

(Z)-Butylidenephthalide Datasheet

4th Edition (Revised in July, 2016)

[Product Information]

Name: (Z)-Butylidenephthalide

Catalog No.: CFN96106

Cas No.: 72917-31-8

Purity: >95%

 $M.F: C_{12}H_{12}O_2$

M.W: 188.22

Physical Description: Oil

Synonyms: cis-Butylidenephthalide.

[Intended Use]

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

[Source]

The roots of Angelica sinensis.

[Biological Activity or Inhibitors]

3-(Z)-butylidenephthalide shows antihyperglycemic activity, it (56.2 mg/kg) decreases

blood glucose levels in NAD-STZ-diabetic mice after an oral sucrose load, suggesting that

its antihyperglycemic effect is due to inhibition of α-glucosidase at the intestinal level; it

also inhibit the activity of yeast-α-glucosidase (IC(50) 2.35 mM) in a noncompetitive

fashion with a K(i) of 4.86 mM .[1]

(Z)-Butylidenephthalide(Bdph) restore temozolomide sensitivity to can

temozolomide-resistant malignant glioma cells by downregulating expression of the DNA

repair enzyme MGMT.[2]

(Z)-Butylidenephthalide was reported to more potently inhibit electrically induced twitch

responses than acetylcholine-induced tonic contraction in isolated guinea-pig ileum

(GPI).[3]

(Z)-Butylidenephthalide has antitumor effects, because of the limitation of the blood-brain

barrier, the Bdph dosage required for treatment of glioma is relatively high; a local-release

system with Bdph incorporated into a biodegradable polyanhydride material, p(CPP-SA;

Bdph-Wafer) can deliver a sufficient concentration of Bdph to the tumor site and effectively

inhibit the tumor growth in the glioma.[4]

[Solvent]

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

[HPLC Method]^[5]

Mobile phase: Chloroform-n-Nexane=50:50;

Flow rate: 1.0 ml/min;

Column temperature: Room Temperature;

The wave length of determination: 254 nm.

[Storage]

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

[References]

- [1] Brindis F, Rodríguez R, Bye R,et al. J. Nat. Prod., 2011, 74(3):314-20.
- [2] Yen S Y, Harn H J, Lin S Z, et al. Eur. J. Cancer, 2012, 48(5):S220-1.
- [3] Chen M, Ko W C. Archiv Für Experimentelle Pathologie Und Pharmakologie, 2016, 389(2):159-66.
- [4] Harn H J, Lin S Z, Lin P C, et al. Neuro-Oncol., 2011, 13(6):635-48.
- [5] Ko W C, Yang J Y, Chen C M. J. Chromatogr. B, 1996, 685(2):379-82.

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