Gliotoxin

Proteasome inhibitor

Immunomodulating mycotoxin which acts by blocking membrane thiol groups. Inhibits the chymotrypsin-like activity of the 20S proteasome in a non-competitive manner. Causes apoptotic cell death in macrophages and thymocytes. Induces Ca²⁺ release from intact rat liver mitochondria. Inhibits the activation of transcription factor NF-κB in response to a variety of stimuli in T and B cells. Displays anti-inflammatory activity in vivo. Antioxidant. Inhibitor of farnesyltransferase and geranylgeranyltransferase I. A negative control analog is available (Prod. No. BML-L122). Displays anti-tumor activity against breast cancer *in vivo*.

Citations: 13

View Online »

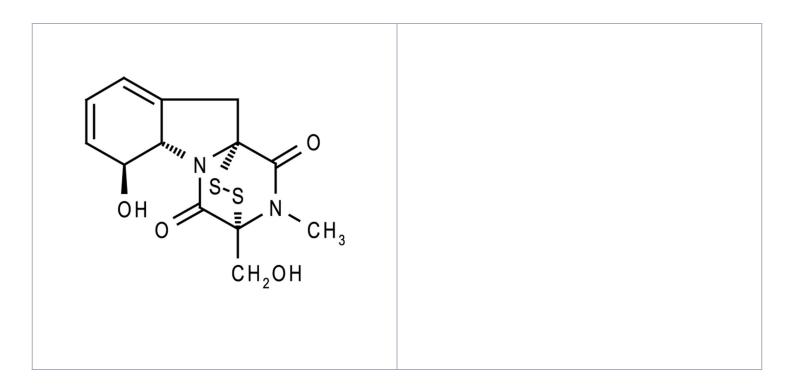
Ordering Information

Order Online »

BML-PI129-0002	2mg
BML-PI129-0010	10mg

Manuals, SDS & CofA

View Online »



Handling & Storage

Use/Stability As indicated on product label or CoA when stored as recommended. Stock solutions are

stable for up to 3 months when stored at -20°C.

Long Term Storage -20°C

Shipping Ambient Temperature

Regulatory Status RUO - Research Use Only

Product Details

Alternative Name $[3R-(3\alpha, 5a\beta,6\beta,10a\alpha)]-2,3,5a,6$ -Tetrahydro-6-hydroxy-3-

(hydroxymethyl)-2-methyl-10H-3,10a-epidithiopyrazino[1,2-a]indole-1,4-dione

Appearance White to yellow crystalline solid.

CAS 67-99-2

Couple Target Proteasome 20S

Couple Type Inhibitor

Formula $C_{13}H_{14}N_2O_4S_2$

MI 14: 4441

MW 326.4

Purity ≥97% (HPLC, TLC)

RTECS KB4725000

Solubility Soluble in DMSO or methanol (10mg/ml).

Source Isolated from Gliocladium fimbriatum



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