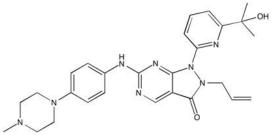
# **Tribioscience**

## Adavosertib (MK-1775), Wee1 checkpoint kinase inhibitor

| Catalog      | Unit  |
|--------------|-------|
| TBI5462-5MG  | 5 mg  |
| TBI5462-25MG | 25 mg |

## **Product Details**

Formal Name: 1,2-Dihydro-1-[6-(1-hydroxy-1-methylethyl)-2-pyridinyl]-6-[[4-(4-methyl-1-piperazinyl)phenyl]amino]-2-(2-propen-1-yl)-3H-pyrazolo[3,4-d]pyrimidin-3-one Alternate Names: MK-1775; AZD 1775 Molecular Formula: C<sub>27</sub>H<sub>32</sub>N<sub>8</sub>O<sub>2</sub> Formula Weight: 500.60 CAS Number: 955365-80-7 **Purity:** >98% Formulation: powder Solubility: Soluble in DMSO (70 mg/ml) Storage: -20°C **Stability:**  $\geq 2$  years.



#### **Applications**

Wee1 checkpoint kinase inhibitor

#### **Functions**

Inhibits Weel tyrosine kinase (IC50 = 5.2 nM) thus preventing phosphorylation of CDC2 and abrogating the  $G_2$  DNA damage checkpoint, sensitizing a variety of tumor cells to DNA damaging agents. Adavosertib also blocks Weel phosphorylation of E3 ubiquitin ligase SKP2 in human cells, ultimately preventing degradation of CDKs and further allowing cell cycle progression. Stimulates anti-tumor immunity and enhances sensitivity to immune checkpoint blockade by activating ERV and the dsRNA pathway. Potentiates sensitivity of tumors to PARP inhibitors.

### **Application Procedures**

First dissolved in DMSO (70 mg/ml), then diluted to aqueous buffer. Solutions in DMSO may be stored at -20°C for up to 3 months.

For research use only.