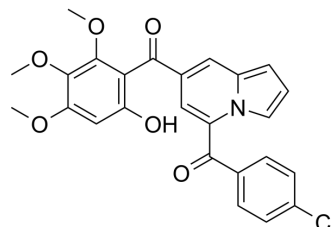


Antitumor agent-72

| | |
|---------------------------|---|
| Cat. No.: | HY-150696 |
| CAS No.: | 2676942-92-8 |
| Molecular Formula: | C ₂₅ H ₂₀ ClNO ₆ |
| Molecular Weight: | 465.88 |
| Target: | Apoptosis; Caspase; PARP |
| Pathway: | Apoptosis; Cell Cycle/DNA Damage; Epigenetics |
| Storage: | Please store the product under the recommended conditions in the Certificate of Analysis. |



BIOLOGICAL ACTIVITY

| | | | | | | | | | | | | | | | | | |
|--------------------|---|------------|---------------------------|----------------|-----------------------------------|------------------|----------|---------|---|------------|-------------|----------------|-----------------------------|------------------|----------|---------|--|
| Description | Antitumor agent-72 (compound 6w) is a potent anticancer agent. Antitumor agent-72 has anticancer activity and induces apoptosis through activation of caspase-3 and cleavage of PARP. Antitumor agent-72 can be used for cancer research ^[1] . | | | | | | | | | | | | | | | | |
| In Vitro | <p>Antitumor agent-72 (compound 6w) (0-10 μM; 72 h; BxPC3, PC3 and MCF7 cells) has anticancer activity and inhibits cell viability in a dose-dependent manner^[1].</p> <p>Antitumor agent-72 (compound 6w) (0-10 μM; 24 h; BxPC3 cells) induces apoptosis through activation of caspase-3 and cleavage of PARP^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Viability Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>BxPC3, PC3 and MCF7 cells</td> </tr> <tr> <td>Concentration:</td> <td>0, 0.03, 0.1, 0.3, 1, 3 and 10 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>72 hours</td> </tr> <tr> <td>Result:</td> <td>Inhibited cell viability in BxPC3, PC3 and MCF7 cells with IC₅₀ values of 0.47 μM, 2.68 μM and 1.82 μM, respectively.</td> </tr> </table> <p>Western Blot Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>BxPC3 cells</td> </tr> <tr> <td>Concentration:</td> <td>0, 0.1, 0.3, 1, 3 and 10 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 hours</td> </tr> <tr> <td>Result:</td> <td>Increased caspase-3 activation and PARP cleavage in BxPC3 cells.</td> </tr> </table> | Cell Line: | BxPC3, PC3 and MCF7 cells | Concentration: | 0, 0.03, 0.1, 0.3, 1, 3 and 10 μM | Incubation Time: | 72 hours | Result: | Inhibited cell viability in BxPC3, PC3 and MCF7 cells with IC ₅₀ values of 0.47 μM, 2.68 μM and 1.82 μM, respectively. | Cell Line: | BxPC3 cells | Concentration: | 0, 0.1, 0.3, 1, 3 and 10 μM | Incubation Time: | 24 hours | Result: | Increased caspase-3 activation and PARP cleavage in BxPC3 cells. |
| Cell Line: | BxPC3, PC3 and MCF7 cells | | | | | | | | | | | | | | | | |
| Concentration: | 0, 0.03, 0.1, 0.3, 1, 3 and 10 μM | | | | | | | | | | | | | | | | |
| Incubation Time: | 72 hours | | | | | | | | | | | | | | | | |
| Result: | Inhibited cell viability in BxPC3, PC3 and MCF7 cells with IC ₅₀ values of 0.47 μM, 2.68 μM and 1.82 μM, respectively. | | | | | | | | | | | | | | | | |
| Cell Line: | BxPC3 cells | | | | | | | | | | | | | | | | |
| Concentration: | 0, 0.1, 0.3, 1, 3 and 10 μM | | | | | | | | | | | | | | | | |
| Incubation Time: | 24 hours | | | | | | | | | | | | | | | | |
| Result: | Increased caspase-3 activation and PARP cleavage in BxPC3 cells. | | | | | | | | | | | | | | | | |

REFERENCES

[1]. Lee Y, et, al. Generation of a poly-functionalized indolizine scaffold and its anticancer activity in pancreatic cancer cells. Bioorg Chem. 2022 Sep;126:105877.

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

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