



trans-Hinokiresinol Datasheet

5th Edition (Revised in January, 2017)

[Product Information]

Name: trans-Hinokiresinol

Catalog No.: CFN99834

Cas No.: 17676-24-3

Purity: > 95%

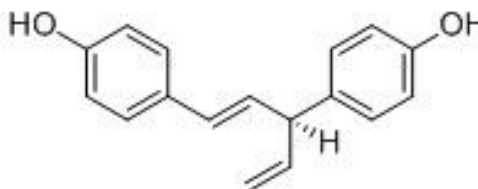
M.F: C₁₇H₁₆O₂

M.W: 252.3

Physical Description: Powder

Synonyms: 4-[(3S)-1-(4-hydroxyphenyl)penta-1,4-dien-3-yl]phenol;

Hinokiresinol; 4,4'-(1Z,3S)-penta-1,4-diene-1,3-diylldiphenol.



[Intended Use]

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

[Source]

The barks of *Cryptomeria japonica*.

[Biological Activity or Inhibitors]

Hinokiresinol (trans-hinokiresinol) and nyasol (cis-hinokiresinol) possess appreciable estrogen receptor binding activity, they can stimulate the proliferation of estrogen-dependent T47D breast cancer cells, and their stimulatory effects could be blocked by an estrogen antagonist, indicating that they are estrogen agonists.^[1]

trans- and cis-Hinokiresinols have similar free radical scavenging and anti-inflammatory activities, they also have anti-ischemic effects, only trans-hinokiresinol can significantly decrease neuronal injury in cultured cortical neurons exposed to oxygen-glucose deprivation followed by re-oxygenation.^[2]

Hinokiresinol is a novel inhibitor of LTB₄ binding to the human neutrophils. ^[3]

Hinokiresinol has antiallergic effect, it inhibits IgE-induced mouse passive cutaneous anaphylaxis reaction.^[4]

[Solvent]

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

[HPLC Method]^[5]

Mobile phase: Acetonitrile-0.03%Phosphoric acid H₂O, gradient elution;

Flow rate: 0.5 ml/min;

Column temperature: 30 °C;

The wave length of determination: 210 nm.

[Storage]

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

[References]

[1] Minami E, Taki M, Takaishi S, *et al. Chem. Pharm. Bull.*, 2000, 48(3):389-92.

[2] Ju C, Hwang S, Cho G S, *et al. Neuropharmacology*, 2013, 67(4):465-75.

[3] Lee H J, Ryu J H. *Planta Med.*, 1999, 65(4):391.

[4] Bae E, Park E H, Baek N, *et al. Planta Med.*, 2006, 72(14):1328-30.

[5] Luo J, Shi S H , Zhang L Q, *et al. Chinese Pharmaceutical Journal*, 2012, 47(22):
1856-9.

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