PRODUCT INFORMATION



Tiliroside

Item No. 34493

CAS Registry No.: 20316-62-5

Formal Name: 5,7-dihydroxy-2-(4-hydroxyphenyl)-3-[[6-O-[(2E)-3-(4-

hydroxyphenyl)-1-oxo-2-propen-1-yl]-β-D-glucopyranosyl]

oxy]-4H-1-benzopyran-4-one

Synonyms: Kaempferol 3-O-β-D-Glucopyranoside-6-p-coumaril ester,

Kaempferol 3-O-β-D-(6"-E-p-coumaroyl)-glucopyranoside

MF: $C_{30}H_{26}O_{13}$ FW: 594.5 **Purity:** ≥98%

 λ_{max} : 268, 317 nm UV/Vis.:

A solid Supplied as: -20°C Storage: ≥4 years Stability:

Plant/Edgeworthia chrysantha Item Origin:

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

Laboratory Procedures

Tiliroside is supplied as a solid. A stock solution may be made by dissolving the tiliroside in the solvent of choice, which should be purged with an inert gas. Tiliroside is soluble in organic solvents such as DMSO and dimethyl formamide. The solubility of tiliroside in these solvents is approximately 30 mg/ml.

Tiliroside is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, tiliroside should first be dissolved in DMSO and then diluted with the aqueous buffer of choice. Tiliroside has a solubility of approximately 0.14 mg/ml in a 1:6 solution of DMSO:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

Description

Tiliroside is a polyketide synthase-derived flavonoid glycoside that has been found in Q. ilex and has diverse biological activities. 1-5 It inhibits carbonic anhydrase VII (CAVII; K; = 4.6 nM), but not CAI, CAII, CAIV, or CAXII ($K_s = >10, >10, 5.46$, and $0.134 \mu M$, respectively), in a cell-free assay.⁵ Tiliroside scavenges DPPH (Item No. 14805) and superoxide radicals (IC $_{50}$ s = 12.8 and 42 μ M, respectively) and is cytotoxic to Jurkat, HepG2, and COLO 205 cells (IC $_{50}$ s = 11.6, 14.3, and 55.4 μ M, respectively).² It reduces ear edema induced by phorbol 12-myristate 13-acetate (TPA; Item No. 10008014) with an ID₅₀ value of 0.357 mg/ear.³ Tiliroside (100 mg/kg) decreases triglyceride accumulation in the liver and skeletal muscle in high-fat diet-fed KKAy diabetic mice.4

References

- 1. Grochowski, D.M., Locatelli, M., Granica, S., et al. A review on the dietary flavonoid tiliroside. Compr. Rev. Food Sci. Food Saf. 17(5), 1395-1421 (2018).
- Rao, Y.K., Geethangili, M., Fang, S.-H., et al. Antioxidant and cytotoxic activities of naturally occurring phenolic and related compounds: a comparative study. Food Chem. Toxicol. 45(9), 1770-1776 (2007).
- Sala, A., Recio, M.C., Schinella, G.R., et al. Assessment of the anti-inflammatory activity and free radical scavenger activity of tiliroside. Eur. J. Pharmacol. 461(1), 53-61 (2003).
- Goto, T., Teraminami, A., Lee, J.-Y., et al. Tiliroside, a glycosidic flavonoid, ameliorates obesity-induced metabolic disorders via activation of adiponectin signaling followed by enhancement of fatty acid oxidation in liver and skeletal muscle in obese-diabetic mice. J. Nutr. Biochem. 23(7), 768-776 (2012).
- 5. Karioti, A., Ceruso, M., Carta, F., et al. New natural product carbonic anhydrase inhibitors incorporating phenol moieties. Bioorg. Med. Chem. 23(22), 7219-7225 (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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