Withaferin A

NF-kB inhibitor

Cell permeable and potent angiogenesis inhibitor from the family of withanolides. Inhibits endothelial cells (HUVEC) sprouting *in vitro* (IC $_{50}$ =12nM) and *in vivo*. Exhibits antitumor, anti-inflammatory, radiosensitizing, and immunosuppressive properties. Potently inhibits NF-κB activation by preventing the TNF-induced activation of IκB kinase β (IKK β). Elicits IKK β hyperphosphorylation concomitant with inhibition of its kinase activity. Affects AP1 transcription and induces cell death. Alters cytoskeletal architecture by covalently binding annexin II and stimulating its basal F-actin cross-linking activity which inhibits the migratory and invasive capability of endothelial cells. Targets the intermediate filament protein vimentin. Displays neuronal regenerative properties. It induced significant regeneration of both axons and dendrites in an A β -induced axonal atrophy model.

Citations: 14

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Ordering Information

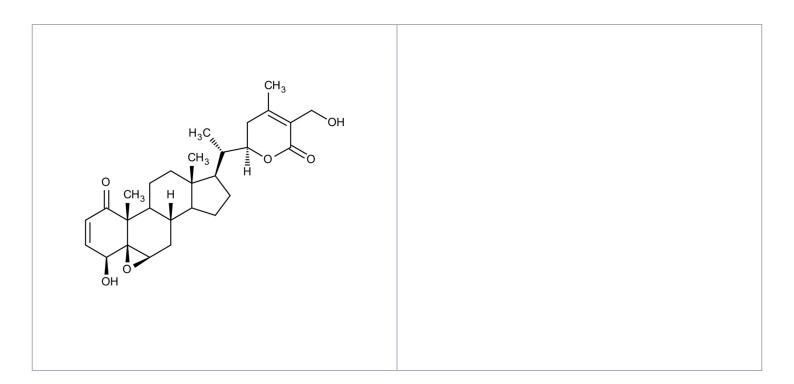
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BML-CT104-0010

10mg

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.

Handling Protect from light.

Long Term Storage -20°C

Shipping Ambient Temperature

Regulatory Status RUO - Research Use Only

Product Details

Alternative Name 5,6-Epoxy-4,27-dihydroxy-1-oxowitha-2,24-dienolide

Appearance Off-white solid.

CAS 5119-48-2

Couple Target NF-kappaB

Couple Type Inhibitor

Formula $C_{28}H_{38}O_6$

MI 14: 10048

MW 470.6

MeltingPoint 241-243°C

Purity ≥99% (HPLC)

RTECS KE7288500

Solubility Soluble in DMSO (20 mg/ml), methanol (10 mg/ml) or

100% ethanol.

Source Isolated from Withania somnifera.

Last modified: May 29, 2024



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