

Leachianone A Datasheet

4th Edition (Revised in July, 2016)

[Product Information]

Name: Leachianone A

Catalog No.: CFN97560

Cas No.: 97938-31-3

Purity: > 95%

M.F: C₂₆H₃₀O₆

M.W: 438.51

Physical Description: Powder

HO, OH OH OH

Synonyms:(2S)-5,7-Dihydroxy-2,3-dihydro-2 α -(2-methoxy-4-hydroxyphenyl)-8-[(R)-5-methyl-2-(1-methylethenyl)-4-hexenyl]-4H-1-benzopyran-4-one;(S)-5,7-Dihydroxy-2-(4-hydroxy-2-Methoxyphenyl)-8-((R)-5-Methyl-2-(prop-1-en-2-yl)hex-4-en-1-yl)chroMan-4-one.

[Intended Use]

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

[Source]

The roots of Sophora flavescens.

[Biological Activity or Inhibitors]

Leachianone A, isolated from Radix Sophorae, possesses a profound cytotoxic activity against human hepatoma cell line HepG2 in vitro, with an IC₍₅₀₎ value of 3.4microg/ml post-48-h treatment, its action mechanism via induction of apoptosis involved both extrinsic and intrinsic pathways.^[1]

Leachianone A is a potential antitoxic agent, it shows inhibitory effects on cadmium-Induced cytotoxicity, furthermore, leachianone A shows dose-dependency in detoxication. [2]

Leachianone A exhibits inhibitory activity against Sodium-dependent glucose cotransporter 2(SGLT2).^[3]

[Solvent]

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

[HPLC Method][3]

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

Flow rate: 9.0 ml/min;

Column temperature: Room Temperature;

The wave length of determination: 254 nm.

[Storage]

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

[References]

[1] Cheung S F, Chung K W, Lui C K, et al. Cancer Lett., 2007, 253(2):224-35.

[2] 양희태, 최화정, 백승화. 동의생리병리학회지, 2008, 22(5):1163-7.

[3] Yang J, Wang C, Lin Q, et al. Journal of Huazhong Normal University, 2014, 48(4):520-4.

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