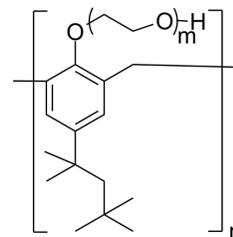


## Tyloxapol

Cat. No.:	HY-B1068
CAS No.:	25301-02-4
Molecular Formula:	$(C_{15}H_{21}O(C_2H_4O)_m)_n$
Target:	Biochemical Assay Reagents
Pathway:	Others
Storage:	4°C, protect from light * In solvent : -80°C, 6 months; -20°C, 1 month (protect from light)



$$m = 6 - 8$$

$$n \leq 5$$

### SOLVENT & SOLUBILITY

In Vitro	<p>H<sub>2</sub>O : 120 mg/mL (Need ultrasonic)</p> <p>Ethanol : 100 mg/mL (Need ultrasonic)</p> <p>DMSO : <math>\geq</math> 38 mg/mL</p> <p>* "<math>\geq</math>" means soluble, but saturation unknown.</p>
In Vivo	<p>1. Add each solvent one by one: PBS</p> <p>Solubility: 100 mg/mL (Infinity mM); Clear solution; Need ultrasonic</p>

### BIOLOGICAL ACTIVITY

Description	<p>Tyloxapol (Triton WR1339) is a nonionic liquid polymer of the alkyl aryl polyether alcohol type, used as a surface active stabilizer. Tyloxapol (Triton WR1339) is used to induce hyperlipidemia in animals<sup>[1][2]</sup>.</p>								
In Vitro	<p>Tyloxapol (100 <math>\mu</math>g/mL) triggers the detachment of HEK293 cells<sup>[2]</sup>.</p> <p>Tyloxapol induces nuclear fragmentation and the appearance of apoptotic nuclei<sup>[2]</sup>.</p> <p>Tyloxapol increases the risk of pulmonary haemorrhage, causes cytotoxicity in epithelial and red blood cells, and induces lysis of human Jurkat T-lymphoblasts<sup>[2]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p>								
In Vivo	<p>Tyloxapol (Triton WR1339, 50 mg/kg) causes significant (<math>P &lt; 0.05</math>) decreases in the activities of the AChE and MAO enzymes in rat plasma and brain<sup>[1]</sup>.</p> <p>?Tyloxapol leads to significant (<math>P &lt; 0.05</math>) reduction in the plasma urea, creatinine, and bilirubin<sup>[1]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1"> <tr> <td>Animal Model:</td> <td>Twenty-one adult male Wistar rats, aged 11–12 weeks weighing 180–200 g<sup>[1]</sup>.</td> </tr> <tr> <td>Dosage:</td> <td>50 mg/kg.</td> </tr> <tr> <td>Administration:</td> <td>Injected intraperitoneally, BW, every other day.</td> </tr> <tr> <td>Result:</td> <td>Caused a significant (<math>P &lt; 0.05</math>) elevation in the levels of TBARS combined with an inhibition</td> </tr> </table>	Animal Model:	Twenty-one adult male Wistar rats, aged 11–12 weeks weighing 180–200 g <sup>[1]</sup> .	Dosage:	50 mg/kg.	Administration:	Injected intraperitoneally, BW, every other day.	Result:	Caused a significant ( $P < 0.05$ ) elevation in the levels of TBARS combined with an inhibition
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of the antioxidant enzymes (GPx, GST, CAT, SOD) in rat plasma, liver, and brain.  
Induced DNA fragmentation and inhibited the activities of acetylcholinesterase and monoaminoxidase in the brain.

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## CUSTOMER VALIDATION

- Oxid Med Cell Longev. 2022 May 24;2022:1889632.

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## REFERENCES

- [1]. Heba Mohamed Abdou, et al. Triton WR-1339-induced hyperlipidemia, DNA fragmentation, neurotransmitters inhibition, oxidative damage, histopathological and morphometric changes: the protective role of soybean oil. The Journal of Basic and Applied Zoology volume 79, Article number: 51 (2018).
- [2]. Julijana Kristl, et al. Surface active stabilizer tyloxapol in colloidal dispersions exerts cytostatic effects and apoptotic dismissal of cells. Toxicol Appl Pharmacol. 2008 Oct 15;232(2):218-25.
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**Caution: Product has not been fully validated for medical applications. For research use only.**

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