

## Wogonin Datasheet

4<sup>th</sup> Edition (Revised in July, 2016)

### [ Product Information ]

**Name:** Wogonin

**Catalog No.:** CFN97089

**Cas No.:** 632-85-9

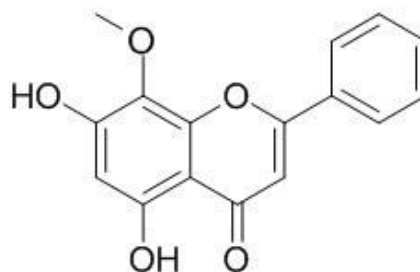
**Purity:** > 98%

**M.F:** C<sub>16</sub>H<sub>12</sub>O<sub>5</sub>

**M.W:** 284.3

**Physical Description:** Yellow powder

**Synonyms:** 5,7-Dihydroxy-8-methoxy-2-phenyl-1-benzopyran-4-one.



### [ Intended Use ]

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

### [ Source ]

The root of *Scutellaria baicalensis Georgi*.

## **[ Biological Activity or Inhibitors ]**

Wogonin is one major constituent of *Scutellaria baicalensis*, possesses potent anticancer activities, the activities are largely due to their abilities to scavenge oxidative radicals, to attenuate NF-kappaB activity, to inhibit several genes important for regulation of the cell cycle, to suppress COX-2 gene expression and to prevent viral infections. [1]

Wogonin and fisetin induce apoptosis in human promyeloleukemic cells, accompanied by a decrease of reactive oxygen species, and activation of caspase 3 and endonuclease, induction of bax protein and suppression of Mcl-1 protein were detected in the process. [2]

Wogonin, baicalin, and baicalein are three major components of *Scutellaria*, they have anti-inflammatory activity against carrageenan-induced paw edema in rats, the baicalin exhibits the greatest inhibition activity. [3]

Wogonin can suppress the VEGF-stimulated migration and tube formation of human umbilical vein endothelial cells (HUVECs), also restrain VEGF-induced tyrosine phosphorylation of vascular endothelial growth factor receptor 2 (VEGFR2), strongly suggests that wogonin might be a promising antitumor drug. [4]

Wogonin induces apoptosis in RPMI 8226, a human myeloma cell line, by downregulating phospho-Akt and overexpressing Bax. [5]

## **[ Solvent ]**

Chloroform, Dichloromethane, DMSO, Acetone, etc.

## **[ HPLC Method ]** [6]

Mobile phase: Methanol-0.1%Phosphoric acid H<sub>2</sub>O gradient elution ;

Flow rate: 1.0 ml/min;

Column temperature: 30 °C;

The wave length of determination: 275 nm.

## **[ Storage ]**

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

## **[ References ]**

- [1] Min L W. *New therapeutic aspects of flavones: Cancer Treatment Reviews*, 2009, 35(35):57-68.
- [2] Lee W R, Shen S C, Lin H Y, *et al. Biochem. Pharmacol.*, 2002, 63(2):225-36.
- [3] Chun-ChingLin, Den-EnShieh. *Am. J. Chinese Med.*, 1996, 24(1):31-6.
- [4] Na L, Ying G, Yun L, *et al. Life Sci.*, 2008, 82(17-18):956-63.
- [5] Zhang M, Liu L P, Chen Y, *et al. Life Sci.*, 2013, 92(1):55-62.
- [6] Lin D, Liang W. *China Pharmacist*, 2009, 12(08):1044-6.

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