

## **Product** Data Sheet

Inhibitors

**Screening Libraries** 

**Proteins** 

FGFR4-IN-6

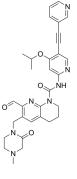
Cat. No.:HY-143881CAS No.:2760970 - 10 - 1Molecular Formula: $C_{31}H_{33}N_7O_4$ Molecular Weight:567.64

Target: FGFR

Pathway: Protein Tyrosine Kinase/RTK

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

Analysis.



**BIOLOGICAL ACTIVITY** 

**Description** FGFR4-IN-6 (Compound 9ka) is a covalently reversible FGFR4 inhibitor with an IC<sub>50</sub> value of 5.4 nM. FGFR4-IN-6 also exhibits good oral pharmacokinetic properties. FGFR4-IN-6 induces significant tumor regressions in a xenograft mouse model of

good oral pharmacokinetic properties. FGFR4-IN-6 induces significant tumor regressions in a xenograft mouse model of Hep3B2.1-7 HCC cell line without an obvious sign of toxicity<sup>[1]</sup>. FGFR4-IN-6 is a click chemistry reagent, it contains an Alkyne group and can undergo copper-catalyzed azide-alkyne cycloaddition (CuAAc) with molecules containing Azide groups.

**REFERENCES** 

[1]. Zhen Zhang, et al. Design, Synthesis, and Biological Evaluation of 2-Formyl Tetrahydronaphthyridine Urea Derivatives as New Selective Covalently Reversible FGFR4 Inhibitors. J Med Chem. 2022 Feb 24;65(4):3249-3265.

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

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