Product Data Sheet

Ac-VLPE-FMK

Storage:

Cat. No.: HY-153614 CAS No.: 2679825-27-3 Molecular Formula: C₂₅H₄₁FN₄O₇ Molecular Weight: 528.61

Target: Cathepsin

Pathway: Metabolic Enzyme/Protease

Powder

-20°C 3 years 2 years

In solvent -80°C 6 months

> -20°C 1 month

BIOLOGICAL ACTIVITY

Description	Ac-VLPE-FMK, a tetrapeptidyl mono-fluoromethyl ketone (m-FMK), is a Cat-B and Cat-L inhibitor. Ac-VLPE-FMK can be used for the research of cancer aggressiveness $^{[1][2]}$.
IC ₅₀ & Target	Cathepsin B cathepsin L
In Vitro	Ac-VLPE-FMK (2 µM) significantly affects the cleavage of probe Ac-PLVQ-AMC by human recombinant CtsB and L ^[1] . Ac-VLPE-FMK (30 min) inhibits the generation of the fluorescence derived from the cleavage of the substrate Ac-PLVQ-AMC in 769-p and A498 cells ^[1] . Ac-VLPE-FMK (2.5-250 µM; 24-72 h) does not affect renal cancer cell viability, but influences cell migration rate, cellular adhesion, colony formation, and markers expression in renal cancer cells ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Rudzińska M, et, al. Cysteine Cathepsins Inhibition Affects Their Expression and Human Renal Cancer Cell Phenotype. Cancers (Basel). 2020 May 21;12(5):1310.

[2]. Citarella A, et, al. Peptidyl Fluoromethyl Ketones and Their Applications in Medicinal Chemistry. Molecules. 2020 Sep 3;25(17):4031.

Caution: Product has not been fully validated for medical applications. For research use only.

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA