## **Product** Data Sheet

## CH-0793076

Cat. No.: HY-107096 CAS No.: 534605-78-2 Molecular Formula:  $C_{26}H_{26}N_4O_4$ Molecular Weight: 458.51

Target: Topoisomerase

Pathway: Cell Cycle/DNA Damage

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.

## **BIOLOGICAL ACTIVITY**

Description	CH-0793076 (TP3076), a hexacyclic camptothecin analog, is active drug and major metabolite of TP300. CH-0793076 inhibits DNA topoisomerase I with an IC $_{50}$ of 2.3 $\mu$ M. CH-0793076 is efficacious against cells expressing BCRP (breast cancer resistance protein) $^{[1]}$ .
IC <sub>50</sub> & Target	Topoisomerase I 2.3 μM (IC <sub>50</sub> )
In Vitro	CH0793076 (TP3076) (6 days at 37°C) shows antiproliferative activity against PC-6/BCRP and PC-6/pRC cells, with IC $_{50}$ s of 0.35 and 0.18 nM, respectively <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	TP300 (1-100 mg/kg; bolus intravenous injection once per week for 3 weeks, for a total of three doses), a water-soluble prodrug of CH0793076, shows more than 50% of tumor growth inhibition in all nine models, regardless of the expression of BCRP (WiDr, HT-29, NCI-H460 and ASPC-1, HCT116, COLO 201, HCT-15, Calu-6 and NCI-N87) <sup>[1]</sup> .  MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## **REFERENCES**

[1]. Endo M, et al. A water soluble prodrug of a novel camptothecin analog is efficacious against breast cancer resistance protein-expressing tumor xenografts. Cancer Chemother Pharmacol. 2010 Jan;65(2):363-71.

Caution: Product has not been fully validated for medical applications. For research use only.

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