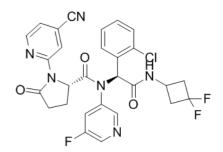


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## **Data Sheet**

Global Supplier of Chemical Probes, Inhibitors & Agonists.

| Product Name      | : | Ivosidenib   |
|-------------------|---|--|
| Cat. No.          | : | PC-45550   |
| CAS No.           | : | 1448347-49-6   |
| Molecular Formula | : | C <sub>28</sub> H <sub>22</sub> ClF <sub>3</sub> N <sub>6</sub> O <sub>3</sub> |
| Molecular Weight  | : | 582.9609   |
| Target            | : | Isocitrate Dehydrogenase (IDH)   |
| Solubility        | : | DMSO: ≥ 39 mg/mL   |
|                   |   |  |



## **Biological Activity**

Ivosidenib (AG-120) is a potent, mutant-selective inhibitor of **IDH1** (isocitrate dehydrogenase type 1) with IC50 of 12 nM for mouse IDH1 R132H.

Ivosidenib (AG-120) shows similar potency against IDH1-R132H (IC50=12 nM); IDH1-R132C (IC50=13 nM); IDH1-R132G (IC50=8 nM); IDH1-R132L (IC50=13 nM); IDH1-R132S (IC50=12 nM), respectively.

Ivosidenib (AG-120) (50 mg/kg and 150 mg/kg, orally gavage) suppresses tumor 2-HG concentration rapidly, with maximum inhibition (92.0% and 95.2% at the 50 mg/kg and 150 mg/kg doses, respectively).

Ivosidenib (AG-120) represent a worthwhile avenue of exploration in the treatment of IDH1-mutated tumors.

## References

Cuyàs E, et al. **Oncotarget.** 2015 May 20;6(14):12279-96.

Cancer Discov. 2015 Jan;5(1):4.

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

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