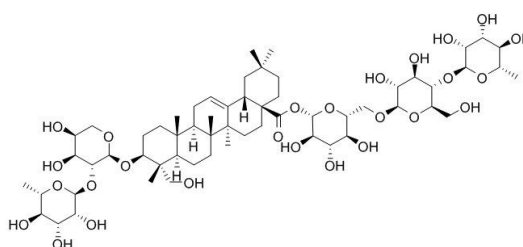


Hederacoside C Datasheet

4th Edition (Revised in July, 2016)**[Product Information]****Name:** Hederacoside C**Catalog No.:** CFN90183**Cas No.:** 14216-03-6**Purity:** >=98%**M.F:** C₅₉H₉₆O₂₆**M.W:** 1221.38**Physical Description:** White powder**Synonyms:** 6-Deoxy- α -L-mannopyranosyl-(1 \rightarrow 4)-D-glucopyranosyl-(1 \rightarrow 6)-1-O-[(3 β)-3-[[[(2 ξ)-2-O-(6-deoxy- α -L-mannopyranosyl)- α -L-erythro-pentopyranosyl]oxy]-23-hydroxy-28-oxoolean-12-en-28-yl]- β -D-glucopyranose; Kalopanaxsaponin B.**[Intended Use]**

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

[Source]The herbs of *Hedera nepalensis*.

[Biological Activity or Inhibitors]

Hederacoside C has antispasmodic activity.^[1]

Hederacoside C only non-competitively inhibits hyaluronidase activity in a dose-dependent fashion, shows comparable IC 50 value is 280.4 uM; it is a potent competitive inhibitor for serine protease porcine pancreatic elastase, shows comparable IC 50 value is 40.6 uM.^[2]

[Solvent]

Pyridine, Methanol, Ethanol, etc.

[HPLC Method]^[3]

Mobile phase: (H₂O: acetonitrile: orthophosphoric acid (85%)=860:140:2) - (Acetonitrile: orthophosphoric acid (85%)=998:2) ,gradient elution ;

Flow rate: 1.5 ml/min;

Column temperature: 40 °C;

The wave length of determination: 205 nm.

[Storage]

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

[References]

[1] Trute A, Gross J, Mutschler E, *et al. Planta Med.*, 1997, 63(2):125-9.

[2] Facino R M, Carini M, Stefani R, *et al. Arch. Pharm.*, 1995, 328(10):720-4.

[3] Khdaier A, Mohammad M K, Tawaha K, *et al. Int. J. Anal. Chem.*, 2010(10):478143.

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