

Cdk1/Cdc2 (phospho-Y15) polyclonal antibody

Catalog: BS5058

Host: Rabbit

Reactivity: Human, Mouse, Rat

BackGround:

Cdc2, an evolutionarily conserved serine/threonine-specific protein kinase, is essential in the cell cycle transition from G2 to M phase. Cdc2 is regulated by association with B-type cyclins and by reversible phosphorylation. Cyclin B binding facilitates the phosphorylation of Cdc2 p34 on three regulatory sites: threonine 14, tyrosine 15, and threonine 161. In higher eukaryotes, Cdc2 is negatively regulated by phosphorylation of two residues located in the ATP-binding site, Thr 14 and Tyr 15. Cdc2 is positively regulated by the cyclin-dependent phosphorylation of Thr 161. Both phosphorylation and de-phosphorylation at Thr 161 are required for progression through the cell cycle.

Product:

1 mg/ml in Phosphate buffered saline (PBS) with 0.05% sodium azide, approx. pH 7.2.

Molecular Weight:

~ 34 kDa

Swiss-Prot:

P06493

Purification&Purity:

The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen and the purity is > 95% (by SDS-PAGE).

Applications:

WB: 1:500~1:1000

IHC: 1:50~1:200

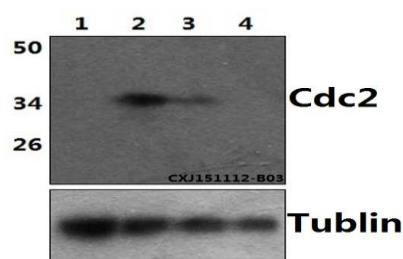
Storage&Stability:

Store at 4 °C short term. Aliquot and store at -20 °C long term. Avoid freeze-thaw cycles.

Specificity:

p-Cdk1/Cdc2 (Y15) polyclonal antibody detects endogenous levels of Cdk1/Cdc2 protein only when phosphorylated at Tyr15.

DATA:



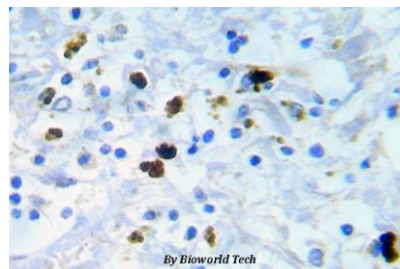
Western blot (WB) analysis of Cdc2 (phospho-Y15) polyclonal antibody at 1:500 dilution

Lane1:Hela whole cell lysate(40ug)

Lane2:Hela treated with UV (10min) whole cell lysate(40ug)

Lane3:Hela treated with UV (20min) whole cell lysate(40ug)

Lane4:Hela treated with UV (30min) whole cell lysate(40ug)



Immunohistochemistry (IHC) analyzes of p-Cdk1/Cdc2 (Y15) pAb in paraffin-embedded human breast carcinoma tissue.

Note:

For research use only, not for use in diagnostic procedure.

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