

# **Calystegine B1 Datasheet**

4<sup>th</sup> Edition (Revised in July, 2016)

### [ Product Information ]

Name: Calystegine B1

Catalog No.: CFN00164

Cas No.: 127414-86-2

**Purity:** > 95%

**M.F:** C<sub>7</sub>H<sub>13</sub>NO<sub>4</sub>

M.W: 175.18

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 root bark of Lycium chinense Mill.

# [ Biological Activity or Inhibitors]

Calystegines occur in many plants of the Convolvulaceae, Solanaceae, and Moraceae

families, calystegines B1 and C1 are potent competitive inhibitors of the bovine, human,

and rat beta-glucosidase activities, with Ki values of 150, 10, and 1.9 microM, respectively

for B1 and 15, 1.5, and 1 microM, respectively, for C1;calystegine B2 is a strong

competitive inhibitor of the alpha-galactosidase activity in all the livers; calystegines A3

and B2 selectively inhibit the rat liver beta-glucosidase activity; the potent inhibition of

mammalian beta-glucosidase and alpha-galactosidase activities in vitro raises the

possibility of toxicity in humans consuming large amounts of plants that contain these

compounds.[1]

[Solvent]

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

[ HPLC Method ]<sup>[2]</sup>

Mobile phase: Acetonitrile- 0. 1% Acetic acid H2O=50:50;

Flow rate: 0.8 ml/min;

Column temperature: 30 °C;

The wave length of determination: 254 nm.

[Storage]

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

[References]

[1] Asano N, Kato A, Matsui K, et al. Glycobiology, 1998, 7(8):1085-8.

[2] Shang L, Ouyang Z, Zhao M, et al. Chinese Journal of Experimental Traditional

Medical Formulae, 2014, 20(2):47-51.

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