

Chk2 (phospho Thr68) Polyclonal Antibody

Catalog No: YP0065

Reactivity: Human; Mouse; Rat

Applications: IF;WB;IHC;ELISA

Target: Chk2

Fields: >>Cell cycle;>>p53 signaling pathway;>>Cellular senescence;>>Human T-cell

leukemia virus 1 infection

Gene Name: CHEK2

Protein Name: Serine/threonine-protein kinase Chk2

O96017

Q9Z265

Human Gene Id: 11200

Human Swiss Prot

No:

Mouse Gene Id: 50883

Mouse Swiss Prot

No:

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

Chk2 around the phosphorylation site of Thr68. AA range:35-84

Specificity: Phospho-Chk2 (T68) Polyclonal Antibody detects endogenous levels of Chk2

protein only when phosphorylated at T68.

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

Source: Polyclonal, Rabbit, IgG

Dilution: IF 1:50-200 WB 1:500 - 1:2000. IHC 1:100 - 1:300. ELISA: 1:20000. Not yet

tested in other applications.

Purification: The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.



Concentration: 1 mg/ml

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

Observed Band: 61kD

Cell Pathway: Cell_Cycle_G1S;Cell_Cycle_G2M_DNA;p53;

Background: In response to DNA damage and replication blocks, cell cycle progression is

this gene is a cell cycle checkpoint regulator and putative tumor suppressor. It contains a forkhead-associated protein interaction domain essential for activation in response to DNA damage and is rapidly phosphorylated in response to replication blocks and DNA damage. When activated, the encoded protein is known to inhibit CDC25C phosphatase, preventing entry into mitosis, and has

halted through the control of critical cell cycle regulators. The protein encoded by

been shown to stabilize the tumor suppressor protein p53, leading to cell cycle arrest in G1. In addition, this protein interacts with and phosphorylates BRCA1, allowing BRCA1 to restore survival after DNA damage. Mutations in this gene

have been linked with Li-Fraumeni syndrome, a highly penetrant familial cancer

phenotype usually associated with inherited mutati

Function: catalytic activity:ATP + a protein = ADP + a

phosphoprotein.,cofactor:Magnesium.,disease:Defects in CHEK2 are associated

with Li-Fraumeni syndrome 2 (LFS2) [MIM:609265]; a highly penetrant familial

cancer phenotype usually associated with inherited mutations in p53/TP53., disease: Defects in CHEK2 are found in some patients with

osteosarcoma (OSRC) [MIM:259500].,disease:Defects in CHEK2 are found in

some patients with prostate cancer (CaP) [MIM:176807].,enzyme

regulation:Rapidly phosphorylated on Thr-68 by MLTK in response to DNA

damage and to replication block. Kinase activity is also up-regulated by

autophosphorylation.,function:Regulates cell cycle checkpoints and apoptosis in response to DNA damage, particularly to DNA double-strand breaks. Inhibits CDC25C phosphatase by phosphorylation on 'Ser-216', preventing the entry into

mitosis. May also play a role in meiosis. Regulates the TP53

Subcellular [Isoform 2 Nucleus.; [

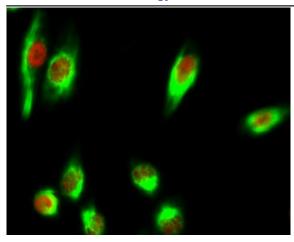
[Isoform 2]: Nucleus. Isoform 10 is present throughout the cell.; [Isoform 4]: Nucleus.; [Isoform 7]: Nucleus.; [Isoform 9]: Nucleus.; [Isoform 12]: Nucleus.; Nucleus, PML body. Nucleus, nucleoplasm. Recruited into PML bodies together

with TP53.

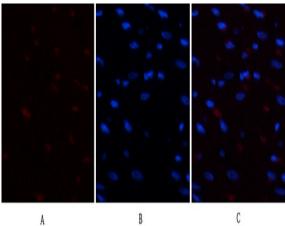
Expression: High expression is found in testis, spleen, colon and peripheral blood leukocytes.

Low expression is found in other tissues.

