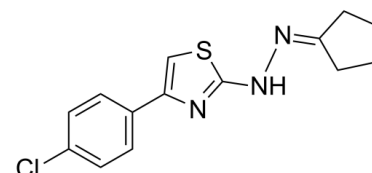


## Data Sheet

<b>Product Name:</b>	CPTH2
<b>Cat. No.:</b>	CS-W013990
<b>CAS No.:</b>	357649-93-5
<b>Molecular Formula:</b>	C <sub>14</sub> H <sub>14</sub> ClN <sub>3</sub> S
<b>Molecular Weight:</b>	291.80
<b>Target:</b>	Apoptosis; Histone Acetyltransferase
<b>Pathway:</b>	Apoptosis; Epigenetics
<b>Solubility:</b>	DMSO : 25 mg/mL (85.68 mM; Need ultrasonic); H <sub>2</sub> O : < 0.1 mg/mL (ultrasonic) (insoluble); Ethanol : 1 mg/mL (3.43 mM; Need ultrasonic)



### BIOLOGICAL ACTIVITY:

CPTH2 is a potent **histone acetyltransferase (HAT)** inhibitor. CPTH2 selectively inhibits the acetylation of histone H3 by **Gcn5**. CPTH2 induces **apoptosis** and decreases the invasiveness of a clear cell renal carcinoma (ccRCC) cell line through the inhibition of **acetyltransferase p300 (KAT3B)**<sup>[1][2]</sup>. *In Vitro*: CPTH2 (100 µM; 12, 24, 48 hours) causes a decrease in cell proliferation after as early as 12 h with a further significant reduction after 48 h stimulation<sup>[1]</sup>. CPTH2 (100 µM; 12 or 48 hours) causes a comparable drop of the activity in both cell lines<sup>[1]</sup>. CPTH2 (100 µM; 48 hours) produces a drastic increase in apoptotic/dead cell population after 48 h<sup>[1]</sup>. CPTH2 (100 µM; 12, 24, 48 hours) shows a reduced acetylation of both global AcH3 histone and H3AcK18<sup>[1]</sup>. CPTH2 (100 µM; 24, 48 hours) is capable to counteract invasion and migration of ccRCC-786-O cells in culture<sup>[1]</sup>. CPTH2 (0.2, 0.5, 1 mM) inhibits the growth of a GCN5 deleted strain and a single catalytic mutant E173H<sup>[2]</sup>. CPTH2 (0.6, 0.8 mM; for 24 hours) inhibits histone H3 acetylation in yeast cell cultures<sup>[2]</sup>. CPTH2 inhibits the Gcn5p dependent functional network<sup>[2]</sup>.

### References:

- [1]. Cocco E, et al. KAT3B-p300 and H3AcK18/H3AcK14 levels are prognostic markers for kidney ccRCC tumor aggressiveness and target of KAT inhibitor CPTH2. Clin Epigenetics. 2018 Apr 4;10:44.
- [2]. Chimenti F, et al. A novel histone acetyltransferase inhibitor modulating Gcn5 network: cyclopentylidene-[4-(4'-chlorophenyl)thiazol-2-yl]hydrazonone. J Med Chem. 2009 Jan 22;52(2):530-6.

### CAIndexNames:

Cyclopentanone, 2-[4-(4-chlorophenyl)-2-thiazolyl]hydrazonone

### SMILES:

C1C=CC=C(C2=CSC(N/N=C3CCCC3)=N2)C=C1

**Caution: Product has not been fully validated for medical applications. For research use only.**

Tel: 610-426-3128

Fax: 888-484-5008

E-mail: sales@ChemScene.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA