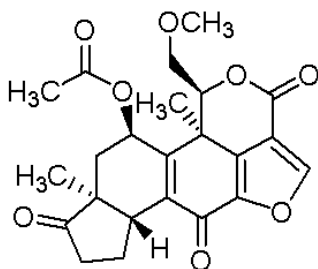


## Product Information

### Wortmannin Ready Made Solution from *Penicillium fumiculosum*

Catalog Number **W3144**  
Storage Temperature  $-20\text{ }^{\circ}\text{C}$

CAS RN 19545-26-7  
Synonyms: Wartmannin, Antibiotic SL-2052



#### Product Description

Molecular formula:  $\text{C}_{23}\text{H}_{24}\text{O}_8$   
Formula weight: 428.43

Wortmannin is a low molecular weight; hydrophobic fungal metabolite with a sterol-like structure produced by *Penicillium fumiculosum*.<sup>1</sup> Inhibition of the PI3K/Akt signal transduction cascade by Wortmannin enhances the apoptotic effects of radiation or serum withdrawal, and blocks the antiapoptotic effect of cytokines.<sup>2,3</sup> PI3K inhibition by Wortmannin also blocks many of the short-term metabolic effects induced by insulin receptor activation.<sup>4</sup> Research has demonstrated Wortmannin inhibits two enzymes from the mitotical division key regulators Polo-like kinase (Plk) family, Plk1 and Plk3.<sup>5,6</sup>

The product is supplied as a 10 mM 0.2  $\mu\text{m}$  filtered solution in dimethyl sulfoxide (DMSO).

Purity: >95% (HPLC)

#### Precautions and Disclaimer

This product is for R&D use only, not for drug, household, or other uses. Please consult the Material Safety Data Sheet for information regarding hazards and safe handling practices.

#### Storage/Stability

Store the product sealed at  $-20\text{ }^{\circ}\text{C}$ , protected from light. Under these conditions, the product is stable for at least 2 years.

#### References

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2. Arcaro, A., and Wymann, M.P., Wortmannin, a widely used phosphoinositide 3-kinase inhibitor, also potently inhibits mammalian polo-like kinase. *Biochem. J.*, **296**, 297–301(1993).
3. Yano, H., et al., Polo-like kinases inhibited by Wortmannin. Labeling site and downstream effects. *J. Biol. Chem.*, **268**, 25846–25856 (1993).
4. Moule, S.K., and Denton, R.M., Multiple signaling pathways involved in the metabolic effects of insulin. *Am. J. Cardiol.*, **80**, 41A-49A (1997).
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6. Liu, Y., Deficiency in chromosome congression by the inhibition of PLK1 polo box domain-dependent recognition. *J. Biol. Chem.*, **282**, 2505-2511 (2007).

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