

Rabbit Anti-CDK1 antibody

SLM-52026R

Product Name CDK1

Chinese Name 周期素依赖性激酶 1Recombinant rabbit monoclonal anti

Alias Cdc 2; Cdc2; CDC28A; CDK 1; CDK-1; CDK1_HUMAN; CDKN1; CELL CYCLE CONTROL PROTEIN 2; CDC2; Cell division control protein 2; Cell division control protein 2 homolog; Cell division cycle 2 protein; Cell division cycle 2 protein kinase; Cell division cycle 2 protein kinase 1; Cell Division Cycle 2 Protein; Cyclin Dependent Kinase 1; Cyclin-dependent kinase 1; DKFZp686L20222; MGC111195; p34 Cdk1; p34 protein kinase; P34

Research Area Tumour Cell biology Signal transduction Cyclin Kinases and Phosphatases

Immunogen Species Rabbit

Clonality Monoclonal

Clone NO. 3E12

React Species Human,Mouse,Rat

Applications WB=1:500-2000,IHC-P=1:100-500,IHC-F=1:100-500,ICC/IF=1:50-200,IF=1:100-500,Flow-Cytometry
(Paraffin sections need antigen repair)
not yet tested in other applications.
optimal dilutions/concentrations should be determined by the end user.

Theoretical molecular weight 34kDa

Cellular localization The nucleus cytoplasmic

Form Liquid

Concentration 1mg/ml

immunogen Recombinant human CDK1 protein, full length

Lsotype IgG

Purification affinity purified by Protein A

Buffer Solution 1M TBS(pH7.4) with 1% BSA, 3% Proclin300 and 50% Glycerol.

Storage Shipped at 4°C. Store at -20 °C for one year. Avoid repeated freeze/thaw cycles.

Attention This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic.

applications.

PubMed

[PubMed](#)

The cell division control protein cdc2, also known as cyclin dependent kinase 1 (Cdk1) or p34/cyclin B complex, plays a key role in the control of the eukaryotic cell cycle, where it is required for entry into S phase and G2 to M phase transition. Cdc2 exists as a complex with both cyclin A and cyclin B. The best characterized of these associations is the Cdc2 p34 cyclin B complex, which is required for the G2 to M phase transition. Activation of Cdc2 is controlled at several steps including cyclin binding and phosphorylation of threonine 161. However, a critical regulatory step in activating cdc2 during progression into mitosis appears to be dephosphorylation of Tyr15 and Tyr14. Phosphorylation at Tyr15 and inhibition of Cdc2 is carried out by WEE1 and MCK1 kinases while Tyr15 dephosphorylation and activation of Cdc2 is carried out by the cdc25 phosphatase. The isoform CDC2deltaT is found in breast cancer tissues. Furthermore, cdc2/Cdk1 is a key mediator of cell death in brain development and degeneration.

Function:

Plays a key role in the control of the eukaryotic cell cycle by modulating the centrosome cycle and mitotic onset; promotes G2-M transition, and regulates G1 progress and G1-S transition via association with multiple interphase cyclins. Required in higher cells for entry into S-phase and mitosis. Phosphorylation of CDC25A/actopaxin, APC, AMPH, APC, BARD1, Bcl-xL/BCL2L1, BRCA2, CALD1, CASP8, CDC20, CDC25A, CDC25C, CC2D1A, CSNK2 proteins/CKII, FZR1/CDH1, CDK7, CEBPB, DMD/dystrophin, EEF1 proteins/EF-1, EZH2, KIF11/EG5, EGFR, FANCG, FOS, GFAP, GOLGA2/GM130, GRASP1, UBE2A/hHR6A, HIST1H1 proteins/histone H1, HMGA1, HIVEP1, LMNA, LMNB, LMNC, LBR, LATS1, MAP1B, MAP4, MARCKS, MCM2, MCM4, MKLP1, NEFH, NFIC, NPC/nuclear pore complex, PITPNM1/NIR2, NPM1, NCL, NUCKS1, NPM1/nucleolar phosphoprotein, ORC1, PRKAR2A, EEF1E1/p18, EIF3F/p47, p53/TP53, NONO/p54NRB, PAPOLA, PLEC/pleckstrin, UL40/R2, RAB4A, RAP1GAP, RCC1, RPS6KB1/S6K1, KHDRBS1/SAM68, ESPL1, SKI, BIRC5/survivin, STIP1, TEX14, beta-tubulins, MAPT/TAU, NEDD1, VIM/vimentin, TK1, FOXO1, RUNX1/AML1 and RUNX2. CDK1/CDC2-cyclin-B controls pronuclear union in interphase fertilization. Essential for early stages of embryonic development. During G2 and early mitosis, CDC25A/B/C-mediated dephosphorylation activates CDK1/cyclin complexes which phosphorylate several substrates that include at least centrosome separation, Golgi dynamics, nuclear envelope breakdown and chromosome condensation. Once chromosomes are condensed and aligned at the metaphase plate, CDK1 activity is switched off by WEE1- and PKMYT1-mediated phosphorylation to allow sister chromatid separation, chromosome decondensation, reformation of the nuclear envelope and cytokinesis. Inactivated by PKR/EIF2AK1. WEE1-mediated phosphorylation upon DNA damage to stop cell cycle and genome replication at G2/M checkpoint thus facilitating DNA repair. Reactivated after successful DNA repair through WIP1-mediated signaling leading to CDC25A/B/C-mediated dephosphorylation and restoring cell cycle progression. In proliferating cells, CDK1-mediated FOXO1 phosphorylation at the G2-M phase represses FOXO1 activity, leading to cell death of postmitotic neurons. The phosphorylation of beta-tubulins regulates microtubule dynamics during mitosis. NEDD1 phosphorylation promotes PLK1-mediated NEDD1 phosphorylation and subsequent targeting of the gamma-tubulin ring complex (gTuRC) to the centrosome, an important step for spindle formation. In addition, CC2D1A phosphorylation regulates CC2D1A subcellular localization and association with SCC1/RAD21 and centriole cohesion during mitosis. The phosphorylation of CDC25A/B/C is essential for cell cycle progression.

