Product Data Sheet



TDCPP

Cat. No.: HY-108712 CAS No.: 13674-87-8 Molecular Formula: $C_9H_{15}Cl_6O_4P$ Molecular Weight: 430.9 Target: Others Pathway: Others

Pure form Storage: -20°C 3 years

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: \geq 62.5 mg/mL (145.05 mM)

* "≥" means soluble, but saturation unknown.

Preparing Stock Solutions	Solvent Mass Concentration	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

- 1. Add each solvent one by one: 15% Cremophor EL >> 85% Saline Solubility: 33.33 mg/mL (77.35 mM); Suspended solution; Need ultrasonic
- 2. Add each solvent one by one: 50% PEG300 >> 50% saline Solubility: 33.33 mg/mL (77.35 mM); Suspended solution; Need ultrasonic
- 3. 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
- 4. 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
- 5. 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 μ g/mL. HCECs show a 16% cell viability loss after exposing to 136 μ g/mL TDCPP. Moreover, TDCPP induces a sharp decrease in viable cells (87%) after exposing to \geq 272 μ g/mL TDCPP. Based on cell viability, the LC₅₀ value for TDCPP is 202 μ g/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 μ g/mL TDCPP, 1.2-folds at 20 μ g/mL but dynamically decreased to 0.4 fold at 200 μ g/mL compare to control. The caspase-3 activity is increased to 2.1 folds of the control at 200 μ g/mL TDCPP[1]. TDCPP inhibits cell growth at lower concentrations (IC₅₀ of 27 μ M), while cell viability and toxicity are affected at higher concentrations (IC₅₀ of 171 μ M and 168 μ 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 μ g/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 g at 4°C for 5 min. Then, 100 μ L of supernatant is mixed with 100 μ L ATP detection working dilution. Luminance is examined by an fluorescence microplate reader^[1].

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Cell Assay [1]

To examine the effects of TDCPP on cell viability, HCECs are planted into 96-well plate (100 μ L/well) at density of 1×10⁵ cells/mL overnight. Then, the medium is changed into fresh medium containing 0.034, 0.34, 3.4, 34, 68, 136, 272, or 340 μ g/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.
- Chemosphere. 2022 Sep 7;136345.
- Gene. 2022 Feb 17;822:146349.

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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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