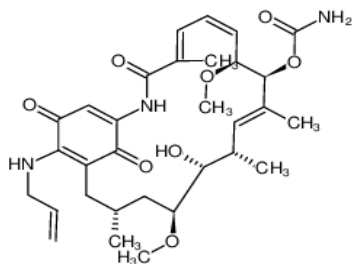


**17-AAG (CAS: 75747-14-7)****Catalog #: EBC51010****Biological Activity**

<b>Synonyms</b>	Tanespimycin
<b>Chemical Name</b>	17-Demethoxy-17-(2-propenylamino)-geldanamycin
<b>Application</b>	17-AAG is a potent heat shock protein 90 (Hsp90) inhibitor for BT474 tumor cells, derived from geldanamycin
<b>CAS No.</b>	75747-14-7
<b>Purity</b>	≥99.0%
<b>Molecular Weight</b>	585.69
<b>Molecular Formula</b>	C <sub>31</sub> H <sub>43</sub> N <sub>3</sub> O <sub>8</sub>
<b>Shipping</b>	Gel Pack
<b>Storage</b>	Store at -20° C
<b>Target &amp; IC<sub>50</sub></b>	Hsp90: = 5 μM

**Molecular Structure****Solubility**

DMSO: 58.0 mg/mL (100 mM)

**PS:** < 1 mg/ml refers to the product insoluble

**Description**

17-AAG is a less toxic and more stable analog of geldanamycin. 17-AAG is an Hsp90 inhibitor that displays a 100-fold higher affinity for Hsp90 derived from tumor cells compared to Hsp90 from normal cells. 17-AAG inhibits Akt activation and expression in tumors and synergizes with a number of anti tumor agents such as taxol2, cisplatin3, and UCN-014. 17-AAG causes the inactivation, destabilization and eventual degradation of HIF-1α. In the prostate, 17-AAG can inhibit the growth of cancer cell lines.

**For Research Use Only. Not For Use In Diagnostic Procedures**

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