

PRODUCT DATA SHEET

AG-CN2-0067

06-Oct-2012

10Z-Hymenialdisine

[Hymenialdisine; 4-(2-Amino-4-oxo-2-imidazolidin-5-ylidene)-2-bromo-4,5,6,7-tetrahydropyrrolo[2,3-c]azepin-8-one]

AG-CN2-0067-C500	
AG-CN2-0067-M001	

Formula MW CAS 500 μg 1 mg

C₁₁H₁₀BrN₅O₂ 324.1 82005-12-7

Handling / Storage

Shipping Short Term Storage Long Term Storage AMBIENT +4°C -20°C

Keep cool and dry. Keep under inert gas. Protect from light.

Use / Stability

Stable for at least 2 years after receipt when stored at -20°C.

MSDS available at www.adipogen.com or upon request.

Product Specifications

Source/Host	Isolated from sponge Axinella carteri.
Purity	≥97% (HPLC) (Contains traces of the 10E-isomer)
Identity	Determined by ¹ H-NMR.
Appearance	Yellow oil.
Solubility	Soluble in DMSO.

Product Description

- Insecticidal and cytotoxic [1].
- Potent NF-κB inhibitor. Inhibits various pro-inflammatory cytokines such as IL-1, IL-2, IL-6, IL-8, TNF-α and nitric oxide (NO) in a variety of cell lines [2, 3, 4, 9].
- ATP-competitive kinase inhibitor. Inhibits DNA damage checkpoint at G2, cyclin-dependent kinases CDK1/cyclin B, CDK2/cyclin A, CDK2/cyclin E, CDK4/cyclin D1, CDK5/p25, GSK-3β and casein kinase 1 (CK1) [5, 6].
- Potent mitogen-activated protein kinase kinase-1 (MEK-1) inhibitor [7].
- MARK (microtubule affinity-regulating kinase) inhibitor [8].

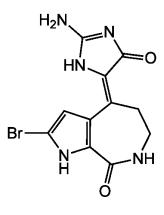
WARNING: Intended for research use only. This product is not intended or approved for human, diagnostics, therapeutic or veterinary use. Use of this product for human or animal testing is extremely hazardous and may result in disease, severe injury, or death. MATERIAL SAFETY DATA: Review the complete Material Safety Data Sheet before use.

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Product Specific References

- 1. Bioactive alkaloids from the tropical marine sponge Axinella carteri: A. Supriyono, et al.; Z. Naturforsch. [C] **50**, 669 (1995)
- 2. The natural product hymenialdisine inhibits interleukin-8 production in U937 cells by inhibition of nuclear factor-kappaB: J.J. Breton & M.C. Chabot-Fletcher; J. Pharmacol. Exp. Ther. **282**, 459 (1997)
- 3. Inhibition of NFkappaB-mediated interleukin-1beta-stimulated prostaglandin E2 formation by the marine natural product hymenialdisine: A. Roshak, et al.; J. Pharmacol. Exp. Ther. **283**, 955 (1997)
- Inhibition of interleukin-1-induced proteoglycan degradation and nitric oxide production in bovine articular cartilage/chondrocyte cultures by the natural product, hymenialdisine: A.M. Badger, et al.; J. Pharmacol. Exp. Ther. 290, 587 (1999)
- 5. Inhibition of cyclin-dependent kinases, GSK-3beta and CK1 by hymenialdisine, a marine sponge constituent: L. Meijer, et al.; Chem. Biol. **7**, 51 (2000)
- 6. Inhibition of the G2 DNA damage checkpoint and of protein kinases Chk1 and Chk2 by the marine sponge alkaloid debromohymenialdisine: D. Curman, et al.; J. Biol. Chem. **276**, 17914 (2001)
- 7. Aldisine alkaloids from the Philippine sponge Stylissa massa are potent inhibitors of mitogen-activated protein kinase kinase-1 (MEK-1): D. Tasdemir, et al.; J. Med. Chem. **45**, 529 (2002)
- 8. Protein kinase MARK/PAR-1 is required for neurite outgrowth and establishment of neuronal polarity: J. Biernat et al.; Mol. Biol. Cell. **13**, 4013 (2002)
- 9. Inhibition of cytokine production by hymenialdisine derivatives: V. Sharma, et al.; J. Med. Chem. **47,** 3700 (2004)

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