Proteins

Product Data Sheet

Indotecan

Cat. No.: HY-18351 CAS No.: 915303-09-2 Molecular Formula: $C_{26}H_{26}N_2O_7$ Molecular Weight: 478.49

Target: Topoisomerase

Pathway: Cell Cycle/DNA Damage Storage: 4°C, stored under nitrogen

* In solvent : -80°C, 6 months; -20°C, 1 month (stored under nitrogen)

SOLVENT & SOLUBILITY

In Vitro

DMSO: < 1 mg/mL (ultrasonic; warming; heat to 60°C) (insoluble or slightly soluble)

BIOLOGICAL ACTIVITY

Description	Indotecan (LMP-400), an indenoisoquinoline derivative, is a potent Topoisomerase I inhibitor, with IC $_{50}$ s of 300, 1200, 560 nM for P388, HCT116, MCF-7 cell lines, respectively. Indotecan prevents the relaxation of supercoiled DNA and can be used for the research of visceral leishmaniasis ^{[1][2]} .		
IC ₅₀ & Target	Topoisomerase I		
In Vitro	Indotecan (48 h) inhibits the proliferation of L. infantum promastigotes, ex vivo-infected splenocytes, and uninfected splenocytes, with IC $_{50}$ s of 0.10 μ M, 0.10 μ M, and 57.16 μ M, respectively ^[2] . Indotecan (1.1-90 μ M; 30 min) induces TopI-DNA complexes and inhibits DNA synthesis in L. infantum cultures ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.		
In Vivo	Indotecan (2.5 mg/kg; i.p. every 2 d for 15 d) depletes the parasitic burden in the spleen and liver of visceral leishmaniasis mice ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.		
	Animal Model:	Female BALB/c mice (4-6 weeks) were infected with metacyclic parasites intravenously	

Animal Model:	Female BALB/c mice (4-6 weeks) were infected with metacyclic parasites intravenously through the tail vein $^{[2]}$
Dosage:	2.5 mg/kg body weight per injection
Administration:	Intraperitoneally every 2 days for 15 days (total, eight doses)
Result:	A drastic reduction of the number of transforming amastigotes recovered from the target organs of drug-treated animals was observed.

REFERENCES

[1]. Balaña-Fouce R, et, al. Indotecan (LMP400) and a Chemother. 2012 Oct;56(10):5264-70.	AM13-55: two novel indenoisoquinolines show potential for treating visceral leishmaniasis. Antimicrob Agents
[2]. Seol Y, et, al. Single-Molecule Supercoil Relaxation Cancer Ther. 2015 Nov;14(11):2552-9.	on Assay as a Screening Tool to Determine the Mechanism and Efficacy of Human Topoisomerase IB Inhibitors. Mol
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