

## **Data Sheet**

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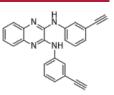
Global Supplier of Chemical Probes, Inhibitors & Agonists.

**Product Name** : CIP2A inhibitor TD52

Cat. No. : PC-23570
CAS No. : 1798328-24-1
Molecular Formula : C<sub>24</sub>H<sub>16</sub>N<sub>4</sub>
Molecular Weight : 360.42

Target : Protein Phosphatase/PTP

**Solubility** : 10 mM in DMSO



## **Biological Activity**

TD52 is a small molecule inhibitor of cancerous inhibitor of protein phosphatase 2A (CIP2A), downregulates CIP2A and p-Akt, reactivates protein phosphatase 2A.

TD52 causes greater reduction in cell viability than erlotinib in all the HCC cell lines tested including HA22T (IC50=0.9  $\mu$ mol/l), Hep3B (IC50=0.9  $\mu$ mol/l), PLC5 (IC50=0.8  $\mu$ mol/l) and Sk-Hep1 (IC50=1.2  $\mu$ mol/l), which is independent of EGFR kinase inhibition.

TD52 downregulates the protein expression of CIP2A and p-Akt in a dose- and time-dependent manner, with no effect on the expression level of PP2A-related subunits, PP2A-A, PP2A-B55 and PP2A-C.

TD52 downregulates transcription of CIP2A via interfering Elk-1 function.

TD52 (10 mg/kg/day) exhibits in vivo antitumor effects on a PLC5 xenograft tumor model.

## References

Yu HC, et al. Cell Death Dis. 2014 Jul 31;5(7):e1359.

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

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