

# **Isoginkgetin Datasheet**

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

### [ Product Information ]

Name: Isoginkgetin

Catalog No.: CFN90174

Cas No.: 548-19-6

**Purity:** > 98%

M.F: C<sub>32</sub>H<sub>22</sub>O<sub>10</sub>

M.W: 566.51

Physical Description: Yellow cryst.

 $\textbf{Synonyms:} 8\text{-}[5\text{-}(5,7\text{-}dihydroxy\text{-}4\text{-}oxo\text{-}1\text{-}benzopyran\text{-}2\text{-}yl)\text{-}2\text{-}methoxyphenyl}]\text{-}5,7\text{-}dihydroxylloopyran\text{-}2\text{-}yl)\text{-}2\text{-}methoxyphenyl}$ 

y-2-(4-methoxyphenyl)-1-benzopyran-4-one.

## [ Intended Use ]

- 1. Reference standards:
- 2. Pharmacological research;
- 3. Food research;
- 4. Cosmetic research;
- 5. Synthetic precursor compounds;
- 6. Care and daily chemicals;
- 7. Intermediates & Fine Chemicals:
- 8. Ingredient in supplements, beverages;
- 9. Others.

[Source]

The leaves of Ginkgo biloba L..

[ Biological Activity or Inhibitors]

Isoginkgetin can inhibits tumor cell invasion by regulating phosphatidylinositol

3-kinase/Akt-dependent matrix metalloproteinase-9 expression.[1]

Isoginkgetin is a general inhibitor of both the major and minor spliceosomes, it inhibits

splicing both in vivo and in vitro at similar micromolar concentrations; isoginkgetin has

been previously described as an anti-tumor agent, suggest that splicing inhibition is the

mechanistic basis of the anti-tumor activity of isoginkgetin, thus, pre-mRNA splicing

inhibitors may represent a novel avenue for development of new anti-cancer agents.[2]

Isoginkgetin enhances adiponectin secretion from differentiated adiposarcoma cells via a

novel pathway involving AMP-activated protein kinase.[3]

Isoginkgetin (0.3mg/kg ip for 3d) can reduce the level of 0 0~-in plasma and erythroeyte

(p0.01) and sometimes increase the activity of SOD in anoxic rats, the action being

stronger than aspirin.[4]

[Solvent]

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

[ HPLC Method ]<sup>[5]</sup>

Mobile phase: Acetonitrile- H2O, gradient eiution;

Flow rate: 1.0 ml/min;

Column temperature: 30 ℃;

The wave length of determination: 270 nm.

[Storage]

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

#### [References]

[1] Yoon S O, Shin S, Lee H J, et al. Mol. Cancer Ther., 2006, 5(11):2666-75.

[2] O'Brien K, Matlin A J, Lowell A M, et al. J. Biol. Chem., 2008, 283(48):33147-54.

[3] Liu G, Grifman M, Macdonald J, et al. J. Endocrinol., 2007, 194(3):569-78.

[4] Pan S. Traditional Chinese Drug Research & Clinical Pharmacology, 1993(2):12-4

[5] Li B, Hu G S, Hu L L, et al. Chinese herbal medicine, 2014, 45 (17): 2552-5.

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