# **Trichostatin A**

#### **HDAC** inhibitor

Trichostatin A (TSA) is a potent and reversible inhibitor of histone deacetylases. In HeLa cells, TSA blocked cell cycle progression at G1 and induced a 12-fold increase in intracellular levels of gelsolin. In cells latently infected with HIV-1, TSA induced the transcriptional activation of the HIV-1 promoter, which resulted in a marked increase in virus production. In NIH 3T3 cells, TSA induced reversion of oncogenic ras-transformed cells to a normal morphology. In Jurkat cells, TSA inhibited IL-2 gene expression (IC  $_{50}$ =73nM) and displayed immunosuppressive activity in a mouse model.

Induces increased acetylation of GATA4, a cardiac-specific transcription factor and increases cardiac muscle cell differentiation. Trichostatin A is a useful tool for induction of hyperacetylation of cellular histones and for further elucidation of their role in gene expression. Induces cell growth arrest at both G1 and G2/M phases. may induce apoptosis in some cases. It can induce accelerated dedifferentiation of primordial germ cells into embryonic germ cells.

Citations: 31

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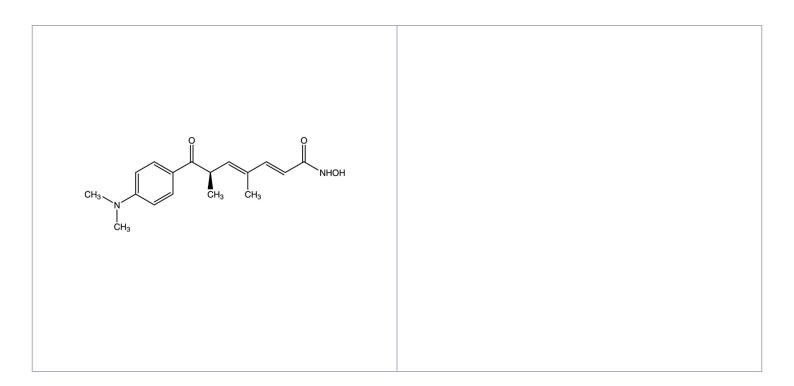
## **Ordering Information**

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BML-GR309-0001	1mg
BML-GR309-0005	5mg

Manuals, SDS & CofA

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### **Handling & Storage**

**Use/Stability** As indicated on product label or CoA when stored as recommended. Store, as supplied,

at -20°C for up to 1 year. Store solutions at -20°C for up to 3 months.

Long Term Storage -20°C

**Shipping** Ambient Temperature

### Regulatory Status RUO - Research Use Only

#### **Product Details**

Alternative Name [R-(E,E)]-7-[4-(Dimethylamino)phenyl]-N-hydroxy-4,6-

dimethyl-7-oxo-2,4-heptadienamide

**Appearance** Off-white to beige solid.

**CAS** 58880-19-6

Couple Target HDAC

Couple Type Inhibitor

Formula  $C_{17}H_{22}N_2O_3$ 

**MW** 302.4

MeltingPoint 150°C

Purity ≥98% (HPLC)

**Solubility** Soluble in DMSO, dimethyl formamide, acetonitrile,

methanol or 100% ethanol

Source Isolated from Streptomyces platensis.

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