

Demethylzeylasteral Datasheet

5th Edition (Revised in January, 2017)

[Product Information]

Name: Demethylzeylasteral

Catalog No.: CFN90136

Cas No.: 107316-88-1

Purity: >=98%

M.F: C₂₉H₃₆O₆

M.W: 480.60



Physical Description: Powder

Synonyms: (9beta, 13alpha, 14beta, 20alpha)-2, 3-Dihydroxy-9, 13-dimethyl-6, 23-dioxo-24,

25,26-trinoroleana-1,3,5(10),7-tetraen-29-oic acid;

A-Friedo-24-noroleana-1,3,5(10),7-tetraen-29-oicacid.

[Intended Use]

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

[Source]

The herbs of Tripterygium wilfordii Hook.f.

[Biological Activity or Inhibitors]

Demethylzeylasteral exhibits dose-dependent inhibitory behaviour towards estradiol glucuronidation.^[1]

Demethylzeylasteral exhibits strong inhibition towards UDP-glucuronosyltransferase (UGT) 1A6 and 2B7, and the inhibition kinetic parameters (Ki) are calculated to be 0.6 uM and 17.3 uM for UGT1A6 and UGT2B7, respectively.^[2]

Demethylzeylasteral has strong immunosuppressive activity, suggests a possible clinical use for demethylzeylasteral as an immunosuppressive agent in the fields of organ transplantation and autoimmune disorders. ^[3]

Demethylzeylasteral has antifertility activity, it concentration-dependently and in a partially reversible manner can inhibit the Ca(2+) current in spermatogenic cells with an IC(50) of 8.8 microg/ml, and it also can inhibit significantly the sperm acrosome reaction initiated by progesterone.^[4]

Demethylzeylasteral shows antimicrobial activity against Gram-positive bacteria and the yeast Candida albicans, it exhibits the fastest inhibition against cell wall synthesis.^[5]

[Solvent]

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

[HPLC Method]^[6]

Mobile phase: 0.1% Formic acid in water- Acetonitrile,gradient elution ; Flow rate: 1.0 ml/min; Column temperature: Room Temperature; The wave length of determination: 268 nm.

[Storage]

 $2-8^{\circ}$ C, Protected from air and light, refrigerate or freeze.

[References]

[1] Liu S L, Zhang S Y, Wang M J, et al. Eur. J. Drug Metab. Ph., 2014, 39(2):99-102.

[2] Zhao J W, Wang G H, Chen M, et al. Molecules, 2012, 17(8):9469-75.

[3] Xu W, Lin Z, Yang C, et al. Int. Immunopharmacol., 2009, 9(7-8):996-1001.

[4] Bai J P, Shi Y L, Fang X, et al. Eur. J. Pharmacol. ,2003,464(1):9-15.

[5] De L L, Beltrán B, Moujir L. Planta Med., 2005, 71(4):313-9.

[6] Wang T, Shen F, Su S, et al. Bmc Complem. Altern. M., 2016, 16(1):476.

[Contact]

Address:	Email: info@chemfaces.com
S5-3 Building, No. 111, Dongfeng Rd.,	Tel: +86-27-84237783
Wuhan Economic and Technological Development Zone,	Fax: +86-27-84254680
Wuhan, Hubei 430056,	Web: www.chemfaces.com
China	Tech Support: service@chemfaces.com