temmink OH, et al. Irinotecan-induced cytotoxicity to colon cancer cells in vitro is stimulated by pre-incubation with trifluorothymidine. Eur J Cancer. 2007 Jan;43(1):175-83.
- [4]. Okayama T, et al. Involvement of concentrative nucleoside transporter 1 in intestinal absorption of trifluorothymidine, a novel antitumor nucleoside, in rats. J Pharmacol Exp Ther. 2012 Feb;340(2):457-62. More references: J Mol Med (Berl). 2019 Aug;97(8):1183-1193.

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