



## **Ansamitocin P-3**

Catalog No: tcsc1568

| Available Sizes   |
|---|
| Size: 5mg   |
| Size: 10mg  |
| Size: 50mg  |
| Size: 100mg   |
| Specifications  |
| <b>CAS No:</b> 66584-72-3   |
| <b>Formula:</b> C <sub>32</sub> H <sub>43</sub> CIN <sub>2</sub> O <sub>9</sub> |
| Pathway: Cell Cycle/DNA Damage;Cytoskeleton;Antibody-drug Conjugate/ADC Related |
| Target: Microtubule/Tubulin; Microtubule/Tubulin; ADC Cytotoxin                 |
| Purity / Grade: >98%  |
| Solubility:<br>10 mM in DMSO  |
| Alternative Names: Antibiotic C 15003P3;Maytansinol butyrate;C15003P3           |
| Observed Molecular Weight: 635.14   |





## **Product Description**

Ansamitocin P-3 is a **microtubule** inhibitor. Ansamitocin P-3 is a macrocyclic antitumor antibiotic.

IC50 & Target: Microtubule<sup>[1]</sup>

In Vitro: Ansamitocin P3 potently inhibits the proliferation of MCF-7, HeLa, EMT-6/AR1 and MDA-MB-231 cells in culture with a half-maximal inhibitory concentration of  $20\pm3$ ,  $50\pm0.5$ ,  $140\pm17$ , and  $150\pm1.1$  pM, respectively. Further, Ansamitocin P3 is found to bind to purified tubulin in vitro with a dissociation constant ( $K_d$ ) of  $1.3\pm0.7$  µM. The binding of Ansamitocin P3 induces conformational changes in tubulin. Ansamitocin P3 inhibits the proliferation of MCF-7, HeLa, EMT-6/AR1 and MDA-MB-231 cells in culture in a concentration dependent manner. Flow cytometric analysis of PI-stained cells suggests that Ansamitocin P3 inhibits the cell cycle progression of MCF-7 cells in G2/M phase. For example, 26, 50 and 70% of the cells are found to be in G2/M phase in the absence and presence of 50 and 100 pM Ansamitocin P3, respectively<sup>[2]</sup>.

All products are for RESEARCH USE ONLY. Not for diagnostic & therapeutic purposes!