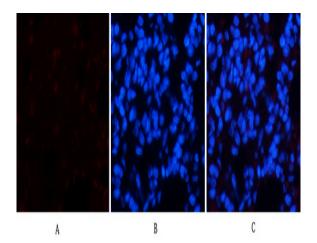
Products Images



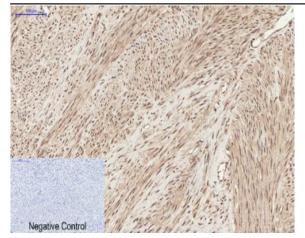
Immunofluorescence analysis of Hela cell. 1,Chk2 (phospho Thr68) Polyclonal Antibody(red) was diluted at 1:200(4° overnight). HAO1 Monoclonal Antibody(Mix)(green) was diluted at 1:200(4° overnight). 2, Goat Anti Rabbit Alexa Fluor 594 Catalog:RS3611 was diluted at 1:1000(room temperature, 50min). Goat Anti Mouse Alexa Fluor 488 Catalog:RS3208 was diluted at 1:1000(room temperature, 50min).



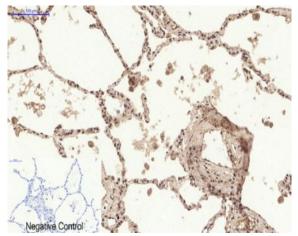
Immunofluorescence analysis of rat-heart tissue. 1,Chk2 (phospho Thr68) Polyclonal Antibody(red) was diluted at 1:200(4°C,overnight). 2, Cy3 labled Secondary antibody was diluted at 1:300(room temperature, 50min).3, Picture B: DAPI(blue) 10min. Picture A:Target. Picture B: DAPI. Picture C: merge of A+B



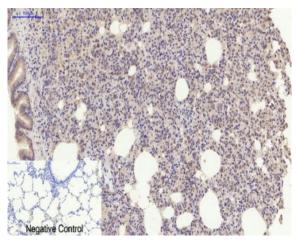
Immunofluorescence analysis of rat-lung tissue. 1,Chk2 (phospho Thr68) Polyclonal Antibody(red) was diluted at 1:200(4°C,overnight). 2, Cy3 labled Secondary antibody was diluted at 1:300(room temperature, 50min).3, Picture B: DAPI(blue) 10min. Picture A:Target. Picture B: DAPI. Picture C: merge of A+B



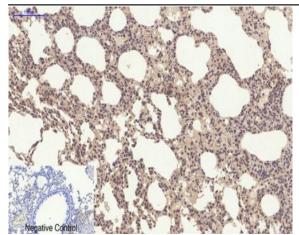
Immunohistochemical analysis of paraffin-embedded Humanuterus tissue. 1,Chk2 (phospho Thr68) Polyclonal Antibody was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room tempeRature, 30min). Negative control was used by secondary antibody only.



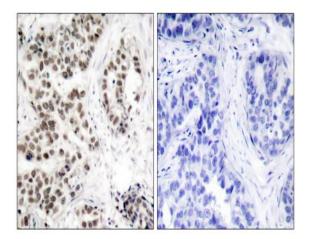
Immunohistochemical analysis of paraffin-embedded Humanlung tissue. 1,Chk2 (phospho Thr68) Polyclonal Antibody was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room tempeRature, 30min). Negative control was used by secondary antibody only.



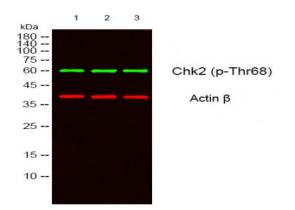
Immunohistochemical analysis of paraffin-embedded Rat-lung tissue. 1,Chk2 (phospho Thr68) Polyclonal Antibody was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room tempeRature, 30min). Negative control was used by secondary antibody only.



Immunohistochemical analysis of paraffin-embedded Mouse-lung tissue. 1,Chk2 (phospho Thr68) Polyclonal Antibody was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room tempeRature, 30min). Negative control was used by secondary antibody only.



Immunohistochemistry analysis of paraffin-embedded human lung carcinoma, using Chk2 (Phospho-Thr68) Antibody. The picture on the right is blocked with the phospho peptide.



Western blot analysis of lysates from 1) 293T, 2) HELA cells, (Green) primary antibody was diluted at 1:1000, 4° over night, secondary antibody(cat:RS23920)was diluted at 1:10000, 37° 1hour. (Red) Actin β Monoclonal Antibody(5B7) (cat:YM3028) antibody was diluted at 1:5000 as loading control, 4° over night,secondary antibody(cat:RS23710)was diluted at 1:10000, 37° 1hour.