## RET-IN-13

Cat. No.: HY-144029 CAS No.: 2684258-54-4 Molecular Formula:  $C_{32}H_{33}F_4N_5O_3$ 

Molecular Weight: 611.63 Target: RET

Pathway: Protein Tyrosine Kinase/RTK

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

Analysis.

## **BIOLOGICAL ACTIVITY**

Description	RET-IN-13 (compound 1), a quinoline compound, is a potent RET inhibitor with IC <sub>50</sub> s of 0.5 nM, 0.9 nM for RET (WT) and RET (V804M), respectively. RET-IN-13 has the potential for tumors or intestinal diseases related to abnormal activation of RET research <sup>[1]</sup> .
In Vitro	RET-IN-13 has low inhibition on CYP1A2 (IC $_{50}$ >50 $\mu$ M), CYP2C9 (IC $_{50}$ =12.5 $\mu$ M), CYP2C19 (IC $_{50}$ =16.9 $\mu$ M), CYP2D6 (IC $_{50}$ =18.8 $\mu$ M) and CYP3A4 (IC $_{50}$ >50 $\mu$ M)[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	RET-IN-13 (compound 1; 0.5 mpk; iv) has a $T_{1/2}$ of 6.93 hours, a CL of 2.68 mL/min·kg, and a $Vd_{ss}$ of 1.52 L/kg for male mice <sup>[1]</sup> . RET-IN-13 (2.5 mpk; po) has a $T_{1/2}$ of 5.16 hours, $C_{max}$ of 0.958 $\mu$ M and an AUC of 12477 $\mu$ M·hr for male mice <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## **REFERENCES**

[1]. Jiansong Wang, et al. Quinoline compounds. WO2021164742A1.

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

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