

Fisetin Datasheet

5th Edition (Revised in January, 2017)

[Product Information]

Name: Fisetin

Catalog No.: CFN98176

Cas No.: 528-48-3

Purity: >=98%

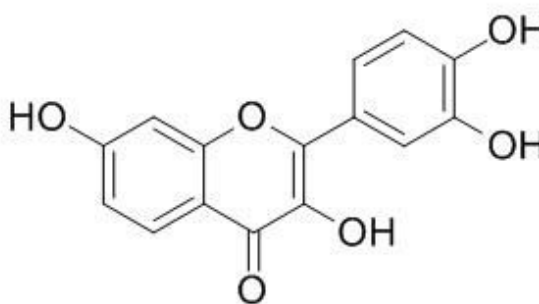
M.F: C₁₅H₁₀O₆

M.W: 286.24

Physical Description: Yellow cryst.

Synonyms: 4H-1-Benzopyran-4-one, 2-(3,4-dihydroxyphenyl)-3,7-dihydroxy-;

3,3',4',7-Tetrahydroxyflavone.



[Intended Use]

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

[Source]

The herbs of *Rhus succedanea* L.

[Biological Activity or Inhibitors]

Fisetin inhibits TH2-type cytokine production by activated human basophils.^[1]

Fisetin functions as a dual inhibitor of mTORC1/2 signaling leading to inhibition of prostate cancer (CaP)-dependent translation and induction of autophagic cell death in PC3 cells, suggests that fisetin could be a useful chemotherapeutic agent in treatment of hormone refractory CaP.^[2]

Fisetin has anti-inflammatory activity in human mast cells (HMC-1) through the down-regulation of mast cell activation.^[3]

Fisetin is an inhibitor of androgen receptor (AR) signaling axis, it could be a useful chemopreventive and chemotherapeutic agent to delay progression of prostate cancer.^[4]

Fisetin could be useful in attenuation of UV radiation-induced oxidative stress and the activation of NF-kappaB and MAPK signaling in human lens epithelial cells, which suggests that fisetin has a potential protective effect against cataractogenesis.^[5]

Fisetin has direct antioxidant activity, it can also increase the intracellular levels of glutathione (GSH), the major endogenous antioxidant, and fisetin has both neurotrophic and anti-inflammatory activity.^[6]

Fisetin shows in vitro antifungal activity(MIC<128 ug/mL) and low toxicity(IC₅₀=158 ug/mL) on animal cells.^[7]

[Solvent]

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

[HPLC Method]^[8]

Mobile phase: 1% Formic acid in water- Acetonitrile, gradient elution;

Flow rate: 1.0 ml/min;

Column temperature: 40°C;

The wave length of determination: 275 nm.

[Storage]

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

[References]

- [1] Higa S, Hirano T, Kotani M, *et al. J. Allergy Clin. Immunol.*, 2003 Jun; 111(6):1299-306.
- [2] Suh Y S, Afaq F, Khan N, *et al. Carcinogenesis*, 2010, 31(8):1424-33.
- [3] Park H H, Lee S, Oh J M, *et al. Pharmacol. Res.*, 2007 Jan; 55(1):31-7.
- [4] Khan N, Asim M, Afaq F, *et al. Cancer Res.*, 2008, 68(20):8555-63.
- [5] Yao K, Zhang L, Zhang Y D, *et al. Molecular Vision*, 2008, 14(219-223):1865-71.
- [6] Chiruta C, Schubert D, Dargusch R, *et al. J. Med. Chem.*, 2012 Jan 12; 55(1):378-89.
- [7] Maysa Paula da Costa, Marize Campos Valadares Bozinis, Wanessa Machado Andrade, *et al. BMC Complement. Altern. Med.*, 2014; 14:245.
- [8] Osman A, Makris D P. *Int. J. Food Sci. Technol.*, 2010, 45(11):2265-71.

[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