**Natural Products** 

OH

OH



# **Sappanone A Datasheet**

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

#### [ Product Information ]

Name: Sappanone A

Catalog No.: CFN92014

Cas No.: 102067-84-5

**Purity:** > 95%

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

**M.W:** 284.3

Physical Description: Cryst.

**Synonyms:**3-[(3,4-Dihydroxyphenyl)methylene]-2,3-dihydro-7-hydroxy-4H-1-benzopyran -4-one.

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#### [Intended Use]

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

### [Source]

The heartwoods of Caesalpinia sappan.

#### [Biological Activity or Inhibitors]

Sappanone A has melanogenesis inhibitory activity, it can dose-dependently inhibit both melanogenesis and cellular tyrosinase activity via repressing tyrosinase gene expression in mouse B16 melanoma cells.<sup>[1]</sup>

Sappanone A exerts its anti-inflammatory effect by modulating the Nrf2 and NF-κB pathways, and may be a valuable compound to prevent or treat inflammatory diseases.<sup>[2]</sup> Sappanone A has antioxidant and anti-inflammatory effects, treatment of sappanone A can improve cisplatin (CP)-induced histopathalogical injury and renal dysfunction through activating Nrf2 and inhibiting NF-κB activation, it is a potential therapeutic drug for treating CP-induced kidney injury.<sup>[3]</sup>

Sappanone A can inhibit ovalbumin (OVA)--induced asthma by activating the Nrf2 signaling pathway, it may have a potential use as a therapeutic agent for asthma.<sup>[4]</sup> Sappanone A can significantly attenuate endothelial cell damage induced by H<sub>2</sub>O<sub>2</sub>.<sup>[5]</sup>

#### [Solvent]

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

#### [ HPLC Method ]

Not data available.

#### [Storage]

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

#### [References]

- [1] Chang T S, Chao S Y, Ding H Y. Int. J. Mol. Sci., 2012, 13(8):10359-67.
- [2] Lee S, Choi S Y, Choo Y Y, et al. Int. Immunopharmacol., 2015, 28(1):328-36.
- [3] Kang L, Zhao H, Chen C, et al. Int. Immunopharmacol., 2016, 38:246-51.
- [4] Liu X, Yu D, Wang T. Int. Arch. Allergy Imm., 2016; 170(3):180-6.
- [5] Zhao H X, Bai H, Li W, et al. Food & Drug, 2010, 12(5):176-80.

## [ Contact ]

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