

Catalog Number: CM00554

## 产品信息

**Catalog Number:**  
CM00554

**CAS号:**  
16858-02-9

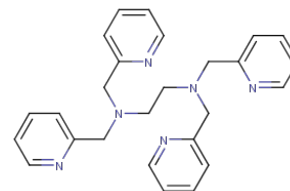
**分子式:**  
 $C_{26}H_{28}N_6$

**主要靶点:**  
Reactive Oxygen  
Species|Apoptosis|Autophagy

**主要通路:**  
凋亡|NF- $\kappa$ B 信号通路|代谢|免疫与  
炎症|自噬

**分子量:**  
424.54

**溶解度:**  
DMSO:6 mg/mL (14.13 mM)



## 体外活性

TPEN通过降低铜、汞和甲基汞引起的Fura-2荧光变化而表现出其效应。特别是在添加铜氯化物（10/30  $\mu$ M）激发的细胞中，TPEN在暴露后3小时加入可以显著降低提升的Fura-2荧光比率至基础水平，在10分钟内降低至119.6 $\pm$ 2.4%或109 $\pm$ 1.5%（ $\Delta$ 比率(F340/F380)的降低）。此外，TPEN通过铜的氧化还原循环针对结肠癌细胞，依赖于剂量和时间减少细胞活性。TPEN引发的细胞死亡也依赖于铜的氧化还原循环，因为铜螯合剂neocuproine抑制了DNA损伤，并降低了pChk1、 $\gamma$ -H2AX和ATM蛋白表达。

## 细胞实验

TPEN is dissolved in DMSO and then diluted with appropriate medium[1]. Human neuroblastoma cell line SH-SY5Y, are grown in Dulbecco's Modified Eagle's Medium (DMEM) mixed 1:1 with Ham's F-12 nutrient mixture containing 10% fetal bovine serum, 100 unit/mL penicillin and 100  $\mu$ g/mL streptomycin at 37°C in a humidified 5% CO<sub>2</sub> atmosphere. Two days before experimentation, cells are seeded at a density of 7 $\times$ 10<sup>4</sup> cells/cm<sup>2</sup> in a 96-well plate. Cells in a 96-well plate are serum-starved for 4 hr; calcium indicator fura-2 is then loaded into the cells by using Calcium kit II fura-2. In brief, SH-SY5Y cells are incubated with 5  $\mu$ M fura-2/AM in the presence of 0.04% Pluronic F-127, a dispersing agent to improve the efficiency of loading with fura-2, and 1.25 mM probenecid, a blocker of organic anion transport to prevent leakage of fura-2 from cells. After 1 hr incubation at 37°C, fura-2 fluorescence is measured at 500 nm emission after excitation at 340 nm (F340) or 380 nm (F380) using an Infinite M200 plate reader at 37°C. The change in [Ca<sup>2+</sup>]<sub>i</sub> is reflected by the ratio of F340 and F380. To determine the changes in fura-2 fluorescence ratio induced by heavy metal compounds, cells are treated with manganese chloride, lead acetate, cadmium chloride, mercuric chloride and MeHg chloride dissolved in distilled water. We confirmed that the cells adhered to the bottom of the plate after 6 hr exposure to heavy metal compounds. The cells are also treated with three Ca<sup>2+</sup> channel blockers, lanthanum chloride dissolved in distilled water, verapamil and 2-APB dissolved in DMSO, 30 min before heavy metal exposure. The heavy metal chelator TPEN is dissolved in DMSO and added 3 hr after the stimulation with heavy metals to determine the contribution of endogenous and exogenous heavy metals on fura-2 fluorescence changes. We measured the effect of TPEN (20  $\mu$ M) on the fura-2 fluorescence ratio after a 10 min treatment with TPEN, since our preliminary experiments showed that the effect of TPEN on fura-2 fluorescence reached maximum and stabilized within 10 min of the treatment[1].

## 储存

Powder: -20°C for 3 years | In solvent: -80°C for 1 year | Shipping with blue ice.