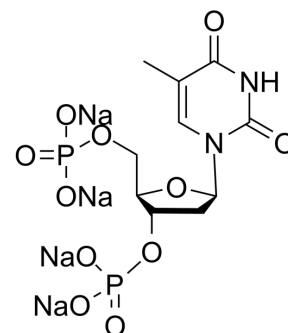


Thymidine 3',5'-diphosphate tetrasodium

Cat. No.:	HY-115581A
CAS No.:	118675-87-9
Molecular Formula:	C ₁₀ H ₁₂ N ₂ Na ₄ O ₁₁ P ₂
Molecular Weight:	490.12
Target:	Apoptosis; MicroRNA
Pathway:	Apoptosis; Epigenetics
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

H₂O : 50 mg/mL (102.02 mM; Need ultrasonic)
DMSO : 3.33 mg/mL (6.79 mM; ultrasonic and warming and heat to 60°C)

Preparing Stock Solutions	Solvent Concentration	Mass		
		1 mg	5 mg	10 mg
	1 mM	2.0403 mL	10.2016 mL	20.4032 mL
	5 mM	0.4081 mL	2.0403 mL	4.0806 mL
	10 mM	0.2040 mL	1.0202 mL	2.0403 mL

Please refer to the solubility information to select the appropriate solvent.

BIOLOGICAL ACTIVITY

Description

Thymidine 3',5'-diphosphate (Deoxythymidine 3',5'-diphosphate) tetrasodium is a selective inhibitor of staphylococcal nuclease and tudor domain containing 1 (SND1, the MicroRNA regulatory complex RISC subunit) and [3,5-²H₂] tyrosyl nuclease. Thymidine 3',5'-diphosphate tetrasodium has anti-tumor activity and can also be used as a catalyst in biochemical reactions^{[1][2]}.

IC₅₀ & Target

Staphylococcal nuclease and tudor domain containing 1, SND1; [3,5-(2)H(2)] Tyrosyl nuclease^{[1][2]}

In Vitro

Thymidine 3',5'-diphosphate tetrasodium (200 μM; 18 h) significantly reduces the expression level of p65 and p65 nuclear translocation in WT and Alb/SND1 (specific transgenic mouse overexpressing SND1) hepatocytes by inhibiting staphylococcal nuclease and tudor domain containing 1 (SND1) enzyme activity. Thymidine 3',5'-diphosphate tetrasodium inhibits the spherical formation of WT and Alb/SND1 hepatocytes^[1].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

In Vivo

Thymidine 3',5'-diphosphate tetrasodium (0.8 mg/kg; i.p.; twice a week for 4 weeks) has insignificant effect on serum liver enzymes (AST, ALT, AP), total protein (TP), albumin (Alb), and globulin (Glo) in WT B6/CBA mice^[1].

Thymidine 3',5'-diphosphate tetrasodium (0.8 mg/kg and 1.6 mg/kg; i.v.; twice a week for 4 weeks) significantly inhibits

tumor growth in WT B6/CBA mice^[1].

Thymidine 3',5'-diphosphate tetrasodium (0.8, 0.16 and 0.32 mg/kg; s.c.; twice a week for 4 weeks) inhibits tumor growth, inflammatory reaction and the expression of tumor initiating cells (TIC) markers in adult male NSG mice. Thymidine 3',5'-diphosphate tetrasodium up-regulates the expression of PTEN, TGFBR2 and CDKN1C apoptosis and selective tumor suppressor genes^[1].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	2 months old WT B6/CBA mice ^[1] .
Dosage:	0.8 mg/kg.
Administration:	Intraperitoneal injection; twice a week for 4 weeks.
Result:	Had insignificant effect on serum liver enzymes, total protein, albumin and globulin and showed biosafety.
Animal Model:	NSG mice ^[1] .
Dosage:	0.8, 0.16 or 0.32 mg/kg.
Administration:	Intravenous injection or subcutaneous injection; twice a week for 4 weeks.
Result:	Inhibited tumor growth, showed antitumor activity and immunomodulatory effects.

REFERENCES

[1]. Nidhi Jariwala, et al. Oncogenic Role of SND1 in Development and Progression of Hepatocellular Carcinoma. Cancer Res. 2017 Jun 15;77(12):3306-3316.

[2]. Cohen J S, et al. Proton magnetic resonance studies of the tyrosine residues of staphylococcal nuclease using [3, 5-2H2] tyrosine[J]. Biochimica et Biophysica Acta (BBA)-Protein Structure, 1971, 236(2): 468-478.

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

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