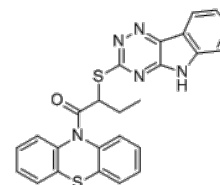


**Product Name** : Inauhzin  
**Cat. No.** : PC-38473  
**CAS No.** : 309271-94-1  
**Molecular Formula** : C<sub>25</sub>H<sub>19</sub>N<sub>5</sub>OS<sub>2</sub>  
**Molecular Weight** : 469.581  
**Target** : MDM2-p53  
**Solubility** : 10 mM in DMSO



## Biological Activity

Inauhzin (INZ) is a small molecule **SIRT1** inhibitor that effectively reactivates p53, promotes p53-dependent apoptosis of human cancer cells without causing apparently genotoxic stress.

Inauhzin (INZ) induced p53 levels as effectively as actinomycin D (ActD, 10 nM), did not apparently affect the interaction between either MDMX and p53, or MDM2 and MDMX, or MDM2 and p53.

Inauhzin inhibited cell growth in a p53-dependent fashion against human lung cancer H460, A549, H1299, colon cancer HCT116, HT29 (IC<sub>50</sub>=5-50 uM), induced p53 level and activity as well as p53-dependent apoptosis.

Inauhzin (INZ) stabilizes p53 without either directly inhibiting MDM2-mediated ubiquitylation or causing genotoxicity.

Inauhzin (INZ) effectively inhibits SIRT1 deacetylase activity at 3 uM, directly binds to SIRT1 in vitro, induces p53 acetylation at lysine 382 at 2 uM in cells.

Inauhzin (INZ) induces p53 and p53-dependent apoptosis in vivo and suppresses the growth of human xenograft tumours.

## References

Zhang Q, et al. *EMBO Mol Med*. 2012 Apr;4(4):298-312.

**Caution: Product has not been fully validated for medical applications. Lab Use Only!**

E-mail: tech@probechem.com