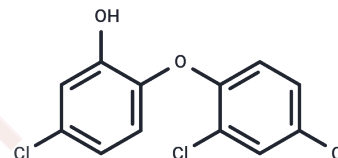


Triclosan

Chemical Properties

| | |
|-------------------|---|
| CAS No. : | 3380-34-5 |
| Formula: | C ₁₂ H ₇ Cl ₃ O ₂ |
| Molecular Weight: | 289.54 |
| Storage: | Powder: -20°C for 3 years In solvent: -80°C for 1 year Actual storage temperature shall be subject to the COA. |



Biological Description

| | |
|---------------|---|
| Description | Triclosan (Irgasan) is an antibacterial and antifungal agent. |
| Targets(IC50) | Apoptosis,Antibacterial,Antibiotic,Autophagy,Antifungal,NADPH |
| In vitro | cancer cells with triclosan at the physiologically related concentrations significantly increases the colony number of the cancer cells assessed by tumor formation assay. Triclosan-treated cells have a mesenchymal-like morphology and decrease the cell-to-cell adhesion. Triclosan-treated cells exhibit decreased E-cadherin while significantly up-regulated levels of EMT markers, namely N-cadherin, vimentin, snail and slug. Triclosan may potentiate cancer cells survival in detached condition and motility via the process of EMT[2]. |
| In vivo | Triclosan plays a pathologic role in promoting tissue fibrogenesis, leading to liver carcinogenesis. It accelerates both HCC development and its long-term effects on liver damage, accompanied by fibrosis and inflammation in mice[1]. |
| Cell Research | CV-1 cells are maintained in DMEM supplemented with 10% FBS, seeded in 96-well plates, and transfected with an expression vector containing a specific nuclear receptor and RXR, along with a luciferase reporter containing the appropriate DNA response element. The assay is first validated with known nuclear receptor ligands as positive controls. For the CAR ligand-binding assay, CV-1 cells are transiently transfected with the expression vector containing the Gal4 DNA-binding domain fused with the ligand-binding domain of murine or human CAR and cotransfected with the luciferase reporter plasmid, mh 100-luc. The day after the transfection, the positive compound of each nuclear receptor or TCS(Triclosan) is added and the cells are incubated for an additional 24 h. The luciferase activities are measured and normalized by β-gal activity.(Only for Reference) |

Solubility Information

| | |
|------------|---|
| Solubility | DMSO: 250 mg/mL (863.44 mM),Sonication is recommended. Ethanol: 57 mg/mL (196.86 mM),Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble) |
|------------|---|

| | |
|---------------------|---|
| In vivo Formulation | <p>10% DMSO+90% Saline: < 10 mg/mL (34.54 mM), Lower concentrations may be soluble, but exact solubility limit is unknown.</p> <p>10% DMSO+40% PEG300+5% Tween 80+45% Saline: 10 mg/mL (34.54 mM), Solution.</p> <p><i>Please add the solvents sequentially, clarifying the solution as much as possible before adding the next one. Dissolve by heating and/or sonication if necessary. Working solution is recommended to be prepared and used immediately. The formulation provided above is for reference purposes only. In vivo formulations may vary and should be modified based on specific experimental conditions.</i></p> |
|---------------------|---|

Preparing Stock Solutions

| | 1mg | 5mg | 10mg |
|-------|-----------|------------|------------|
| 1 mM | 3.4538 mL | 17.2688 mL | 34.5375 mL |
| 5 mM | 0.6908 mL | 3.4538 mL | 6.9075 mL |
| 10 mM | 0.3454 mL | 1.7269 mL | 3.4538 mL |
| 50 mM | 0.0691 mL | 0.3454 mL | 0.6908 mL |

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

Reference

Yueh MF, et al. Proc Natl Acad Sci U S A. 2014, 111(48):17200-5.

Hao Z, Wu Q, Li Z, et al. Maternal exposure to triclosan constitutes a yet unrecognized risk factor for autism spectrum disorders. Cell Research. 2019, 29(10): 866-869

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Hao Z, Wu Q, Li Z, et al. Maternal exposure to triclosan constitutes a yet unrecognized risk factor for autism spectrum disorders. Cell Research. 2019, 29(10): 866-869.

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