

## **Data Sheet**

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 $\begin{tabular}{lll} \textbf{Product Name} & :G007\text{-LK} \\ \textbf{Cat.No.} & :URK\text{-V}1005 \\ \textbf{CAS No.} & :1380672\text{-}07\text{-}0 \\ \textbf{Molecular Formula} & :C_{25}H_{16}\text{CIN}_7O_3S \\ \end{tabular}$ 

Molecular Weight :529.958 Target :PARP

Solubility :

## **Biological Activity**

G007-LK is a potent, specific, metabolically stable TNKS1/2 inhibitor ectively; reduces Wnt/ $\beta$ -catenin signaling by preventing poly(ADP-ribosyl)ation-dependent AXIN degradation, thereby promoting  $\beta$ -catenin destabilization; effectively blocks ligand-driven Wnt/ $\beta$ -catenin signaling in cell culture and displays approximately 50% inhibition of APC mutation-driven signaling in most CRC cell lines, inhibits the g of mouse intestinal organoids cultured ex vivo with IC50 of 80 nM; inhibits in vivo tumor growth in a subset of APC-mutant CRC xenograft models.

## References

- 1. Voronkov A, et al. J Med Chem. 2013 Apr 11;56(7):3012-23.
- 2. Lau T, et al. Cancer Res. 2013 May 15;73(10):3132-44.

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