Apicidin

HDAC inhibitor

Potent inhibitor of histone deacetylase (HDAC). Inhibits proliferation. Induces cell cycle arrest at the G1 phase. At 100nM it induces a long lasting hyperacetylation of histone H4 while that induced by trichostatin is transient. Stimulates apoptosis. Apoptosis is induced via induction of Fas/Fas ligand. Displays potent antiangiogenic effects and dramatically decreases HIF-1 α protein levels and transcriptional activity in human and mouse tumor cell lines. Antiprotozoal.

Citations: 20

View Online »

Ordering Information

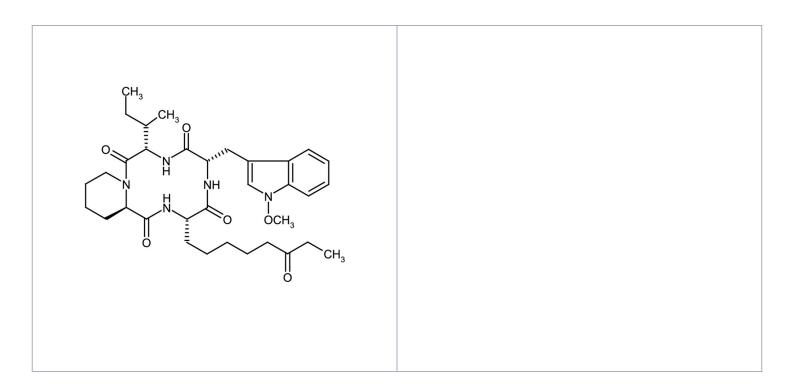
Order Online »

BML-GR340-0005

5mg

Manuals, SDS & CofA

View Online »



Handling & Storage

Use/Stability As indicated on product label or CoA when stored as recommended. Stable for at least

1 year after receipt when stored, as supplied, at -20°C. Stock solutions are stable for up

to 3 months at -20°C.

Long Term Storage -20°C

Shipping Blue Ice

Regulatory Status RUO - Research Use Only

Product Details

Alternative Name cyclo-L-(2-Amino-8-oxodecanoyl)-L-(N-methoxy-

 $tryptophan)\hbox{-$L$-isoleucyl-D-pipecolinyl, } Cyclo(N\hbox{-}O\hbox{-methyl-L-}$

tryptophanyl-L-isoleucinyl-D-pipecolinyl-L-2-amino-8-

oxodecanoyl)

Appearance White powder.

CAS 183506-66-3

Couple Target HDAC

Couple Type Inhibitor

Formula $C_{34}H_{49}N_5O_6$

MW 623.8

Purity ≥99% (HPLC)

Solubility Soluble in DMSO or 100% ethanol.

Last modified: May 29, 2024

