

# **Angelicin Datasheet**

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

#### [ Product Information ]

Name: Angelicin

Catalog No.: CFN98854

Cas No.: 523-50-2

**Purity:** > 98%

M.F: C<sub>11</sub>H<sub>6</sub>O<sub>3</sub>

M.W: 186.2

Physical Description: Powder

**Synonyms:** 2-Furo[2,3-h][1]benzopyranone; Isopsoralen.

#### [ Intended Use ]

1. Reference standards;

2. Pharmacological research;

3. Synthetic precursor compounds;

4. Intermediates & Fine Chemicals;

5. Others.

#### [Source]

The seeds of Psoralea grandulosa.

## [ Biological Activity or Inhibitors]

Angelicin is a furocoumarin found in Psoralea corylifolia L. fruit, can block the

phosphorylation of IκBα, NFκBp65, p38 MAPK, and JNK in lipopolysaccharide-induced

acute lung injury model, suggests that angelicin was potentially advantageous to prevent

inflammatory diseases by inhibiting NF-κB and MAPK pathways, it might be a potential

new agent for prevention of inflammatory reactions and diseases in the clinic.[1]

Angelicin, compared with cytosine arabinoside, mithramycin and cisplatin, is a powerful

inducer of erythroid differentiation and -globin mRNA accumulation of human leukemia

K562 cells, it is a potential therapeutic approach in hematological disorders, including

-beta-thalassemia and sickle cell anemia.[2]

A novel angelicin derivative 6a was identified to inhibit influenza A (H1N1) virus induced

Cytopathic effect in Madin-Darby canine kidney cell culture in low micromolar range, these

compounds act as anti-influenza agents by inhibiting ribonucleoprotein (RNP) complex

associated activity and have the potential to be developed further, which could form the

basis for developing additional defense against influenza pandemics.[3]

Angelicin is structurally related to psoralens, a well-known chemical class of

photosensitizers used for its antiproliferative activity in treatment of different skin disease,

angelicin is an effective apoptosis-inducing natural compound of human SH-SY5Y

neuroblastoma cells which suggests that this compound may have a role in future

therapies for human neuroblastoma cancer.[4]

[Solvent]

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

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

Mobile phase: Methanol: H2O=52:48;

Flow rate: 1.0 ml/min:

Column temperature: Room Temperature;

The wave length of determination: 246 nm.

### [Storage]

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

#### [References]

[1] Liu F, Sun G Q, Gao H Y, et al. J. Surg. Res., 2013, 185(1):300-9.

[2] Lampronti I, Bianchi N M, Fibach E, et al. Eur. J. Haematol., 2003, 71(3):189-95.

[3] Yeh J, Coumar M, Horng J, et al. J. Med. Chem., 2010, 53(4):1519-33.

[4] Rahman M A, Kim N H, Yang H, et al. Mol. Cell Biochem., 2012, 369(1-2):95-104.

[5] Yan R, Liu Z, Luo J. Chinese Medicine Modern Distance Education of China, 2008,6(11):1329-30.

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