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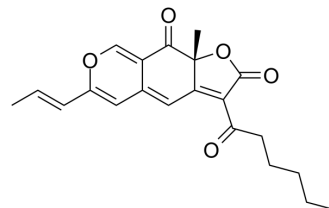
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Rubropunctatin

| | |
|--------------------|--|
| Cat. No.: | HY-N7766 |
| CAS No.: | 514-67-0 |
| Molecular Formula: | C ₂₁ H ₂₂ O ₅ |
| Molecular Weight: | 354.4 |
| Target: | Apoptosis |
| Pathway: | Apoptosis |
| Storage: | -20°C, protect from light * In solvent : -80°C, 6 months; -20°C, 1 month (protect from light) |



BIOLOGICAL ACTIVITY

| | | | | | | | | | | | | | | | | | |
|--------------------|---|------------|------------------------|----------------|------------------------------|------------------|----------|---------|--|------------|---------------|----------------|-----------------|------------------|--------------------|---------|---|
| Description | Rubropunctatin, an orange azaphilone pigment, is isolated from the extracts of <i>Monascus pilosus</i> -fermented rice (red-mold rice). Rubropunctatin has anti-inflammatory, immunosuppressive and antioxidative effects, and also exhibits anti-tumor activity ^{[1][2][3]} . | | | | | | | | | | | | | | | | |
| In Vitro | <p>Rubropunctatin (1.5-30 μM; 24 h) shows selective tumoricidal effect on the human gastric carcinoma BGC-823 cells and no significant toxicity toward normal epithelial cell GES-1^[1].</p> <p>Rubropunctatin (5-30 μM; 6-24 h) induces apoptosis in a dose- and time-dependent manner in BGC-823 cells^[1].</p> <p>Rubropunctatin (0.75-8.0 μg/ml) exhibits DDPH radical scavenging activity, inhibition of super oxide radical generation and ferric reducing activity^[2].</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>BGC-823 and GES-1cells</td> </tr> <tr> <td>Concentration:</td> <td>1.5, 3, 6, 12, 15, 18, 30 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 hours</td> </tr> <tr> <td>Result:</td> <td>Decreased the viability of BGC-823 cells with an IC₅₀ of 12.57 μM for 24 h. Did not show obvious cytotoxic effects on the normal cells.</td> </tr> </table> <p>Cell Cycle Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>BGC-823 cells</td> </tr> <tr> <td>Concentration:</td> <td>0, 5, 10, 30 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>0, 6, 12, 24 hours</td> </tr> <tr> <td>Result:</td> <td>Increased the percentage of cells in sub-G1 phase in a dose- and time-dependent manner.</td> </tr> </table> | Cell Line: | BGC-823 and GES-1cells | Concentration: | 1.5, 3, 6, 12, 15, 18, 30 μM | Incubation Time: | 24 hours | Result: | Decreased the viability of BGC-823 cells with an IC ₅₀ of 12.57 μM for 24 h. Did not show obvious cytotoxic effects on the normal cells. | Cell Line: | BGC-823 cells | Concentration: | 0, 5, 10, 30 μM | Incubation Time: | 0, 6, 12, 24 hours | Result: | Increased the percentage of cells in sub-G1 phase in a dose- and time-dependent manner. |
| Cell Line: | BGC-823 and GES-1cells | | | | | | | | | | | | | | | | |
| Concentration: | 1.5, 3, 6, 12, 15, 18, 30 μM | | | | | | | | | | | | | | | | |
| Incubation Time: | 24 hours | | | | | | | | | | | | | | | | |
| Result: | Decreased the viability of BGC-823 cells with an IC ₅₀ of 12.57 μM for 24 h. Did not show obvious cytotoxic effects on the normal cells. | | | | | | | | | | | | | | | | |
| Cell Line: | BGC-823 cells | | | | | | | | | | | | | | | | |
| Concentration: | 0, 5, 10, 30 μM | | | | | | | | | | | | | | | | |
| Incubation Time: | 0, 6, 12, 24 hours | | | | | | | | | | | | | | | | |
| Result: | Increased the percentage of cells in sub-G1 phase in a dose- and time-dependent manner. | | | | | | | | | | | | | | | | |
| In Vivo | <p>Rubropunctatin (8-32 mg/kg; i.v. for 5 times) has anti-tumor effect mice^[1].</p> <p>Rubropunctamine inhibits 12-O-tetradecanoylphorbol-13-acetate (TPA)-induced inflammation in mice, with an ID₅₀ of 0.11 mg/ear^[3].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> | | | | | | | | | | | | | | | | |

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|-----------------|---|
| Animal Model: | Male nude mice (5 weeks) are inoculated with BGC-823 cells ^[1] |
| Dosage: | 8, 32 mg/kg |
| Administration: | I.v. five times (day 1st, 4th, 7th, 10th, and 13th) |
| Result: | Diminished the tumor volume by 11.1% (8 mg/kg) and 24.2% (32 mg/kg). Reduced the tumor weight by 23.5% (8 mg/kg) and 37.7% (32 mg/kg). No significant difference was observed on the body weight. |

REFERENCES

- [1]. Zheng Y, et, al. Anti-cancer effect of rubropunctatin against human gastric carcinoma cells BGC-823. *Appl Microbiol Biotechnol*. 2010 Nov;88(5):1169-77.
- [2]. Dhale MA, et, al. Protective and antioxidative effect of rubropunctatin against oxidative protein damage induced by metal catalyzed reaction. *Int J Biol Macromol*. 2018 Sep;116:409-416.
- [3]. Akihisa T, et, al. Azaphilones, furanoisophthalides, and amino acids from the extracts of *Monascus pilosus*-fermented rice (red-mold rice) and their chemopreventive effects. *J Agric Food Chem*. 2005 Feb 9;53(3):562-5.
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Caution: Product has not been fully validated for medical applications. For research use only.

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