



## Veliparib (dihydrochloride)

**Catalog No: tcsc0077** 

Available Sizes
Size: 5mg
Size: 10mg
Size: 50mg
Size: 100mg
Size: 200mg
Size: 500mg
Size: 1g
Specifications
<b>CAS No:</b> 912445-05-7
Formula: C <sub>13</sub> H <sub>18</sub> Cl <sub>2</sub> N <sub>4</sub> O
Pathway: Epigenetics;Cell Cycle/DNA Damage;Autophagy
Target: PARP;PARP;Autophagy
Purity / Grade: >98%
<b>Solubility:</b> DMSO : ≥ 3.2 mg/mL (10.09 mM): H2O : ≥ 50 mg/mL (157.62 mM)





## **Alternative Names:**

ABT-888 dihydrochloride

## **Observed Molecular Weight:**

317.21

## **Product Description**

Veliparib (dihydrochloride) is a potent inhibitor of **PARP1** and **PARP2** with **K**<sub>i</sub> of 5.2 nM and 2.9 nM in cell-free assays, respectively.

IC50 & Target: Ki: 5.2 nM (PARP1), 2.9 nM (PARP2)[1]

In Vitro: Veliparib is inactive to SIRT2 (>5  $\mu$ M)<sup>[1]</sup>. Veliparib inhibits the PARP activity with EC<sub>50</sub> of 2 nM in C41 cells<sup>[2]</sup>. Veliparib can decrease the PAR levels in both irradiated and nonirradiated H460 cells. Veliparib reduces clonogenic survival and inhibits DNA repair by PARP-1 inhibition in H460 cells. Veliparib increases apoptosis and autophagy in H460 cells when combination with radiation <sup>[3]</sup>. Veliparib inhibits PARP activity in H1299, DU145 and 22RV1 cells and the inhibition is independent of p53 function. Veliparib (10  $\mu$ M) suppresses the surviving fraction (SF) by 43% in the clonogenic H1299 cells. Veliparib shows effective radiosensitivity in oxic H1299 cells. Veliparib can attenuate the SF of hypoxic-irradiated cells including H1299, DU145 and 22RV1<sup>[4]</sup>.

In **Vivo:** The oral bioavailability of Veliparib is 56%-92% in mice, SD rats, beagle dogs, and cynomolgus monkeys after oral administration<sup>[1]</sup>. Veliparib (25 mg/kg, i.p.) can improve tumor growth delay in a NCI-H460 xenograft model. Combination with radiation, veliparib decreases the tumor vessel formation<sup>[3]</sup>. Veliparib reduces intratumor PAR levels by more than 95% at a dose of 3 and 12.5 mg/kg in A375 and Colo829 xenograft models and the suppression can be maintained over time<sup>[4]</sup>.

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