

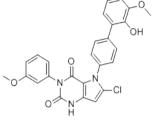
Data Sheet

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 $\begin{tabular}{lll} \textbf{Product Name} & :GSK621 \\ \textbf{Cat.No.} & :URK-V624 \\ \textbf{CAS No.} & :1346607-05-3 \\ \textbf{Molecular Formula} & :C_{26}H_{20}CIN_3O_5 \\ \textbf{Molecular Weight} & :489.912 \\ \textbf{Target} & :AMPK \\ \end{tabular}$

Solubility :



Biological Activity

GSK621 (GSK-621) is a potent, specific AMPK agonist (activator), induces cytotoxicity in AML (IC50=13-30 uM) but not in normal hematopoietic cells.

GSK621 is more potent than A-769662 at inducing AMPK activation, as measured by the level of ACC phosphorylation, markedly increases phosphorylation at AMPKa T172 in AML cell lines (MOLM-14, HL-60, and OCI-AML3) and primary AML samples.

Cytotoxicity in AML cells from GSK621 involves the eIF2a/ATF4 signaling pathway that specifically results from mTORC1 activation.

References

- 1. Sujobert P, et al. Cell Rep. 2015 Jun 9;11(9):1446-57.
- 2. Jiang H, et al. PLoS One. 2016 Aug 17;11(8):e0161017.
- 3. Chen L, et al. Biochem Biophys Res Commun. 2016 Nov 25;480(4):515-521.
- 4. Liu W, et al. Oncotarget. 2017 Feb 7;8(6):10543-10552.

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