

Technical Data Sheet: Z-VAD-FMK

Catalog Number SML18A or SML18B

Z-Val-Ala-Asp(OMe)-FMK; Z-VAD(OMe)-FMK; Synonyms

Size

Description Z-VAD-FMK is a cell-permeable pan-caspase inhibitor that irreversibly binds to the

catalytic site of caspase proteases and can inhibit induction of apoptosis. Z-VAD-FMK has been shown to block TGF-beta1-induced apoptosis and inhibit CPP32-like protease activity

in lysates in the pretreatment of hepatocytes³.

467.49 Molecular Weight

Molecular Formula C22H30FN3O7

Chemical Name L-Alaninamide, N-[(phenylmethoxy)carbonyl]-L-valyl-N-[(1S)-3-fluoro-1-(2-methoxy-2-

oxoethyl)-2-oxopropyl]-

CAS Number 187389-52-2

Caspase **Target**

Appearance White to off-white (Solid))

Purity ≥98% by HPLC

Solubility and Reconstitution Soluble in DMSO up to 213.91mM, for example:)

> 10 mg/106.954 mL = 0.093 mg/mL = 0.2 mM10 mg/21.3908 mL = 0.467 mg/mL = 1 mM10 mg/4.2782 mL = 2.337 mg/mL = 5 mM10 mg/2.1391 mL = 4.674 mg/mL = 10 mM

Storage Temperature and Stability Powder: -20°C 3 years

> 4°C 2 years

-80°C In solvent: 6 months

-20°C 1 month

References

- 1. Kawasaki M, et al. (2000) Protection from lethal apoptosis in lipopolysaccharide-induced acute lung injury in mice by a caspase inhibitor. Am J Pathol. 157(2):597-603.
- 2. Park S, et al. (2004) Neurovascular protection reduces early brain injury after subarachnoid hemorrhage. Stroke. 35(10):2412-7.
- Inayat-Hussain SH, et al. (1997) Processing/activation of CPP32-like proteases is involved in transforming growth factor beta1-induced apoptosis in rat hepatocytes. Hepatology. 25(6):1516-26.