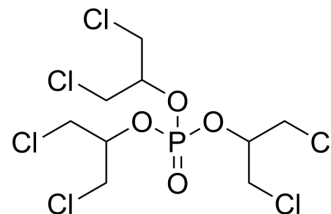


TDCPP

Cat. No.:	HY-108712		
CAS No.:	13674-87-8		
Molecular Formula:	C ₉ H ₁₅ Cl ₆ O ₄ P		
Molecular Weight:	430.9		
Target:	Biochemical Assay Reagents		
Pathway:	Others		
Storage:	Pure form	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



SOLVENT & SOLUBILITY

In Vitro

Ethanol : 100 mg/mL (232.07 mM; Need ultrasonic)
 DMSO : ≥ 62.5 mg/mL (145.05 mM)
 * "≥" means soluble, but saturation unknown.

Preparing Stock Solutions	Solvent Concentration	Mass		
		1 mg	5 mg	10 mg
	1 mM	2.3207 mL	11.6036 mL	23.2072 mL
	5 mM	0.4641 mL	2.3207 mL	4.6414 mL
	10 mM	0.2321 mL	1.1604 mL	2.3207 mL

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

In Vivo

- Add each solvent one by one: 15% Cremophor EL >> 85% Saline
Solubility: 33.33 mg/mL (77.35 mM); Suspended solution; Need ultrasonic
- Add each solvent one by one: 50% PEG300 >> 50% saline
Solubility: 33.33 mg/mL (77.35 mM); Suspended solution; Need ultrasonic
- Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline
Solubility: ≥ 2.08 mg/mL (4.83 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)
Solubility: ≥ 2.08 mg/mL (4.83 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil
Solubility: ≥ 2.08 mg/mL (4.83 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

TDCPP is a chlorinated analog of tris(2,3-dibromopropyl)phosphate (Tris) which is one of the most detected

organophosphorus flame retardants (OPFRs) in the environment.

In Vitro

Exposure to TDCPP does not significantly affect cell viability until at concentration $>68 \mu\text{g}/\text{mL}$. HCECs show a 16% cell viability loss after exposing to $136 \mu\text{g}/\text{mL}$ TDCPP. Moreover, TDCPP induces a sharp decrease in viable cells (87%) after exposing to $\geq 272 \mu\text{g}/\text{mL}$ TDCPP. Based on cell viability, the LC_{50} value for TDCPP is $202 \mu\text{g}/\text{mL}$ using a nonlinear regression. Compare to controls, TDCPP-exposed cells exhibit a concentration-dependent increase in apoptosis. Anti-apoptotic Bcl-2 protein expression is increased to 1.4 fold after exposing to $2 \mu\text{g}/\text{mL}$ TDCPP, 1.2-folds at $20 \mu\text{g}/\text{mL}$ but dynamically decreased to 0.4 fold at $200 \mu\text{g}/\text{mL}$ compare to control. The caspase-3 activity is increased to 2.1 folds of the control at $200 \mu\text{g}/\text{mL}$ TDCPP^[1]. TDCPP inhibits cell growth at lower concentrations (IC_{50} of $27 \mu\text{M}$), while cell viability and toxicity are affected at higher concentrations (IC_{50} of $171 \mu\text{M}$ and $168 \mu\text{M}$, respectively)^[2]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

PROTOCOL

Kinase Assay ^[1]

The cellular ATP contents are determined in HCECs grown in DMEM containing 0, 2, 20, or $200 \mu\text{g}/\text{mL}$ TDCPP using a luciferase-based ATP assay kit according to the manufacturer's guideline. Briefly, after 24 h exposure, HCECs are lysed with lysis buffer. Lysates are then centrifuged at $12,000 \text{ g}$ at 4°C for 5 min. Then, $100 \mu\text{L}$ of supernatant is mixed with $100 \mu\text{L}$ ATP detection working dilution. Luminance is examined by an fluorescence microplate reader^[1].

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

Cell Assay ^[1]

To examine the effects of TDCPP on cell viability, HCECs are planted into 96-well plate ($100 \mu\text{L}/\text{well}$) at density of 1×10^5 cells/mL overnight. Then, the medium is changed into fresh medium containing 0.034, 0.34, 3.4, 34, 68, 136, 272, or $340 \mu\text{g}/\text{mL}$ of TDCPP and solvent vehicle (0.1%, v/v) and incubated for 24 h. Cell viability is detected using CCK-8 cell viability assay kit according to the manufacturer's instructions. After exposure, cellular morphology is observed and recorded by an inverted microscopy^[1].

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

CUSTOMER VALIDATION

- J Hazard Mater. 2022 Jul 15;434:128824.
- Sci Total Environ. 2025 Jan 10;961:178429.
- Environ Pollut. 2024 Mar 8:123740.
- Chemosphere. 2022 Sep 7;136345.
- J Cereb Blood Flow Metab. 2024 Jun 9:271678X241260100.

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REFERENCES

[1]. Xiang P, et al. Effects of organophosphorus flame retardant TDCPP on normal human corneal epithelial cells: Implications for human health. Environ Pollut. 2017 Nov;230:22-30.

[2]. Killilea DW, et al. Flame retardant tris(1,3-dichloro-2-propyl)phosphate (TDCPP) toxicity is attenuated by N-acetylcysteine in human kidney cells. Toxicol Rep. 2017 May 17;4:260-264.

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

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