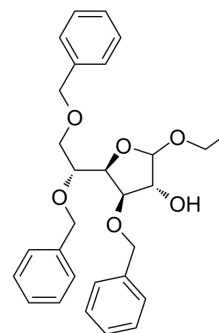


Tribenoside

Cat. No.:	HY-108249
CAS No.:	10310-32-4
Molecular Formula:	C ₂₉ H ₃₄ O ₆
Molecular Weight:	478.58
Target:	Others
Pathway:	Others
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Tribenoside is a vasoprotective agent, can be used for the research of hemorrhoids. Tribenoside has mild anti-inflammatory, analgesic, and wound healing properties ^[1] .									
In Vitro	<p>Tribenoside (1 nM-100 μM) exhibits cytotoxic effect against HeLa cells with the EC₅₀ of 13.74 μM^[2]. MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Viability Assay^[2]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>HeLa cells</td> </tr> <tr> <td>Concentration:</td> <td>1 nM, 10 nM, 100 nM, 1 μM, 10 μM, 100 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>72 hours</td> </tr> <tr> <td>Result:</td> <td>Reduced HeLa cells viability in a dose-responsive manner.</td> </tr> </table>		Cell Line:	HeLa cells	Concentration:	1 nM, 10 nM, 100 nM, 1 μM, 10 μM, 100 μM	Incubation Time:	72 hours	Result:	Reduced HeLa cells viability in a dose-responsive manner.
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Concentration:	1 nM, 10 nM, 100 nM, 1 μM, 10 μM, 100 μM									
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Result:	Reduced HeLa cells viability in a dose-responsive manner.									
In Vivo	<p>Tribenoside (oral administration in doses of 500 and 1,200 mg/kg weekly) significantly decreases in the development of osteoporosis^[3]. 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>male C57 black mice^[3]</td> </tr> <tr> <td>Dosage:</td> <td>500 and 1,200 mg/kg weekly</td> </tr> <tr> <td>Administration:</td> <td>Oral administration</td> </tr> <tr> <td>Result:</td> <td>Led to a significant reduction in the overall arthrotic involvement.</td> </tr> </table>		Animal Model:	male C57 black mice ^[3]	Dosage:	500 and 1,200 mg/kg weekly	Administration:	Oral administration	Result:	Led to a significant reduction in the overall arthrotic involvement.
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REFERENCES

- [1]. Yamato Kikkawa, et al. The influence of Tribenoside on expression and deposition of epidermal laminins in HaCaT cells. *Biol Pharm Bull.* 2010;33(2):307-10.
- [2]. Yu-Chen Lo, et al. Computational Cell Cycle Profiling of Cancer Cells for Prioritizing FDA-Approved Drugs with Repurposing Potential. *Sci Rep.* 2017 Sep 12;7(1):11261.

[3]. G Wilhelmi, et al. Suitability of the C57 black mouse as an experimental animal for the study of skeletal changes due to ageing, with special reference to osteo-arthritis and its response to tribenoside. *Pharmacology*. 1976;14(4):289-96.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA