Talotrexin

Cat. No.: HY-10824 CAS No.: 113857-87-7 Molecular Formula: $C_{27}H_{27}N_{9}O_{6}$ Molecular Weight: 573.56

Target: Antifolate; Dihydrofolate reductase (DHFR)

Pathway: Cell Cycle/DNA Damage; Metabolic Enzyme/Protease

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.

Product Data Sheet

BIOLOGICAL ACTIVITY

Description	Talotrexin (PT523), an analog of <u>Aminopterin</u> (HY-14518), is a nonpolyglutamatable classic antifolate. Talotrexin is a RFC (reduced folate carrier) specific inhibitor and selectively inhibits RFC transport. Talotrexin shows antitumor activity by targeting DHFR to inhibit tumor growth $^{[1][2]}$.	
In Vitro	Talotrexin shows no significant PCFT or FR α inhibitory activity ^[1] . Talotrexin (1-1000 nM) inhibits proliferation of R1-11/Tet-on-RFC HeLa cells ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	
In Vivo	Talotrexin (0-35 mg/kg, IV injection, once weekly for 4 weeks) alone or combined with <u>Paclitaxel</u> (HY-B0015, 7.5 mg/kg) inhibits A549 tumor growth in mice ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	
	Animal Model:	NSCLC xenografts in athymic female nude mice (6-8 weeks) ^[2]
	Dosage:	15, 25, and 35 mg/kg, combined with <u>Paclitaxel</u> (HY-B0015, 7.5 mg/kg)
	Administration:	IV injection, once weekly for 4 weeks
	Result:	Inhibited A549 tumor growth.

REFERENCES

[1]. O'Connor C, et al. Folate transporter dynamics and therapy with classic and tumor-targeted antifolates. Sci Rep. 2021 Mar 18;11(1):6389.

[2]. Choy G S, et al. Combination talotrexin (PT-523) with paclitaxel in A549 non-small cell lung cancer (NSCLC) athymic nude mice xenografts[J]. Cancer Research, 2007, 67(1_Annual_Meeting).

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$

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