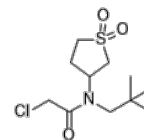


**Product Name** : Sulfopin  
**Cat. No.** : PC-22076  
**CAS No.** : 2451481-08-4  
**Molecular Formula** : C<sub>11</sub>H<sub>20</sub>ClNO<sub>3</sub>S  
**Molecular Weight** : 281.80  
**Target** : Other Targets  
**Solubility** : 10 mM in DMSO



CAS: 2451481-08-4

## Biological Activity

Sulfopin is a potent, selective and covalent inhibitor of Pin1 with  $K_i$  of 17 nM in FP assay, targets Pin1's active site Cys113, blocks Myc-driven tumors.

Sulfopin inhibits the catalytic activity of Pin1 with an apparent  $K_i$  of 211 nM in chymotrypsin-coupled peptidyl-prolyl isomerization assay.

Sulfopin is highly selective for Pin1 Cys113 over other 161 cysteine residues.

Sulfopin could phenocopy Pin1 knockout.

Sulfopin significantly resensitized HeLa cells to irradiation in a dose-dependent manner, and decreases IRAK1 Thr209 phosphorylation at concentrations as low as 100 nM.

Sulfopin (1  $\mu$ M) shows a Pin1-dependent viability effect following long-term exposure.

Sulfopin downregulates Myc transcriptional activity and increases c-Myc protein levels in Mino B cells, blocks neuroblastoma in zebrafish.

Sulfopin (20 or 40 mg/kg) inhibits pancreatic cancer progression in mice, significantly extends survival.

## References

Dubiella C, et al. Nat Chem Biol. 2021 Sep;17(9):954-963.

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

E-mail: tech@probechem.com