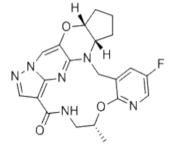


## **Data Sheet**

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Global Supplier of Chemical Probes, Inhibitors & Agonists

 $\begin{tabular}{lll} \textbf{Product Name} & :TPX-0046 \\ \textbf{Cat.No.} & :URK-V871 \\ \textbf{CAS No.} & :2359650-19-2 \\ \textbf{Molecular Formula} & :C_{21}H_{21}FN_6O_3 \\ \textbf{Molecular Weight} & :424.436 \\ \textbf{Target} & :c-RET \\ \textbf{Solubility} & : \\ \end{tabular}$ 



## **Biological Activity**

TPX-0046 (TPX0046) is a potent, selective next-generation RET/SRC inhibitor, demonstrates low nanomolar potency against WT and 18 RET mutations/fusions, as well as SRC, and is VEGFR2/KDR-sparing.

TPX-0046 inhibited RET phosphorylation (IC50 < 10 nM) in tumor cell lines (LC2/ad, CCDC6-RET; TT, RET C634W) and Ba/F3 engineered RET models (WT, G810R).

TPX-0046 inhibited KIF5B-RET Ba/F3, LC2/ad, and TT cells with IC50 values of 1 nM in cell proliferation assays.

TPX-0046 notently inhibited Ba/F3 RET engineered cells with SEMs (e.g. G810C/R/S) with mean proliferation IC5

TPX-0046 potently inhibited Ba/F3 RET engineered cells with SFMs (e.g. G810C/R/S) with mean proliferation IC50 of 1-17 nM.

TPX-0046 demonstrated marked in vivo anti-tumor efficacy in RET-driven cell-derived and patient-derived xenograft tumor models, a single dose of 5 mg/kg

TPX-0046 inhibited >80% of RET phosphorylation.

TPX-0046 is a unique next-generation RET inhibitor that possesses potent in vitro and in vivo activity against a diverse range of RET alterations, including SFM-mediated resistance.

## References

1. A.Drilon, et al. Annals of Oncology Volume 30, Supplement 5, October 2019, Pages v190-v191

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