

Produktinformation



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PRODUCT INFORMATION



Chrysoeriol

Item No. 33395

CAS Registry No.: 491-71-4

Formal Name: 5,7-dihydroxy-2-(4-hydroxy-3-

methoxyphenyl)-4H-1-benzopyran-4-one

Synonyms: 3'-methoxy Apigenin, Luteolin 3'-methyl ether

MF: $C_{16}H_{12}O_6$ 300.3 FW: **Purity:** ≥98% UV/Vis.:

 λ_{max} : 398 nm Supplied as: A crystalline solid

Storage: -20°C Stability: ≥2 years Item Origin: Synthetic

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.

Laboratory Procedures

Chrysoeriol is supplied as a crystalline solid. A stock solution may be made by dissolving the chrysoeriol in the solvent of choice, which should be purged with an inert gas. Chrysoeriol is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide (DMF). The solubility of chrysoeriol in ethanol is approximately 1 mg/ml and approximately 30 mg/ml in DMSO and DMF.

Chrysoeriol is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, chrysoeriol should first be dissolved in DMSO and then diluted with the aqueous buffer of choice. Chrysoeriol has a solubility of approximately 0.25 mg/ml in a 1:3 solution of DMSO:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

Description

Chrysoeriol is a flavonoid that has been found in Capsicum and has diverse biological activities. 1-4 It is active against the Gram-positive bacteria E. faecalis, B. subtilis, and S. aureus (MICs = 1, 1, and 0.25 µg/ml, respectively) and the Gram-negative bacteria P. aeruginosa, K. pneumoniae, and E. coli (MICs = 0.12, 0.25, and 0.06 μg/ml, respectively). Chrysoeriol (7.5, 15, and 30 μM) induces cell cycle arrest at the G₁ phase and autophagy in A549 lung cancer cells.² It reduces LPS-induced production of IL-6, IL-1 β , and TNF- α in RAW 264.7 cells when used at a concentration of 20 μM.³ Chrysoeriol (20 mg/kg) reduces plasma glucose, total cholesterol, free fatty acid, and phospholipid levels in a rat model of diabetes induced by streptozotocin (STZ; Item No. 13104).4

References

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- 2. Wei, W., He, J., Ruan, H., et al. In vitro and in vivo cytotoxic effects of chrysoeriol in human lung carcinoma are facilitated through activation of autophagy, sub-G1/G0 cell cycle arrest, cell migration and invasion inhibition and modulation of MAPK/ERK signalling pathway. J. BUON 24(3), 936-942 (2019).
- Wu, J.-Y., Chen, Y.-J., Bai, L., et al. Chrysoeriol ameliorates TPA-induced acute skin inflammation in mice and inhibits NF-kB and STAT3 pathways. Phytomedicine 68, 153173 (2020).
- 4. Baskaran, K., Pugalendi, K.V., and Saravanan, R. Antidiabetic and antihyperlipidemic activity of chrysoeriol in diabetic rats, role of HMG CoA reductase, LCAT and LPL: In vivo and in silico approaches. J. Pharm. Res. 9(9), 597-605 (2015).

WARNING
THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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