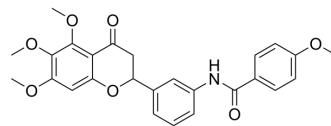


## Wnt/ $\beta$ -catenin-IN-1

<b>Cat. No.:</b>	HY-157990
<b>Molecular Formula:</b>	C <sub>26</sub> H <sub>25</sub> NO <sub>7</sub>
<b>Molecular Weight:</b>	463.48
<b>Target:</b>	Wnt; $\beta$ -catenin
<b>Pathway:</b>	Stem Cell/Wnt
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Wnt/ $\beta$ -catenin-in-1 (compounds 17) is a Wnt/ $\beta$ -catenin signaling pathway inhibitor. Wnt/ $\beta$ -catenin-IN-1 can induce apoptosis of colon cancer cells, has broad-spectrum anticancer activity, and can be used for the research of a variety of solid tumors <sup>[1]</sup> .																
<b>In Vitro</b>	<p>Wnt/<math>\beta</math>-catenin-IN-1 (compounds 17) (10 <math>\mu</math>M, 48 h) has broad-spectrum anticancer activity, with an average growth inhibition rate of 81.9% against 52 tumor cell lines in the trial<sup>[1]</sup>.</p> <p>Wnt/<math>\beta</math>-catenin-IN-1 (compounds 17) (10 <math>\mu</math>M, 48 h) is less cytotoxic to normal human cell lines (CCD841, MRC5) and is somewhat selective to cancer cells<sup>[1]</sup>.</p> <p>Wnt/<math>\beta</math>-catenin-IN-1 (compounds 17) (0/0.5/1/2 <math>\mu</math>M, 24 h/48 h) dose-dependently triggers cell cycle arrest and increases the population of HCT116 colon cancer cells in G2/M cell cycle phase<sup>[1]</sup>.</p> <p>Wnt/<math>\beta</math>-catenin-IN-1 (compounds 17) (0/0.5/1/2 <math>\mu</math>M, 24 h) is dose-dependent in reducing CDK8 level in HCT colorectal cancer cells<sup>[1]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Viability Assay<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>HCT116, HCT15, HT29, KM12, SW620, KM12, COLO205, HCC2998, CCD841, MRC5, et al.</td> </tr> <tr> <td>Concentration:</td> <td>0.05 <math>\mu</math>M</td> </tr> <tr> <td>Incubation Time:</td> <td>48 h</td> </tr> <tr> <td>Result:</td> <td>Showed broad-spectrum anticancer activity, less toxicity to normal cells, and certain selectivity to cancer cells.</td> </tr> </table> <p>Cell Cycle Analysis<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>HCT116, HCT15, HT29, KM12, SW620, KM12, COLO205, HCC2998, CCD841, MRC5, et al.</td> </tr> <tr> <td>Concentration:</td> <td>0.05 <math>\mu</math>M</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h</td> </tr> <tr> <td>Result:</td> <td>Triggered G2/M cell cycle arrest by activating the p53-p21 pathway.</td> </tr> </table>	Cell Line:	HCT116, HCT15, HT29, KM12, SW620, KM12, COLO205, HCC2998, CCD841, MRC5, et al.	Concentration:	0.05 $\mu$ M	Incubation Time:	48 h	Result:	Showed broad-spectrum anticancer activity, less toxicity to normal cells, and certain selectivity to cancer cells.	Cell Line:	HCT116, HCT15, HT29, KM12, SW620, KM12, COLO205, HCC2998, CCD841, MRC5, et al.	Concentration:	0.05 $\mu$ M	Incubation Time:	24 h	Result:	Triggered G2/M cell cycle arrest by activating the p53-p21 pathway.
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## REFERENCES

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[1]. Hassan AHE, et al. Discovery of a stilbenoid-flavanone hybrid as an antitumor Wnt/ $\beta$ -catenin signaling pathway inhibitor. *Bioorg Chem.* 2024 Feb 5;145:107178.

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

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