Applications

IHC,IF,ELISA



WEE1 (Phospho Ser53) Rabbit pAb

Reactivity

CatalogNo: YP1064

Key Features

Host Species

Rabbit
 Human, Mouse, Rat

MW Isotype • 72kD (Calculated) • IgG

Recommended Dilution Ratios

IHC 1:100-1:300 ELISA 1:20000 IF 1:50-200

Storage

Storage* -15°C to -25°C/1 year(Do not lower than -25°C)

Formulation Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Basic Information

Clonality Polyclonal

Immunogen Information

Immunogen The antiserum was produced against synthesized peptide derived from human WEE1

around the phosphorylation site of Ser53. AA range:19-68

Specificity Phospho-Wee 1 (S53) Polyclonal Antibody detects endogenous levels of Wee 1 protein

only when phosphorylated at S53. The name of modified sites may be influenced by many factors, such as species (the modified site was not originally found in human samples) and the change of protein sequence (the previous protein sequence is incomplete, and the protein sequence may be prolonged with the development of protein sequencing technology). When naming, we will use the "numbers" in historical reference to keep the sites consistent with the reports. The antibody binds to the following modification

sequence (lowercase letters are modification sites):EDsAF

| Target Information

Gene name

WEE1

Protein Name

Wee1-like protein kinase

Organism	Gene ID	UniProt ID
Human	<u>7465;</u>	<u>P30291;</u>
Mouse	22390;	<u>P47810;</u>
Rat	<u>308937</u> ;	<u>Q63802</u> ;

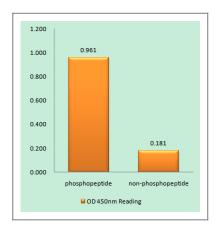
Cellular Localization Nucleus.

Tissue specificity Amygdala, Blood, Epithelium, Human uterus endothel primary cell culture, Placenta, Skin,

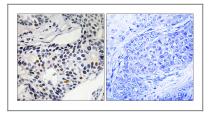
Function

Catalytic activity:ATP + a [protein]-L-tyrosine = ADP + a [protein]-L-tyrosine phosphate.,cofactor:Binds 2 magnesium ions per subunit.,enzyme regulation:Synthesis is increased during S and G2 phases, presumably by an increase in transcription; activity is decreased by phosphorylation during m phase. Protein levels fall in M phase as a result of decreased synthesis combined with degradation. Activity seems to be negatively regulated by phosphorylation upon entry into mitosis, although N-terminal phosphorylation might also regulate the protein stability via protection from proteolysis or might regulate the subcellular location., Function: May act as a negative regulator of entry into mitosis (G2 to M transition) by protecting the nucleus from cytoplasmically activated cyclin B1-complexed CDC2 before the onset of mitosis. Its activity increases during S and G2 phases and decreases at M phase when it is hyperphosphorylated. A correlated decrease in protein level occurs at M/G1 phase, probably due to its degradation. Specifically phosphorylates and inactivates cyclin B1-complexed CDC2 reaching a maximum during G2 phase and a minimum as cells enter M phase. Phosphorylation of cyclin B1-CDC2 occurs exclusively on 'Tyr-15' and phosphorylation of monomeric CDC2 does not occur., PTM: Phosphorylated during M and G1 phases. Also autophosphorylated., PTM: Ubiquitinated and degraded at the onset of G2/M phase., similarity: Belongs to the protein kinase superfamily., similarity: Belongs to the protein kinase superfamily. Ser/Thr protein kinase family. WEE1 subfamily, similarity: Contains 1 protein kinase domain.,

Validation Data



Enzyme-Linked Immunosorbent Assay (Phospho-ELISA) for Immunogen Phosphopeptide (Phospho-left) and Non-Phosphopeptide (Phospho-right), using WEE1 (Phospho-Ser53) Antibody



Immunohistochemistry analysis of paraffin-embedded human breast carcinoma, using WEE1 (Phospho-Ser53) Antibody. The picture on the right is blocked with the phospho peptide.

| Contact information

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Please scan the QR code to access additional product information:

WEE1 (Phospho

WEE1 (Phospho Ser53) Rabbit pAb

For Research Use Only. Not for Use in Diagnostic Procedures.

Antibody | ELISA Kits | Protein | Reagents