

Z-VAD-FMK

Cat. No. 2163

Z-Val-Ala-Asp(OMe)-FMK

Chemical Name: Benzyloxycarbonyl-Val-Ala-Asp(OMe)-
fluoromethylketone

Biological Activity

Potent, selective, cell-permeable and irreversible inhibitor of phosphatidylinositol 3-kinase (PI 3-kinase) ($IC_{50} = 2 - 4$ nM). Also potently inhibits polo-like kinase 1 (PLK1) ($IC_{50} = 5.8$ nM).

Technical Data

M.Wt:

428.44

Formula:

$C_{23}H_{24}O_8$

Solubility:

Soluble to 50 mM in DMSO and to 5 mM in ethanol

Storage:

Desiccate at $-20^{\circ}C$

CAS No:

19545-26-7

The technical data provided above is for guidance only.

For batch specific data refer to the Certificate of Analysis.

References

Arcaro and Wymann (1993) Wortmannin is a potent phosphatidylinositol 3-kinase inhibitor: the role of phosphatidylinositol 3,4,5-trisphosphate in neutrophil responses. *Biochem.J.* **296** 297. PMID: [8257416](#).

Powis et al (1994) Wortmannin, a potent and selective inhibitor of phosphatidylinositol-3-kinase. *Cancer Res.* **54**2419. PMID: [8162590](#).

Schultz et al (1995) *In vitro* and *in vivo* antitumor activity of the phosphatidylinositol-3-kinase inhibitor wortmannin. *Anticancer Res.* **15** 1135. PMID: [7653991](#).

Liu et al (2005) Wortmannin, a widely used phosphoinositide 3-kinase inhibitor, also potently inhibits mammalian polo-like kinase. *Chem.Biol.* **12** 99. PMID: [15664519](#).

If you know of a relevant citation for this product [please let us know](#).

Keywords: Wortmannin, supplier, Potent, inhibitors, PLK1, PI, 3-kinase, irreversible, Polo-like, Kinases, PI3K, Mitosis, Phosphoinositide, SL2052

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