Geldanamycin

HSP90 inhibitor

Potent antitumor antibiotic. Inhibitor of pp60src tyrosine kinase and of cmyc gene expression in murine lymphoblastoma cells. Inhibits the transforming activity of abl, erbB, fps, src, and yes. Binds specifically to heat shock protein 90 (HSP90) and to its endoplasmic reticulum homolog GP96 (GRP94). Capable of destabilizing several oncogene and protooncogene products. Potent inhibitor of the nuclear hormone receptor family. Protects against α-synuclein toxicity to dopaminergic neurons in Drosophila. Destabilizes mutant p53 protein from a number of breast, leukemic, and prostate cell lines. Inhibits basal and hypoxia-induced expression of c-Jun (IC_{50} =75nM) and abolishes hypoxia-induced increase in c-Jun N-terminal kinase (JNK) activity. Inhibits telomerase activity through inhibition of HSP90, a chaperone required for the assembly and activation of telomerase in human cells. Hsp90 inhibitors geldanamycin or 17-DMAG reduces the uptake of Chaperone Mediated Autophagy (CMA) substrates by isolated lysosomes.~10-fold more potent than herbimycin A (Prod. No. BML-EI227)

Citations: 39

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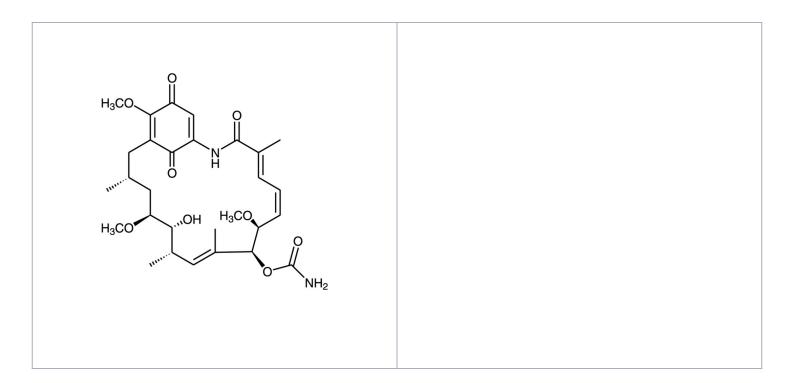
Ordering Information

Order Online »

BML-El280-0001	1mg
BML-EI280-0005	5mg

Manuals, SDS & CofA

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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

Appearance Yellow solid.

CAS 30562-34-6

Couple Target HSP90, Src kinase

Couple Type Inhibitor

Formula $C_{2q}H_{40}N_2O_q$

MW 560.6

Purity ≥95% (HPLC)

RTECS LX8920000

Solubility Soluble in DMSO (10mg/ml); insoluble in water.

Technical Info / Product Notes Replacement for ADI-HPK-102

Last modified: May 29, 2024

