

Catalog Number: CM02933

## 产品信息

**Catalog Number:**  
CM02933

**CAS号:**  
22862-76-6

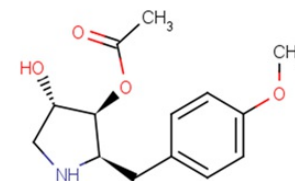
**分子式:**  
 $C_{14}H_{19}NO_4$

**主要靶点:**  
Antibacterial|Antibiotic|Apoptosis|DNA/RNA  
Synthesis|JNK

**主要通路:**  
微生物学|DNA损伤和修复|凋  
亡|MAPK信号通路|细胞周期

**分子量:**  
265.3

**溶解度:**  
DMSO:26.5 mg/mL (100  
mM), Ethanol:13.3 mg/mL (50 mM)



## 体外活性

Anisomycin ( $3 \mu M$ ) decreases protein synthesis in MDA16 and MDA-MB-468 cells, and reduces colony formation by MDA-MB-468 cells. Anisomycin causes an increase in the number of apoptotic cells in MDA-MB-468 cultures, but not in MDA16 cultures. Anisomycin activates JNK phosphorylation in MDA-MB-468 cells.[2] In U251 and U87 cells, anisomycin ( $0.01-8 \mu M$ ) inhibits the cell growth in time- and concentration-dependent manners with the IC<sub>50</sub> (48 h) values of 0.233 and 0.192  $\mu mol/L$ , respectively. Anisomycin ( $4 \mu M$ ) causes 21.5% and 25.3% of apoptosis proportion in U251 and U87 cells, respectively, and activates p38 MAPK and JNK, while inactivated ERK1/2. Anisomycin ( $4 \mu M$ ) reduces the level of PP2A/C subunit in a time-dependent manner in U251 and U87 cells.[3] Anisomycin inhibits EAC cell proliferation in concentration-dependent manner.[4]

## 体内活性

Peritumoral administration of anisomycin (5 mg/kg) significantly suppresses Ehrlich ascites carcinoma (EAC) growth resulting in the survival of approximately 60% of the mice 90 days after EAC inoculation.[4]

## 细胞实验

For the assay, EAC cells are plated in 96-well plates at a density of 10,000 cells/well/200  $\mu L$  of medium. The cells are treated with the different concentrations of anisomycin for 48 h. Adriamycin (500 ng/mL) is used as a positive control. 0.5 mg/mL of MTT is added to each well. 4 h later, the formazan product of MTT reduction is dissolved in DMSO, and absorbance is measured at 570 nm using a Model 680 microplate reader.(Only for Reference)

## 描述

Anisomycin is an antibiotic isolated from various Streptomyces species. It interferes with protein and DNA synthesis by inhibiting peptidyl transferase or the 80S ribosome system.

## 储存

Powder: -20°C for 3 years | In solvent: -80°C for 2 years