# **Screening Libraries**

# **Tubulin polymerization-IN-39**

Cat. No.: HY-151982 Molecular Formula:  $C_{21}H_{21}N_5O_5$ **Molecular Weight:** 423.42

Microtubule/Tubulin Target:

Cell Cycle/DNA Damage; Cytoskeleton Pathway:

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

Analysis.

**Product** Data Sheet

# **BIOLOGICAL ACTIVITY**

Description

Tubulin polymerization-IN-39 is a tubulin polymerization inhibitor (IC<sub>50</sub>: 4.9 µM). Tubulin polymerization-IN-39 occupies the colchicine-binding site. Tubulin polymerization-IN-39 inhibits cancer cell proliferation<sup>[1]</sup>.

In Vitro

Tubulin polymerization-IN-39 (compound 4k) shows antiproliferative activity toward HeLa, HCT116, A549, and T47D, with the IC<sub>50</sub> values of 0.31, 1.28, 3.99 and 10.32  $\mu$ M, respectively<sup>[1]</sup>.

Tubulin polymerization-IN-39 (0.15-0.6  $\mu$ M, 24 h) increases percentages of HeLa cells at the G2/M phase [1]. Tubulin polymerization-IN-39 (0.15-0.6 μM, 24 h) induces HeLa cells apoptosis by increasing cleaved PARP<sup>[1]</sup>.

Tubulin polymerization-IN-39 (0.2-0.8 µM, 24 h) inhibits the migration and the tube formation of HUVEC<sup>[1]</sup>.

Tubulin polymerization-IN-39 (10  $\mu$ M) inhibits tubulin polymerization by 78%<sup>[1]</sup>.

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Cell Viability Assay<sup>[1]</sup>

Cell Line:	HeLa cells
Concentration:	0.15, 0.3, 0.6 μM
Incubation Time:	24 h
Result:	Increased cyclin B1 protein level and cleaved PARP.  Decreased p-cdc2 (the inactive form of cdc2).
Cell Migration Assay [1]	

Cell Line:	HUVEC			
Concentration:	0.2, 0.4, 0.8 μΜ			
Incubation Time:	24 h			
Result:	Dose-dependently inhibited HUVEC migration.  Prevented HUVEC motility and migration (wound healing assay).			

## **REFERENCES**

1]. Chen L, et al. Rational design, synthesis and biological evaluation of novel 2-(substituted amino)-[1,2,4]triazolo[1,5-a]pyrimidines as novel tubulin polymerization nhibitors. Eur J Med Chem. 2022 Dec 15;244:114864.						
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