

N-p-trans-Coumaroyltyramine Datasheet

4th Edition (Revised in July, 2016)

[Product Information]

Name: N-p-trans-Coumaroyltyramine

Catalog No.: CFN98494

Cas No.: 36417-86-4

Purity: > 95%

M.F: C₁₇H₁₇NO₃

M.W: 283.3

Physical Description: Powder

Synonyms:

HO NH OH

[Intended Use]

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

[Source]

The herbs of Exochorda racemosa.

[Biological Activity or Inhibitors]

N-p-coumaroyltyramine, an α-glucosidase inhibitor, isolated from methanol extracts of

Welsh onion (Allium fistulosum), the inhibitory activity of it against a yeast enzyme is as

high as Ki 8.4×10⁻⁷ M.^[1]

N-p-coumaroyl tyramine is an inhibitor on acetylcholinesterase (AChE), it inhibits AChE

activity in a dose-dependent manner with IC₅₀ value of 34.5 microg/mL (122 microM).^[2]

N-trans-p-coumaroyltyramine exhibits potent inhibition of cell proliferation, platelet

aggregation, and shows antioxidant activity. [3]

N-trans-p-coumaroyltyramine and N-trans-pcoumaroyloctopamine exhibit a strong

suppressive effect on phagocytosis response upon activation with serum opsonized

zymosan in the range of $IC_{50} = 0.5-7.2$ uM, they display weak cytotoxic activity against the

human Caucasian prostate adenocarcinoma cell line PC-3, with IC₅₀ values ranging from

69.8 to 99.0 uM.[4]

N-trans-p-coumaroyltyramine, N-trans-caffeoyltyramine, and N-trans-feruloyltyramine as

the main active constituents of a methanolic extract from aerial parts of Polygonum

hyrcanicum (Polygonaceae) show activity against Trypanosoma brucei rhodesiense

(IC50s ranging from 2.2 to 13.3 microM).^[5]

[Solvent]

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

[HPLC Method]^[6]

Mobile phase: Methanol- 0.1% Formic acid in water, gradient elution;

Flow rate: 1.0 ml/min;

Column temperature: Room Temperature;

The wave length of determination: 300 nm.

[Storage]

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

[References]

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- [3] Neelam S, Gokara M, Sudhamalla B, et al. J. Phys. Chem. B, 2010, 114(8):3005-12.
- [4] Happi E N, Waffo A F, Wansi J D, et al. Planta Med., 2011, 77(9):934-8.
- [5] Moradi-Afrapoli F, Yassa N, Zimmermann S, et al. Nat. Prod. Commun., 2012, 7(6): 753-5.
- [6] Sun J, Gu Y F, Li M M, et al. China Journal of Chinese Materia Medica, 2014, 39(12):2300-4.

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