

# **Dracorhodin perchlorate Datasheet**

5<sup>th</sup> Edition (Revised in January, 2017)

## [ Product Information ]

Name: Dracorhodin perchlorate

Catalog No.: CFN90486

Cas No.: 125536-25-6

**Purity: >=98%** 

M.F: C<sub>17</sub>H<sub>15</sub>ClO<sub>7</sub>

M.W: 366.75

Physical Description: Red powder

**Synonyms:** Methane, 5-methoxy-6-methyl-2-phenylchromenylium-7-ol.

#### [ Intended Use ]

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

#### [Source]

The herbs of Daemonorops draco Bl.

### [ Biological Activity or Inhibitors]

Dracorhodin perchlorate can inhibit PI3K/Akt and NF-kB activation, up-regulate the

expression of p53, and enhance apoptosis in cancer cells [1]

Dracorhodin perchlorate alters the intracellular redox status, changes the balance of

Bcl-X(L) and Bax protein expression, and induces apoptosis through caspase pathways in

HeLa cells.[2]

Dracorhodin perchlorate can inhibit high glucose-induced connective tissue growth factor

expression in human mesangial cells, and this may be its mechanism of prevention and

treatment on renal fibrosis in diabetic nephropathy (DN). [3]

Dracorhodin perchlorate has been used as a medicine to treat chronic wounds, the effects

of it on wound healing have association with the Ras/MAPK signaling pathway.[4]

[Solvent]

Pyridine, Methanol, Ethanol, etc.

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

Mobile phase: Acetonitrile-0.05 M Sodium dihydrogen phosphate solution=37:63;

Flow rate: 1.0 ml/min;

Column temperature: 30 ℃;

The wave length of determination: 440 nm.

[Storage]

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

[References]

[1] Rasul A, Ding C, Li X,et al. Apoptosis, 2012, 17(10):1104-19.

[2] Xia M, Wang D, Wang M, et al. J. Pharmacol. Sci., 2004, 95(2):273-83.

[3] Wang Y H, Wang Q S, Liu J G, et al. Zhongguo Zhong yao za zhi, 2009, 34(7):896-9.

[4] Li F, Jiang T, Liu W, et al. Mol. Med. Rep., 2016, 14(2):1667-72.

[5] He Y, Ding N, Wang R Z, et al. Journal of Chinese Pharmaceutical Sciences, 2015,

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