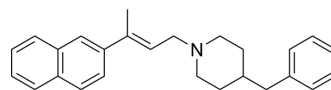


## RC-106

|                           |   |
|---------------------------|---|
| <b>Cat. No.:</b>          | HY-158197   |
| <b>CAS No.:</b>           | 1346216-50-9  |
| <b>Molecular Formula:</b> | C <sub>26</sub> H <sub>29</sub> N   |
| <b>Molecular Weight:</b>  | 355.52  |
| <b>Target:</b>            | Sigma Receptor; Proteasome  |
| <b>Pathway:</b>           | Neuronal Signaling; Metabolic Enzyme/Protease   |
| <b>Storage:</b>           | Please store the product under the recommended conditions in the Certificate of Analysis. |



### BIOLOGICAL ACTIVITY

|                    |  |   |
|--------------------|--|---|
| <b>Description</b> | RC-106 is a proteasome inhibitor (IC <sub>50</sub> : 35 μM) and Sigma receptor modulator with anticancer activity. RC-106 has antiproliferative activity against cancer cells including glioblastoma (GB) and multiple myeloma (MM) <sup>[1]</sup> . |   |
| <b>In Vitro</b>    | RC-106 (60 μM; 24 h) inhibits cancer cell activity <sup>[1]</sup> .<br>MCE has not independently confirmed the accuracy of these methods. They are for reference only.<br>Cell Viability Assay <sup>[1]</sup>  |   |
|                    | Cell Line:   | U87-MG cells  |
|                    | Concentration:   | 40, 60 μM   |
|                    | Incubation Time:   | 24 h  |
|                    | Result:  | Showed a moderate activity against U87-MG cells at 60 μM, reducing the cell viability to 34% <sup>⊠</sup> |

### REFERENCES

[1]. Listro R, et al. Exploring the RC-106 Chemical Space: Design and Synthesis of Novel (E)-1-(3-Arylbut-2-en-1-yl)-4-(Substituted) Piperazine Derivatives as Potential Anticancer Agents. *Front Chem.* 2020 Jun 30;8:495.

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

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