Product Detail

of Bcl-xL/BCL2L1 after prolonged G2 arrest upon DNA damage triggers apoptosis. In contrast, phosphorylation during mitosis prevents its activation by proteolysis and subsequent apoptosis. Phosphorylation occurs in cancer cell lines, as well as in primary breast tissues and lymphocytes. Phosphorylation promotes H3K27me3 maintenance and epigenetic gene silencing. CALD1 phosphorylation promotes Schwann cell migration during peripheral nerve regeneration.

Subunit:

Forms a stable but non-covalent complex with a regulatory subunit and with a cyclin. Interacts with cyclins (CCNB1, CCNB2 and CCNB3) to form a serine/threonine kinase holoenzyme complex also known as maturation promoting factor (MPF). The cyclin subunit imparts substrate specificity to the complex. Forms CDK1-cyclin-D and CDK1-cyclin-E complexes that phosphorylate RB1 in vitro. Binds to REB1 and other transcription factors such as FOXO1 and RUNX2. Promotes G2-M transition when in complex with cyclin-B. Interacts with DLGAP5. Binds to the CDK inhibitors CDKN1A/p21 and CDKN1B/p27. CDKN1A/p21 is unable to complex with cyclin-B1 and also fails to bind to CDKN1A/p21. Interacts with catalytic subunit of CCNB1 and RALBP1 during mitosis to form an endocytotic complex during interphase. Associates with cyclins-A and B1 during S-phase in regenerating hepatocytes. Interacts with FANCC. Interacts with CDK1; this interaction recruits CDK1 to centrosomes.

Subcellular Location:

Nucleus. Cytoplasm. Mitochondrion. Cytoplasm, cytoskeleton, centrosome. Note=Cytoplasmic during interphase. Reversibly translocated from cytoplasm to nucleus when phosphorylated before G2-M transition. Accumulates in mitochondria when associated with cyclin-B1. Accumulates in mitochondria in G2-arrested cells upon DNA-damage.

Tissue Specificity:

Isoform 2 is found in breast cancer tissues.

Post-translational modifications:

Phosphorylation at Thr-161 by CAK/CDK7 activates kinase activity. Phosphorylation at Thr-14 and Tyr-15 by PKMYT1 prevents nuclear translocation. Phosphorylation at Tyr-15 by WEE1 and WEE2 inhibits protein kinase activity and acts as a negative regulator of entry into mitosis (G2 to M transition). Phosphorylation by PKMYT1 and WEE1 takes place during mitosis to keep CDK1-cyclin-B complex inactive until the end of G2. By the end of G2, PKMYT1 and WEE1 are inactivated, but CDC25A and CDC25B are activated. Dephosphorylation by active CDC25A and CDC25B at Thr-14 and Tyr-15 activates CDK1 activation at the G2-M transition. Phosphorylation at Tyr-15 by WEE2 during oogenesis is required to maintain meiotic arrest in oocytes during the germinal vesicle (GV) stage, a long period of quiescence that dictyate prophase I, leading to prevent meiotic reentry. Phosphorylation by WEE2 is also required for metaphase II exit during egg activation to ensure exit from meiosis in oocytes and promote pronuclear formation. Phosphorylated at Tyr-4 by PKR/EIF2AK2 upon genotoxic stress. This phosphorylation leads to CDK1 polyubiquitination and subsequent proteolysis, thus leading to G2 arrest. In response to UV irradiation, phosphorylation at Tyr-15 by PRKCD activates the G2/M DNA damage checkpoint. Polyubiquitinated upon genotoxic stress.

