

## Catalog # 10-4823 Adavosertib

CAS# 955365-80-7

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; MK-1775; AZD1775

## Lot # X109622

Inhibits Wee1 tyrosine kinase ( $IC_{50} = 5.2$  nM) thus preventing phosphorylation of CDC2 and abrogating the  $G_2$  DNA damage checkpoint, sensitizing a variety of tumor cells to DNA damaging agents.<sup>1</sup> Adavosertib also blocks Wee1 phosphorylation of E3 ubiquitin ligase SKP2 in human cells, ultimately preventing degradation of CDKs and further allowing cell cycle progression.<sup>2</sup> Stimulates anti-tumor immunity and enhances sensitivity to immune checkpoint blockade by activating ERV and the dsRNA pathway.<sup>3</sup> Potentiates sensitivity of tumors to PARP inhibitors.<sup>4</sup>

- 1) Hirai et al. (2009) Small-molecule inhibition of Wee1 kinase by MK-1775 selectively sensitizes p53-deficient tumor cells to DNA-damaging agents; Mol. Cancer Ther. **8** 2992
- 2) Pan et al. (2021) A novel WEE1 pathway for replication stress responses; Nat. Plants 7 209
- 3) Guo et al. (2022) WEE1 inhibition induces anti-tumor immunity by activating ERV and the dsRNA pathway; J. Exp. Med. **219** e20210789
- 4) Seo et al. (2021) Inhibition of WEE1 Potentiates Sensitivity to PARP Inhibitor in Biliary Tract Cancer; Cancer Res. Treat. Epub ahead of print

## PHYSICAL DATA

 $\begin{tabular}{lll} Molecular Weight: & 500.61 \\ Molecular Formula: & $C_{27}H_{32}N_8O_2$ \\ Purity: & $>98\% \ by \ HPLC$ \\ NMR: (Conforms) \\ \end{tabular}$ 

DMSO (70 mg/ml)

Physical Description: Yellow solid

Solubility:

Storage and Stability: Store as supplied at -20°C for up to 2 years from the date of purchase. Solutions in

DMSO may be stored at -20°C for up to 3 months.

Materials provided by Focus Biomolecules are for laboratory research use only and are not intended for human or veterinary applications.