

Tectorigenin Datasheet

4th Edition (Revised in July, 2016)

[Product Information]

Name: Tectorigenin

Catalog No.: CFN99920

Cas No.: 548-77-6

Purity: > 98%

M.F: C₁₆H₁₂O₆

M.W: 300.26

Physical Description: Yellow cryst.

Synonyms: 5,7-Dihydroxy-3-(4-hydroxyphenyl)-6-methoxy-1-benzopyran-4-one.

HO OH O

[Intended Use]

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

[Source]

The rhizomes of Iris tectorrum.

[Biological Activity or Inhibitors]

Tectorigenin (Tg) and tectoridin (Td) are the major compounds isolated from the rhizomes

of iridaceous plant Belamcanda chinensis which is well known as a chinese traditional

medicine for the treatment of inflammatory diseases; tectorigenin

IFN-y/LPS-induced inflammatory responses in murine macrophage RAW 264.7 cells, it

appears to have the potential to prevent inflammation.[1]

Tectorigenin and several other phytochemicals downregulate PDEF, PSA and IGF-1

receptor mRNA expression in vitro, the downregulation of PDEF, PSA, hTERT and IGF-1

receptor gene expression by tectorigenin demonstrates the antiproliferative potential of

these agents, they may be new and established targets for therapies in prostate cancer.[2]

Tectorigenin and kaikasaponin III have hypoglycemic and hypolipidemic effects in the

streptozotocin-induced diabetic rat and their antioxidant activity.[3]

Tectorigenin-paclitaxel-induces nuclear translocation of NFkB and the phosphorylation of

IkB and IKK, suggests that tectorigenin could sensitize paclitaxel-resistant human ovarian

cancer cells through inactivation of the Akt/IKK/IkB/NFkB signaling pathway, and promise

a new intervention to chemosensitize paclitaxel-induced cytotoxicity in ovarian cancer.[4]

Tectorigenin has inhibitory effect of the activities of plasma ALT, the effects is much more

potent than that of a commercially available dimethyl diphenyl bicarboxylate; orally

administered tectoridin shows hepatoprotective activity; tectorigenin also protects against

the cytotoxicity of HepG2 cells induced by t-BHP, this protection may have originated from

the inhibition of apoptosis; tectorigenin may be hepatoprotective and tectoridin should be

a prodrug that is transformed to tectorigenin.^[5]

Tectorigenin inhibits the in vitro proliferation and enhances miR-338* expression of

pulmonary fibroblasts in rats with idiopathic pulmonary fibrosis. [6]

[Solvent]

Chloroform, Dichloromethane, Ethyl Acetate, DMSO, Acetone, etc.

[HPLC Method]^[7]

Mobile phase: Methanol- Acetonitrile- H2O=2:1:2;

Flow rate: 1.0 ml/min;

Column temperature: Room Temperature;

The wave length of determination: 265 nm.

[Storage]

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

[References]

[1] Pan C H, Kim E S, Sang H J, et al. Arch. Pharm. Res., 2008, 31(11):1447-56.

[2] Thelen P, Scharf J G, Burfeind P, et al. Carcinogenesis, 2005, 26(8):1360-7.

[3] Lee K T, Sohn I C, Dong H K, et al. Arch. Pharm. Res., 2000, 23(5):461-6.

[4] Genead R, Fischer H, Hussain A, et al. Carcinogenesis, 2012, 33(12):2488-98.

[5] Lee H U, Bae E A, Kim D H. J. Pharmacol. Sci., 2005, 97(97):541-4.

[6] Zhang H, Liu X, Shi C, et al. J. Ethnopharmacol., 2010, 131(1):165-73.

[7] Zhao N, Liu D, Wang Y P, et al. J. Pharm. Practice, 2012, 3.

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