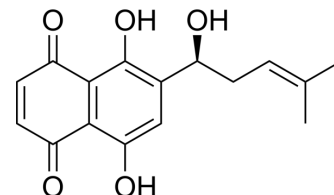


## Alkannin

Cat. No.:	HY-119874
CAS No.:	23444-65-7
Molecular Formula:	C <sub>16</sub> H <sub>16</sub> O <sub>5</sub>
Molecular Weight:	288.3
Target:	Pyruvate Kinase
Pathway:	Metabolic Enzyme/Protease
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Alkannin is a potent and specific inhibitor of tumor-specific pyruvate kinase-M2 (PKM2). Alkannin does not inhibit PKM1 and pyruvate kinase-L (PKL). Alkannin acts as a potential anticancer agent <sup>[1]</sup> .								
<b>In Vitro</b>	<p>Alkannin is a potent and specific inhibitor to PKM2, an enzyme that dictates the last rate-limiting step of glycolysis, which is essential for cancer cells' proliferation and survival. In the absence of D-fructose-1,6-bisphosphate (FBP), the IC<sub>50</sub> of Alkannin is 0.3 μM. In the presence of FBP (125 μM), the IC<sub>50</sub> of Alkannin is 0.9 μM, respectively. Alkannin effectively inhibits the cellular glycolytic flux in cancer cells dominantly expressing PKM2<sup>[1]</sup>.</p> <p>Alkannin (2.5-20 μM, 1 hour) inhibit the rates of cellular lactate production and glucose consumption<sup>[1]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Western Blot Analysis<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>MCF-7 and A549 express PKM2 but not PKM1 and PKL</td> </tr> <tr> <td>Concentration:</td> <td>0-20 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>1 hour</td> </tr> <tr> <td>Result:</td> <td>Inhibited the cellular glycolytic rate in a concentration-dependent manner.</td> </tr> </table>	Cell Line:	MCF-7 and A549 express PKM2 but not PKM1 and PKL	Concentration:	0-20 μM	Incubation Time:	1 hour	Result:	Inhibited the cellular glycolytic rate in a concentration-dependent manner.
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Concentration:	0-20 μM								
Incubation Time:	1 hour								
Result:	Inhibited the cellular glycolytic rate in a concentration-dependent manner.								

### REFERENCES

[1]. J Chen, et al. Shikonin and its analogs inhibit cancer cell glycolysis by targeting tumor pyruvate kinase-M2. *Oncogene*. 2011 Oct 20;30(42):4297-306.

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

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