

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

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PRODUCT INFORMATION



BH3I-1

Item No. 18763

CAS Registry No.: 300817-68-9

Formal Name: 5-[(4-bromophenyl)methylene]-a-

(1-methylethyl)-4-oxo-2-thioxo-3-

thiazolidineacetic acid

MF: $C_{15}H_{14}BrNO_3S_2$

400.3 FW: ≥95% **Purity:**

Stability: ≥2 years at -20°C Supplied as: A crystalline solid λ_{max} : 279, 382 nm UV/Vis.:

Laboratory Procedures

For long term storage, we suggest that BH3I-1 be stored as supplied at -20°C. It should be stable for at least two years.

BH3I-1 is supplied as a crystalline solid. A stock solution may be made by dissolving the BH3I-1 in the solvent of choice. BH3I-1 is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide (DMF), which should be purged with an inert gas. The solubility of BH3I-1 in ethanol is approximately 5 mg/ml and approximately 30 mg/ml in DMSO and DMF.

BH3I-1 is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, BH3I-1 should first be dissolved in DMSO and then diluted with the aqueous buffer of choice. BH3I-1 has a solubility of approximately 0.5 mg/ml in a 1:1 solution of DMSO:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

Description

The family of Bcl-2 proteins plays pivotal roles in either promoting or preventing apoptosis. Bcl-2 family members contain one or more of four characteristic Bcl-2 homology (BH) domains, which are crucial for function. For example, anti-apoptotic Bcl-2 family proteins prevent death signaling by heterodimerizing with pro-death proteins at their BH3 domains. BH3I-1 is a cell permeable inhibitor that blocks the binding of BH3 peptides to Bcl-xL, inducing apoptosis.² It inhibits interactions of BH3 domain-containing proteins with Bcl-xL, Bcl-2, and Bcl-W, inducing apoptosis in Bcl-2 or Bcl-W expressing cells with K, values of 43.4 and 124 µM, respectively.^{3,4} BH3I-1 enhances radiation sensitivity in non-small cell lung cancer cells.³

References

- 1. Ni Chonghaile, T. and Letai, A. Mimicking the BH3 domain to kill cancer cells. Oncogene 27(Suppl 1), S149-S157 (2008).
- 2. Degterev, A., Lugovskoy, A., Cardone, M., et al. Identification of small-molecule inhibitors of interaction between the BH3 domain and Bcl-xL. Nat. Cell. Biol. 3(2), 173-182 (2001).
- 3. Roa, W., Chen, H., Alexander, A., et al. Enhancement of radiation sensitivity with BH3I-1 in non-small cell lung cancer. Clin. Invest. Med. 28(2), 55-63 (2005).
- 4. Porter, J.R., Helmers, M.R., Wang, P., et al. Profiling small molecule inhibitors against helix-receptor interactions: The Bcl-2 family inhibitor BH3I-1 potently inhibits p53/hDM2. Chem. Commun. 46(42), 8020-8022 (2010).

WARNING
THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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