

Tiliroside Datasheet

4th Edition (Revised in July, 2016)

[Product Information]

Name: Tiliroside

Catalog No.: CFN98026

Cas No.: 20316-62-5

Purity: > 95%

M.F: C₃₀H₂₆O₁₃

M.W: 594.52

Physical Description: Yellow powder

HO OH OH OH

Synonyms:(2E)-3-(4-Hydroxyphényl)acrylatede[(2R,3S,4S,5R,6S)-6-{[5,7-dihydroxy-2-(4-hydroxyphényl)-4-oxo-4H-chromén-3-yl]oxy}-3,4,5-trihydroxytétrahydro-2H-pyran-2-yl]m éthyle;4H-1-Benzopyran-4-one,5,7-dihydroxy-2-(4-hydroxyphenyl)-3-[[6-O-[(2E)-3-(4-hydroxyphenyl)-1-oxo-2-propenyl]-.beta.-D-glucopyranosyl]oxy]-.

[Intended Use]

- 1. Reference standards;
- 2. Pharmacological research;
- 3. Synthetic precursor compounds;
- 4. Intermediates & Fine Chemicals;
- 5. Others.

[Source]

The herbs of Agrimonia pilosa Ledeb.

[Biological Activity or Inhibitors]

Tiliroside, a dietary flavonol glycoside from the flowers of Tilia argentea (linden), shows a

hepatoprotective effect against D-galactosamine (D-GalN)/lipopolysaccharide

(LPS)-induced liver injury in mice.[1]

Tiliroside has anti-inflammatory and antioxidant activity, it can significantly reduce the

oedema and leukocyte infiltration induced by 12-O-tetradecanoylphorbol 13-acetate

(TPA).[2]

Tiliroside shows very potent anti-complement activity (IC50=5.4 x 10(-5) M) on the

classical pathway of the complement system. [3]

Tiliroside and gnaphaliin are antioxidants against in vitro Cu 2+ -induced LDL oxidation in

the same order of magnitude compared to that of the reference drug, probucol.[4]

Tiliroside has anti-diabetic effects, the effects at least partially mediated through inhibitory

effects on carbohydrate digestion and glucose uptake in the gastrointestinal tract. [5]

Tiliroside has shown in vivo anti-inflammatory activity; it also can inhibit

neuroinflammation in BV2 microglia through a mechanism involving TRAF-6-mediated

activation of NF-kB and p38 MAPK signalling pathways, these activities are possibly due,

in part, to the antioxidant property of this compound. [6]

[Solvent]

Pyridine, Methanol, Ethanol, etc.

[HPLC Method]^[7]

Mobile phase: Acetonitrile- 0.1% Acetic acid H2O, gradient elution;

Flow rate: 1.0 ml/min;

Column temperature: 25 °C;

The wave length of determination: 310 nm.

[Storage]

2-8℃, Protected from air and light, refrigerate or freeze.

[References]

[1] Matsuda H, Ninomiya K, Shimoda H, et al. Bioorg. Med. Chem. 2002, 10(3):707-12.

[2] Sala A, Recio M C, Schinella G R, et al. Eur. J. Pharmacol., 2003, 461(1):53-61.

[3] Jung K Y, Oh S R, Park S H, et al. Biol. Pharmaceut. Bull., 1998, 21(10):1077-8.

[4] Schinella G R, Tournier H A, Máñez S, et al. Fitoterapia, 2007, 78(1):1-6.

[5] Goto T, Horita M, Nagai H, et al. Mol. Nutr.Food Res., 2012, 56(3):435-45.

[6] Velagapudi R, Aderogba M, Olajide O A.B.B.A.Gen. Subjects, 2014, 1840(12):3311-9.

[7] Song L, Lu J, Yan P F, et al. China Journal of Chinese Materia Medica, 2009, 34(3):301-3.

[Contact]

Address:

S5-3 Building, No. 111, Dongfeng Rd., Wuhan Economic and Technological Development Zone, Wuhan, Hubei 430056,

China

Email: info@chemfaces.com

Tel: +86-27-84237783
Fax: +86-27-84254680
Web: www.chemfaces.com

Tech Support: service@chemfaces.com