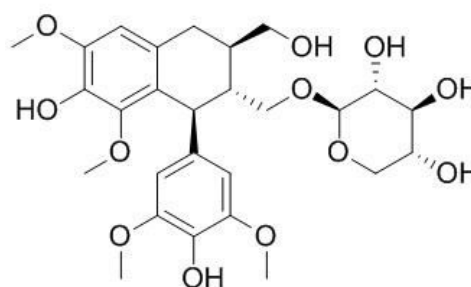


Lyoniside Datasheet

4th Edition (Revised in July, 2016)**[Product Information]****Name:** Lyoniside**Catalog No.:** CFN98451**Cas No.:** 34425-25-7**Purity:** > 98%**M.F:** C₂₇H₃₆O₁₂**M.W:** 552.6**Physical Description:** Powder**Synonyms:**[[[(1S)-1α-(3,5-Dimethoxy-4-hydroxyphenyl)-3α-(hydroxymethyl)-6,8-dimethoxy-7-hydroxytetralin-2β-yl]methyl]β-D-xylopyranoside;(+) -Lyoniside.**[Intended Use]**

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

[Source]The stem barks of *Canarium bengalense*.

[Biological Activity or Inhibitors]

Lyoniside and saracoside are cytotoxic to promastigotes and intracellular amastigotes, they effectively kill *L. donovani* amastigotes inside macrophages in vitro, demonstrate strong anti-leishmanial efficacies in BALB/c mice model of leishmaniasis, suggests that these two compounds potential anti-leishmanial candidates. ^[1]

The synergistic action of lyoniside and triterpene acids was demonstrated in inhibitory effect exerted on germination and growth of *Pinus sylvestris*. ^[2]

[Solvent]

Pyridine, Methanol, Ethanol, etc.

[HPLC Method]^[3]

Mobile phase: 0.1% Trifluoroacetic acid (TFA) H₂O-Methanol, gradient elution ;

Flow rate: 1.0 ml/min;

Column temperature: Room Temperature;

The wave length of determination: 280 nm.

[Storage]

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

[References]

[1] Saha S, Mukherjee T, Chowdhury S, *et al. Biochem. Pharmacol.*, 2013, 86(12): 1673-87.

[2] Szakiel A, Voutquenne-Nazabadioko L, Henry M. *Phytochem. Lett.*, 2011, 4(2):138-43.

[3] Lee B, Jin B W, Yun B R, *et al. Pharm. Mag.*, 2014, 10(10):195-9.

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