



# Ginsenoside Rg2 Datasheet

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

## [ Product Information ]

**Name:** Ginsenoside Rg2

**Catalog No.:** CFN99968

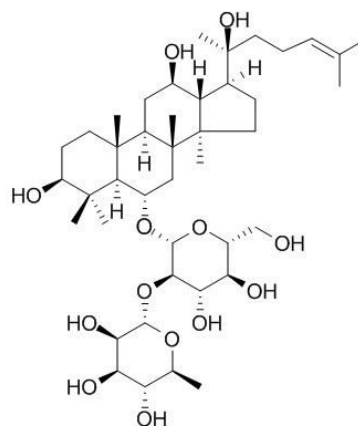
**Cas No.:** 52286-74-5

**Purity:** > 98%

**M.F:** C<sub>42</sub>H<sub>72</sub>O<sub>13</sub>

**M.W:** 785.02

**Physical Description:** White powder



**Synonyms:** (2S,3R,4R,5R,6S)-2-[[[(2R,3R,4S,5S,6R)-2-[[[(6R,10R,12S,14R)-3,12-dihydroxy-17-[(E,2S)-2-hydroxy-6-methyloct-5-en-2-yl]-4,4,10,14-tetramethyl-1,2,3,5,6,7,8,9,11,12,13,15,16,17-tetradecahydrocyclopenta[a]phenanthren-6-yl]oxy]-4,5-dihydroxy-6-(hydroxymethyl)-

## [ Intended Use ]

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

## **[ Source ]**

The roots of *Panax ginseng* C. A. Mey.

## **[ Biological Activity or Inhibitors ]**

Ginsenoside Rg2 suppresses the hepatic glucose production via AMPK-induced phosphorylation of GSK3 $\beta$  and induction of SHP gene expression, suggests that it has therapeutic potential for type 2 diabetic patients.<sup>[1]</sup>

Ginsenoside Rg2 inhibits nicotinic acetylcholine receptor-mediated Na<sup>+</sup> influx and channel activity; it also inhibits the 5-HT-induced inward peak current (I<sub>5-HT</sub>) in dose dependent and reversible manner, the half-inhibitory concentrations (IC<sub>50</sub>) of ginsenoside Rg2 is 22.3  $\pm$  4.6  $\mu$ M, suggests that it might regulate the 5-HT<sub>3A</sub> receptors that are expressed in *Xenopus* oocytes.<sup>[2]</sup>

Ginsenoside Rg2 can reduce LPS-mediated THP-1 monocyte adhesion to HUVEC, in a concentration-dependent manner, it may provide direct vascular benefits with inhibition of leukocyte adhesion into vascular wall thereby providing protection against vascular inflammatory disease.<sup>[3]</sup>

Ginsenoside Rg2 protects cells against UVB-induced genotoxicity by increasing DNA repair, in possible association with modulation of protein levels involved in p53 signaling pathway.<sup>[4]</sup>

Ginsenoside Rg2 improves learning and memory through mechanisms related to anti-apoptosis in MID rats, indicates that it may represent a potential neurorestorative treatment strategy for vascular dementia or other ischemic insults.<sup>[5]</sup>

Ginsenoside Rg2 has protective effects against H<sub>2</sub>O<sub>2</sub>-induced injury and apoptosis in H9c2 cells.<sup>[6]</sup>

## **[ Solvent ]**

Pyridine, Methanol, Ethanol, Hot water, etc.

## **[ HPLC Method ]<sup>[7]</sup>**

Mobile phase: Methanol -H<sub>2</sub>O=65:35 ;

Flow rate: 1.0 ml/min;

Column temperature: 25 °C;

The wave length of determination: 203 nm.

## **[ Storage ]**

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

## **[ References ]**

- [1] Yuan H D, Kim D Y, Quan H Y, *et al. Chem.-Biol. Interact.*, 2012, 195(1):35-42.
- [2] Choi S, Lee J H, Oh S, *et al. Mol. Cells*, 2003, 15(1):108-13.
- [3] Cho Y S, Chan H K, Ha T S, *et al. Korean J. Physiol. Pham.* 2013, 17(2):133-7.
- [4] Ha S E, Shin D H, Kim H D, *et al. N.-S. Arch. Pharmacol.*, 2010, 382(1):89-101.
- [5] Gong Z H, Liu M X, Gong L L, *et al. Prog. in Modern Biomed.*, 2010,10(06):1069-75.
- [6] Fu W, Sui D, Yu X, *et al. Int. J. Clin. Exp. Med.*, 2015, 8(11):19938-47.
- [7] Yu M, Mi H, Jiao L. *China Pharmacist*, 2005, 8(12):1017-9.

## **[ Contact ]**

### **Address:**

S5-3 Building, No. 111, Dongfeng Rd.,  
Wuhan Economic and Technological Development Zone,  
Wuhan, Hubei 430056,  
China

**Email:** [info@chemfaces.com](mailto:info@chemfaces.com)

**Tel:** +86-27-84237783

**Fax:** +86-27-84254680

**Web:** [www.chemfaces.com](http://www.chemfaces.com)

**Tech Support:** [service@chemfaces.com](mailto:service@chemfaces.com)