

Ginsenoside Rb3 Datasheet

4th Edition (Revised in July, 2016)

[Product Information]

Name: Ginsenoside Rb3

Catalog No.: CFN99966

Cas No.: 68406-26-8

Purity: > 98%

M.F: $C_{53}H_{90}O_{22}$

M.W: 1079.27

Physical Description: White powder

Synonyms: Gypenoside IV; β -D-Glucopyranoside, $(3\beta,12\beta)$ -3-[(2-O- β -D-

[3β-[[2-O-(β-D-Glucopyranosyl)-β-D-glucopyranosyl]oxy]-12β-hydroxy-5α-dammar-24-en-

20-yl]6-O-(β -D-xylopyranosyl)- β -D-glucopyranoside .

[Intended Use]

- 1. Reference standards;
- 2. Pharmacological research;
- 3. Food and cosmetic research;
- 4. Synthetic precursor compounds;
- 5. Intermediates & Fine Chemicals;
- 6. Ingredient in supplements, beverages;
- 7. Aromatics:
- 8. Others.

[Source]

The root and rhizome of Panax ginseng C. A. Mey.

[Biological Activity or Inhibitors]

Ginsenoside Rb3 has protective effects on oxygen and glucose deprivation-induced ischemic injury in PC12 cells.^[1]

Ginsenoside Rb3 possesses the effect against isoproterenol-induced myocardial injury and heart function impairment, and that the mechanism of pharmacological action was related to the antioxidant activity of ginsenoside Rb3 at least in part. [2]

Ginsenoside Rb3 is extracted from the plant Panax ginseng and plays important roles in cardiovascular diseases, including myocardial ischemia-reperfusion (I/R) injury, the protective effect of ginsenoside Rb3 on the OGD-Rep injury is attributed to the inhibition of JNK-mediated NF-kB activation, suggesting that ginsenoside Rb3 has the potential to serve as a novel therapeutic agent for myocardial I/R injury.^[3]

Ginsenoside Rb3 may have antidepressant-like effects, brain-derived neurotrophic factor and the monoamine neurotransmitters 5-hydroxytryptamine, dopamine, and norepinephrine are involved in ginsenoside Rb3's antidepressant-like effects. [4]

Ginsenoside Rb3 significantly attenuates the changes of creatine kinase activity and lactate dehydrogenase activity.^[5]

Ginsenoside Rb3 can exert a neuroprotective role on hippocampal neurons, a role which was partly mediated by the facilitation of Ca2+-dependent deactivation of NMDA receptors, and the resultant reduction of intracellular free Ca2+ level.^[6]

Ginsenoside Rb3 reduces fasting blood glucose level, food intake, water intake, improved oral glucose tolerance, and repaired injured pancreas tissues of alloxan-induced diabetic mice, suggests that ginsenoside possesses the potential of the clinical use in preventing and treating diabetes.^[7]

[Solvent]

Pyridine, DMSO, Ethanol, Methanol.

[HPLC Method][8]

Mobile phase: Acetonitrile-0.2% Phosphoric acid H2O, gradient eiution;

Flow rate: 1.0 ml/min;

Column temperature: 40 °C;

The wave length of determination: 203 nm.

[Storage]

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

[References]

[1] Jun-rong, Yi-fu, Shen, et al. Acta Pharmacol. Sin., 2010, 31(3):273-80.

[2] Tian W, Yu X, Qu S, et al. Eur. J. Pharmacol., 2010, 636(1-3):121-5.

[3] Ma L, Liu H, Xie Z, et al. Plos One, 2014, 9(8):e103628-e103628.

[4] Cui J, Jiang L, Xiang H. J. Psychopharmacol., 2011, 26(5):697-713.

[5] Shi Y, Han B, Yu X, et al. Pharm. Biol., 2011, 49(9):900-6.

[6] Peng L L, Hong M S, Zheng L J, et al. Am. Chinese Med., 2009, 37(4):759-70.

[7] Bu Q T, Zhang W Y, Chen Q C, et al. Med. Chem., 2012, 8(5):934-41.

[8] Yin S, Wu H, Xu F, et al. Acta Academiae Medicinae Militaris Tertiae, 2010, 32(7):658-60.

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