

**Product Name** : L-002

**Synonyms** : NSC 764414;L002

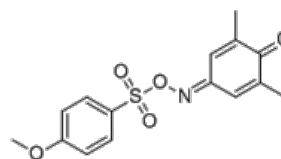
**Cat No.** : M14048

**CAS Number** : 321695-57-2

**Molecular Formula** : C<sub>15</sub>H<sub>15</sub>NO<sub>5</sub>S

**Formula Weight** : 321.35

**Chemical Name** : 2,6-dimethyl-2,5-cyclohexadiene-1,4-dione 4-[O-[(4-methoxyphenyl)sulfonyl]oxime]



**Description** : L-002 (NSC 764414, L002) is a novel potent, specific acetyltransferase p300 (KAT3B) inhibitor with IC<sub>50</sub> of 1.98 μM; also shows weak inhibitory activity against PCAF (KAT2B) and GCN5 (KAT2A) with IC<sub>50</sub> of 34.7 and 33.9 μM, displays no inhibition against Tip60, MYST2 and MYST4 (IC<sub>50</sub>>100 μM), as well as a panel of HDACs and HMTs; inhibits acetylation of histones and p53, and suppresses STAT3 activation in cell-based assays; exhibits extreme sensitivity against leukemia, breas and lymphoma cell lines; potently suppresses tumor growth and histone acetylation of MDA-MB-468 xenografts.

**Pathway** : Chromatin/Epigenetic

**Target** : HAT

**Receptor** : HAT

**Solubility** : —

**SMILES** : CC1=CC(=NOS(=O)(=O)C2=CC=C(C=C2)OC)C=C(C1=O)C

**Storage** : (-20°C)

**Stability** : ≥ 2 years

**Reference** :

1. Rai R, et al. Epigenetics. 2017;12(11):1004-1013. 2. Yang H, et al. Mol Cancer Ther. 2013 May;12(5):610-20.