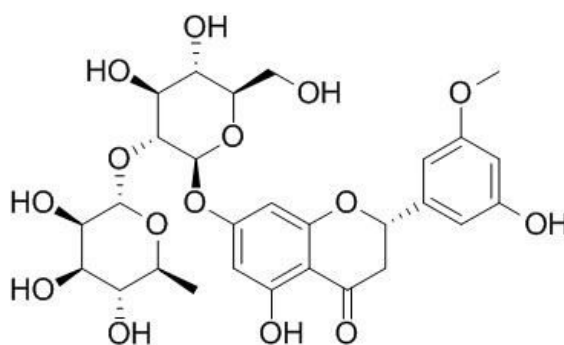


Neohesperidin Datasheet

4th Edition (Revised in July, 2016)**[Product Information]****Name:** Neohesperidin**Catalog No.:** CFN99125**Cas No.:** 13241-33-3**Purity:** > 98%**M.F:** C₂₈H₃₄O₁₅**M.W:** 610.56**Physical Description:** Powder

Synonyms: (2S)-7-[[[(2S,3R,4S,5S,6R)-4,5-dihydroxy-6-(hydroxymethyl)-3-[[[(3R,4R,5R,6S)-3,4,5-trihydroxy-6-methyl-2-oxanyl]oxy]-2-oxanyl]oxy]-5-hydroxy-2-(3-hydroxy-4-methoxyphenyl)-3,4-dihydro-2H-1-benzopyran-4-one.

[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. Others.

[Source]

The peel of *Citrus aurantium* L.

[Biological Activity or Inhibitors]

Neohesperidin is a natural new nutrition sweetener, widely existing in plants of dry citrus peel, which can be derived from extraction; since the sweetness is 1,300-1,500 times greater than that of sugar, neohesperidin are widely used in fruit juices, wines, beverages, bakeries and pharmaceutical formulations, and are particularly suitable for consumption by diabetic patients.^[1]

Neohesperidin exhibits antioxidant activity ($IC_{50}=22.31\mu\text{g/mL}$) in the 1,1-diphenyl-2-picrylhydrazyl (DPPH) radical-scavenging assay; neohesperidin (50mg/kg) significantly inhibits 55.0% of HCl/ethanol-induced gastric lesions, and increases the mucus content; In pylorus ligated rats, neohesperidin (50 mg/kg) significantly decreases the volume of gastric secretion and gastric acid output, and increases the pH; suggests that neohesperidin isolated from PF may be useful for the treatment and/or protection of gastritis. ^[2]

Neohesperidin has free radical scavenging activity, can induce cellular apoptosis in human breast adenocarcinoma MDA-MB-231 cells via activating the Bcl-2/Bax-mediated signaling pathway.^[3]

Neohesperidin can attenuate cerebral ischemia-reperfusion injury via the inhibition of neuronal and oxidative stress through the regulation of the apoptotic pathway and activating the Akt/Nrf2/HO-1 pathway.^[4]

Neohesperidin and hesperidin are the major flavanones isolated from bittersweet orange, they have potent anti-inflammatory effects in various inflammatory models, they significantly aggravate gastric damage caused by indomethacin administration as evidenced by increased ulcer index and histopathological changes of stomach.^[5]

Neohesperidin, Albiflorin, and Aloeemodin have a potent inhibitory effect on A β 1-40 and A β 1-42 aggregation, and have neuroprotective effect on primary hippocampal cells against β -Amyloid induced toxicity.^[6]

[Solvent]

Pyridine, DMSO, Ethanol, Methanol, Hot water.

[HPLC Method]^[7]

Mobile phase: Acetonitrile- 0.2% Formic acid H₂O, gradient elution ;

Flow rate: 1.0 ml/min;

Column temperature: 40 °C;

The wave length of determination: 285 nm.

[Storage]

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

[References]

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- [3] Xu F, Zang J, Chen D, *et al.* *Nat. Prod .Commun.*, 2012, 7(11):1475-8.
- [4] Wang J J, Cui P. *J. Asian Nat. Prod. Res.*, 2013, 15(9):1023-37.
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- [6] Ho S L, Poon C Y, Lin C, *et al.* *Curr. Alzheimer Res.*, 2015, 12(5):424-33.
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