

## Damnacanthal Datasheet

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

### [ Product Information ]

**Name:** Damnacanthal

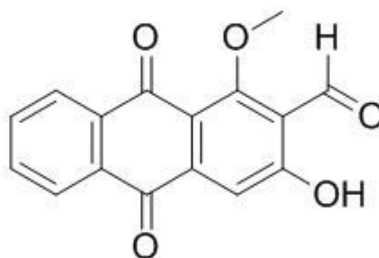
**Catalog No.:** CFN98723

**Cas No.:** 477-84-9

**Purity:** > 95%

**M.F:** C<sub>16</sub>H<sub>10</sub>O<sub>5</sub>

**M.W:** 282.25



**Physical Description:** Yellow powder

**Synonyms:** 3-Hydroxy-1-methoxy-9,10-dioxo-9,10-dihydro-anthracene-2-carbaldehyde.

### [ Intended Use ]

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

### [ Source ]

The fruit of *Morinda citrifolia* Linn.

### [ Biological Activity or Inhibitors ]

Damnacanthal, an anthraquinone compound, is isolated from the roots of *Morinda citrifolia*

L. (noni), which has been used for traditional therapy in several chronic diseases including cancer, damnacanthal increases antitumorigenic activity in human colorectal cancer cells and that C/EBP $\beta$  plays a role in damnacanthal-induced NAG-1 expression.<sup>[1]</sup>

Damnacanthal is a highly potent, selective inhibitor of p56lck tyrosine kinase activity.<sup>[2]</sup>

Damnacanthal activates p38 MAPK by mediates apoptosis in SKHep 1 cells through the DR5/TRAIL and TNFR1/TNF-alpha and p53 pathways. <sup>[3]</sup>

Damnacanthal has antinociceptive and anti-inflammatory actions in mice.<sup>[4]</sup>

Damnacanthal can act as an immunomodulatory agent which may be very useful for maintaining a healthy immune system.<sup>[5]</sup>

## **[ Solvent ]**

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

## **[ HPLC Method ]<sup>[6]</sup>**

Mobile phase: 0.5% Acetic acid in water- Acetonitrile, gradient elution ;

Flow rate: 1.3 ml/min;

Column temperature: 30 °C;

The wave length of determination: 250 nm.

## **[ Storage ]**

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

## **[ References ]**

[1] Nualsanit T, Rojanapanthu P, Gritsanapan W, *et al. J.Nutr. Biochem.*, 2012, 23(8): 915-23.

[2] Faltynek C R, Schroeder J, Mauvais P, *et al. Biochemistry*, 1995, 34(38):12404-10.

[3] Lin F L, Hsu J L, Chou C H, *et al. Eur. J.Pharmacol.*, 2011, 650(1):120-9.

[4] Okusada K, Nakamoto K, Nishida M, *et al. Biol. Pharmaceut. Bull.*, 2011, 34(1):103-7.

[5] Alitheen N B, Manaf A A, Yeap S K, *et al. Pharm. Biol.*, 2010, 48(4):446-52.

[6] Kiathevest K, Goto M, Sasaki M, *et al. Sep. Purif. Technol.*, 2009, 66(1):111-7.

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