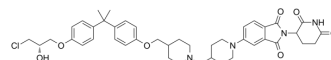


## BWA-522

<b>Cat. No.:</b>	HY-149433		
<b>Molecular Formula:</b>	C <sub>43</sub> H <sub>51</sub> ClN <sub>4</sub> O <sub>7</sub>		
<b>Molecular Weight:</b>	771.34		
<b>Target:</b>	Androgen Receptor; Apoptosis		
<b>Pathway:</b>	Vitamin D Related/Nuclear Receptor; Apoptosis		
<b>Storage:</b>	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : 90 mg/mL (116.68 mM; Need ultrasonic)

Concentration	Mass		
	1 mg	5 mg	10 mg
1 mM	1.2964 mL	6.4822 mL	12.9645 mL
5 mM	0.2593 mL	1.2964 mL	2.5929 mL
10 mM	0.1296 mL	0.6482 mL	1.2964 mL

Please refer to the solubility information to select the appropriate solvent.

### BIOLOGICAL ACTIVITY

#### Description

BWA-522 is an orally available small molecule protein-targeting chimera (PROTACs) with significant degradation effect on AR-FL and AR-V7. BWA-522 antagonizes the N-terminal domain (AR-NTD) of the androgen receptor (Androgen Receptor) and induces apoptosis in PC cells. BWA-522 inhibits tumor growth in LNCaP xenograft model studies (60 mg/kg, po; TGI=76%). The efficiencies of BWA-522 in degrading AR-V7 and AR-FL were 77.3% (1 μM) and 72.0% (5 μM) in VCaP and LNCaP cells, respectively<sup>[1]</sup>.

### REFERENCES

[1]. Zhang B, et al. Discovery of BWA-522, a First-in-Class and Orally Bioavailable PROTAC Degradator of the Androgen Receptor Targeting N-Terminal Domain for the Treatment of Prostate Cancer. J Med Chem. 2023 Aug 24;66(16):11158-11186..

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

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