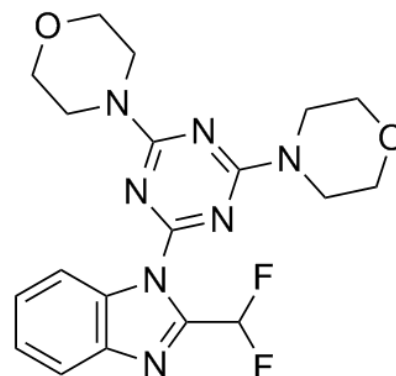


**Product Name** : ZSTK474  
**Cat. No.** : PC-45869  
**CAS No.** : 475110-96-4  
**Molecular Formula** : C<sub>19</sub>H<sub>21</sub>F<sub>2</sub>N<sub>7</sub>O<sub>2</sub>  
**Molecular Weight** : 417.4125  
**Target** : PI3K  
**Solubility** : 10 mM in DMSO



## Biological Activity

ZSTK474 (ZSTK-474) is a potent inhibitor of **class I PI3K isoforms** with IC<sub>50</sub> of 17 nM, 53 nM, and 6 nM for p110β, p110γ, and p110δ, respectively.

ZSTK474 (ZSTK474) shows potent antiproliferative activity against a panel of 39 human cancer cell lines with mean GI<sub>50</sub> of 0.32 μM.

ZSTK474 (ZSTK474) is more effectively than that of LY294002 or wortmannin.

ZSTK474 (ZSTK474) induces apoptosis in OVCAR3 cells, and induces complete G1-phase arrest but not apoptosis in A549 cells at 10 μM.

ZSTK474 (ZSTK474) completely inhibits the growth of A549, PC-3, and WiDr xenografts in mice at 400mg/kg; and induces the regression of A549 xenograft tumors.

## References

Yaguchi S, et al. *J Natl Cancer Inst*, 2006, 98(8), 545-556.

Kong D, et al. *Cancer Sci*, 2007, 98(10), 1638-1642.

Kong D, et al. *Eur J Cancer*, 2009, 45(5), 857-865.

**Caution: Product has not been fully validated for medical applications. Lab Use Only!**

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