Product Data Sheet

Derrone

Cat. No.: HY-N3737 **CAS No.:** 76166-59-1

Molecular Formula: $C_{20}H_{16}O_{5}$ Molecular Weight: 336.34

Target: Aurora Kinase; PERK; Reactive Oxygen Species

Pathway: Cell Cycle/DNA Damage; Epigenetics; Immunology/Inflammation; Metabolic

Enzyme/Protease; NF-кВ

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.

BIOLOGICAL ACTIVITY

Description	Derrone, a prenylated isoflavones, is an Aurora kinase inhibitor, with IC ₅₀ values of 6 and 22.3 μ M against Aurora B and Aurora A, respectively. Derrone shows anti-tumor activity ^{[1][2]} .
IC ₅₀ & Target	IC $_{50}$: 6 μM (Aurora kinase B), 22.3 μM (Aurora kinase A) $^{[1]}$
In Vitro	Derrone (30-60 µM, 15 days) significantly inhibits the formation and growth of MCF7 tumor spheroids, with the tumor spheroid growth inhibition (% TGI) of 17.5% and 65.4% for 30 and 60 µM Derrone, respectively ^[1] . Derrone shows the autophagic features, such as the conversion of LC3-I to LC3-II, the formation of autophagosome and the downregulation of SQSTM1/p62 (p62) ^[2] . Derrone induces autophagic cell death through intracellular ROS and sustained ERK phosphorylation in A549 cells ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Hoang NT, et al. In Vitro Characterization of Derrone as an Aurora Kinase Inhibitor. Biol Pharm Bull. 2016 Jun 1;39(6):935-45.

[2]. Kang MJ, et al. Derrone induces autophagic cell death through induction of ROS and ERK in A549 cells. PLoS One. 2019 Jun 19;14(6):e0218659.

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