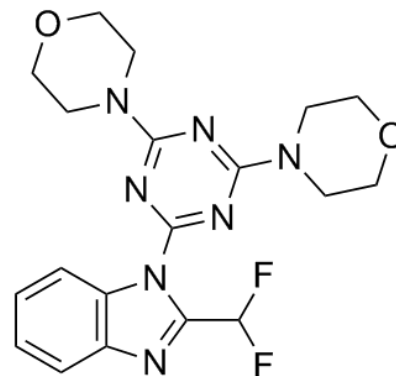


Product Name : ZSTK-474
Cat. No. : PC-45869
CAS No. : 475110-96-4
Molecular Formula : C₁₉H₂₁F₂N₇O₂
Molecular Weight : 417.4125
Target : PI3K
Solubility : 10 mM in DMSO



Biological Activity

ZSTK-474 (ZSTK474) is a potent inhibitor of **class I PI3K isoforms** with IC₅₀ of 17 nM, 53 nM, and 6 nM for p110β, p110γ, and p110δ, respectively.

ZSTK-474 (ZSTK474) shows potent antiproliferative activity against a panel of 39 human cancer cell lines with mean GI₅₀ of 0.32 μM.

ZSTK-474 (ZSTK474) is more effectively than that of LY294002 or wortmannin.

ZSTK-474 (ZSTK474) induces apoptosis in OVCAR3 cells, and induces complete G₁-phase arrest but not apoptosis in A549 cells at 10 μM.

ZSTK-474 (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

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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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