## **Product** Data Sheet

## ALK/ROS1-IN-1

Cat. No.: HY-130794 CAS No.: 2365497-07-8 Molecular Formula:  $C_{30}H_{35}F_3N_6O_3$ 

Molecular Weight: 584.63

Target: ALK; ROS

Pathway: Protein Tyrosine Kinase/RTK

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

Analysis.

## **BIOLOGICAL ACTIVITY**

Description	ALK/ROS1-IN-1 (compound 2e) is a potent and selective anti crizotinib-resistant ALK/ROS1 dual inhibitor, with IC $_{50}$ s of 0.174 $\mu$ M and 0.530 $\mu$ M for ALK and ROS1 enzyme, respectively.
IC <sub>50</sub> & Target	IC50: 0.174 $\mu$ M (ALK), 0.530 $\mu$ M (ROS1) $^{[1]}$
In Vitro	ALK/ROS1-IN-1 displays potent anti-proliferative activity against ALK-addicted H3122 and ROS1-addicted HCC78 cell lines $(IC_{50}=6.27~\mu\text{M}~\text{and}~10.71~\mu\text{M},\text{respectively})^{[1]}.$ ALK/ROS1-IN-1 shows impressive enzyme activity against clinically Crizotinib-resistant ALK <sup>L1196M</sup> with an IC <sub>50</sub> value of 41.3 nM <sup>[1]</sup> . ALK/ROS1-IN-1 shows potent inhibitory activity in Ba/F3 cell line expressing ROS1 mutants, with IC <sub>50</sub> s of 137.7, 104.7 nM and 233.9 for wide-type, G2032R mutant and L2026M mutant, respectively <sup>[1]</sup> . ALK/ROS1-IN-1 has no significant effect on inducing apoptosis of HCC78 cell line <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## **REFERENCES**

[1]. Liu S, et al. Design, synthesis and biological evaluations of 2-amino-4-(1-piperidine) pyridine derivatives as novel anti crizotinib-resistant ALK/ROS1 dual inhibitors. Eur J Med Chem. 2019 Oct 1;179:358-375.

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

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