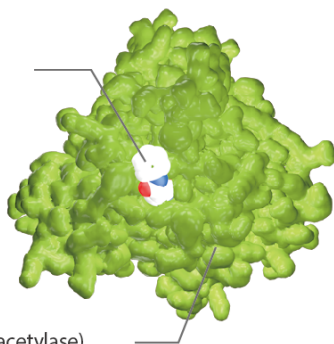


Wee1

HDAC Inhibitor:
Vorinostat (SAHA)



HDAC (Histone deacetylase)

H2B at tyrosine 37 residue which regulates global expression of histones.

Wee1 is a nuclear kinase belonging to the Ser/Thr family of protein kinases in the fission yeast *Schizosaccharomyces pombe* (*S. pombe*). Wee1 has a molecular mass of 96 kDa and it is a key regulator of cell cycle progression. Wee1 influences cell size by inhibiting the entry into mitosis, through inhibiting Cdk1. Wee1 has homologues in many other organisms, including mammals. Wee1 inhibits Cdk1 by phosphorylating it on two different sites, Tyr15 and Thr14. Cdk1 is crucial for the cyclin-dependent passage of the various cell cycle checkpoints. At least three checkpoints exist for which the inhibition of Cdk1 by Wee1 is important: G₂/M checkpoint, Cell size checkpoint, DNA damage checkpoint. Wee1 is shown to phosphorylate histone

Wee1 Inhibitors & Modulators

Adavosertib

(AZD1775; MK-1775)

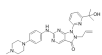
Cat. No.: HY-10993

Bioactivity: Adavosertib (AZD-1775; MK-1775) is a potent **Wee1** inhibitor with an **IC₅₀** of 5.2 nM.

Purity: 99.96%

Clinical Data: Phase 2

Size: 10mM x 1mL in DMSO,
5 mg, 10 mg, 50 mg, 100 mg



PD0166285

Cat. No.: HY-13925

Bioactivity: PD0166285 is a **WEE1** inhibitor and a weak **Myt1** inhibitor with **IC₅₀s** of 24 and 72 nM, respectively.

Purity: 99.46%

Clinical Data: No Development Reported

Size: 10mM x 1mL in DMSO,
5 mg, 10 mg, 25 mg, 50 mg, 100 mg

