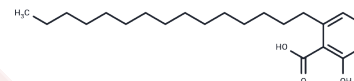


## Anacardic Acid

## Chemical Properties

CAS No. :	16611-84-0
Formula:	C <sub>22</sub> H <sub>36</sub> O <sub>3</sub>
Molecular Weight:	348.52
Storage:	Powder: -20°C for 3 years   In solvent: -80°C for 1 year Actual storage temperature shall be subject to the COA.



## Biological Description

Description	Anacardic Acid (6-pentadecylsalicylic Acid) is an effective inhibitor of p300 and p300/CBP-associated factor histone acetyltransferases. It also has antimicrobial activity, antibacterial activity, and inhibits prostaglandin synthase, lipoxygenase, and
Targets(IC50)	Epigenetic Reader Domain,Histone Acetyltransferase,Antibacterial
In vitro	Anacardic acid is an inhibitor of several Clinically targeted enzymes such as HATs, NF-κB, LOX-1, xanthine oxidase, tyrosinase, and ureases. Anacardic acid is also an effective inhibitor of Tip60 in vitro. It inhibits the Tip60-dependent acetylation and activation of the ATM protein kinase in HeLa cells and sensitizes the cells to the cytotoxic effects of radiation.
In vivo	Anacardic acid decreases over-expression of Gata4, α-MHC, and cTnT in fetal mouse hearts exposed to ethanol. Anacardic acid inhibits binding of PCAF, P300 to the promoter of Gata4.
Kinase Assay	HAT and HDAC Activity Assays :After homogenization of cardiac tissues, nucleoproteins are extracted using a Nuclear Extract Kit according to the manufacturer's instructions. HAT and HDAC activities of the nuclear protein extracts are determined using a colorimetric assay included in the HAT and HDAC assay kits.
Cell Research	HeLa and 293T cells are grown in Dulbecco's modified Eagle's medium/10% fetal calf serum. SQ20B and SCC35 squamous cell carcinoma cell lines are cultured in modified eagles medium/10% fetal calf serum. For anacardic acid (EMD Biosciences, CA) exposure, cells are preincubated with anacardic acid (0-100 μM) for 40 min, irradiated, then allowed to recover for 4 h. Cells are then switched to fresh media and allowed to grow for 10 days.
Animal Research	Animal Models: 9 to 11 week old Kunming mice. Formulation: Dissolved in DMSO,co-injected with ethanol. Dosages: 0,1.25,2.5,5,10 mg/kg. Administration: i.p.

## Solubility Information

Solubility	Ethanol: 17.4 mg/mL (49.93 mM),Sonication is recommended. DMSO: 252 mg/mL (723.06 mM),Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
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## A DRUG SCREENING EXPERT

In vivo Formulation	10% DMSO+90% Corn Oil: 3.3 mg/mL (9.47 mM),Sonication is recommended. <i>Please add the solvents sequentially, clarifying the solution as much as possible before adding the next one. Dissolve by heating and/or sonication if necessary. Working solution is recommended to be prepared and used immediately. The formulation provided above is for reference purposes only. In vivo formulations may vary and should be modified based on specific experimental conditions.</i>
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### Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	2.8693 mL	14.3464 mL	28.6928 mL
5 mM	0.5739 mL	2.8693 mL	5.7386 mL
10 mM	0.2869 mL	1.4346 mL	2.8693 mL
50 mM	0.0574 mL	0.2869 mL	0.5739 mL

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

### Reference

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