

## Demethylwedelolactone Datasheet

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

### [ Product Information ]

**Name:** Demethylwedelolactone

**Catalog No.:** CFN90521

**Cas No.:** 6468-55-9

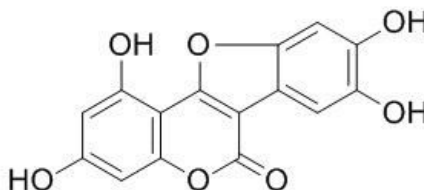
**Purity:** >=98%

**M.F:** C<sub>15</sub>H<sub>8</sub>O<sub>7</sub>

**M.W:** 300.22

**Physical Description:** Powder

**Synonyms:** 1,3,8,9-Tetrahydroxycoumestan;; Norwedelolactone; Isodemethyl-wedelolactone; 1,3,8,9-Tetrahydroxy-6H-benzofuro-[3,2-c]chromen-6-one.



### [ Intended Use ]

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

### [ Source ]

The herbs of *Eclipta prostrata*.

### [ Biological Activity or Inhibitors ]

Demethylwedelolactone and wedelolactone can inhibit the anchorage-independent growth and suppress cell motility and cell invasion of MDA-MB-231 cells, they also can reduce the activity and expression of matrix metalloproteinases (MMPs) involved in blocking the I $\kappa$ B- $\alpha$ /NF $\kappa$ B and MEK/ERK signaling pathways in MDA-MB-231 cells; suggests that demethylwedelolactone derivatives exert anti-invasive growth effect on breast cancer cells.<sup>[1]</sup>

Demethylwedelolactone and wedelolactone have trypsin inhibitory effect, IC(50) values are 3.0 and 2.9 microg/mL respectively.<sup>[2]</sup>

Demethylwedelolactone and wedelolactone can dose dependently inhibit the de-granulation of mast cells induced by Compound 48/80 (C 48/80), they also inhibit the production of NO, pro-inflammatory cytokines such as TNF- $\alpha$ , IL-1 $\beta$  and IL-6 and the expression of costimulatory molecules such as CD40, CD80 and CD86 in LPS-stimulated macrophages; suggests they have immunomodulatory effects, can be exploited as alternative new therapeutics for various inflammatory diseases.<sup>[3]</sup>

Demethylwedelolactone is the major constituent of the butanolic and purified butanolic extracts (PBEs) of *Eclipta prostrata*, has anti-venom potential; both extracts partially inhibit the hemorrhagic activity but display very low anti-phospholipase A activity and do not inhibit proteolytic activity of Malayan pit viper (MPV) venom.<sup>[4]</sup>

## **[ Solvent ]**

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

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

Mobile phase: Methanol- 0.2% Formic acid in water=57:43;

Flow rate: 1.0 ml/min;

Column temperature: 30 °C;

The wave length of determination: 351 nm.

## **[ Storage ]**

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

## **[ References ]**

- [1] Lee Y J, Lin W L, Chen N F, *et al. Eur. J.Med.Chem.*, 2012, 56(10):361-7.
- [2] Syed S D, Deepak M, Yogisha S, *et al. Phytother. Res.*, 2003, 17(4):420-1.
- [3] Maji A K, Mahapatra S, Banerji P, *et al. Oriental Pharmacy & Experimental Medicine*, 2015, 15(1):23-31.
- [4] Pithayanukul P, Laovachirasuwan S, Bavovada R, *et al. J. Ethnopharmacol.*, 2004, 90(2-3):347-52.
- [5] Shailajan S, Menon S, Singh D, *et al. Pharmacogn. J.*, 2016, 8(2):132-9.

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