Similarity:

Belongs to the protein kinase superfamily. CMGC Ser/Thr protein kinase family. CDC2/CDKX

Contains 1 protein kinase domain.

SWISS:
P06493

Gene ID:
983

Database links:

[Entrez Gene: 12534](#) Mouse

[Entrez Gene: 54237](#) Rat

[Omim: 116940](#) Human

[SwissProt:](#) Human

[SwissProt: P11440](#) Mouse

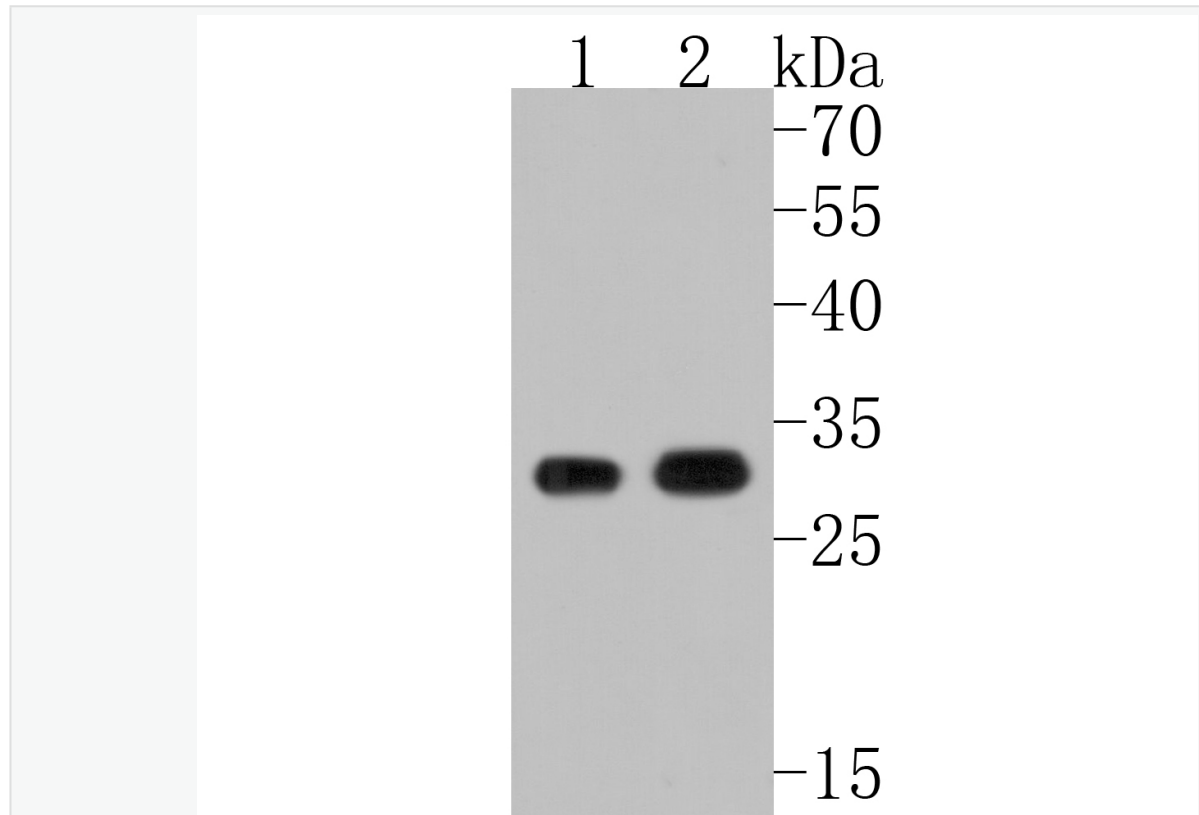
[SwissProt: P39951](#) Rat

[Unigene: 334562](#) Human

[Unigene: 281367](#) Mouse

[Unigene: 6934](#) Rat

**Product
Picture**



Sample:

Lane 1: HepG2 (Human) Cell Lysate at 30 ug

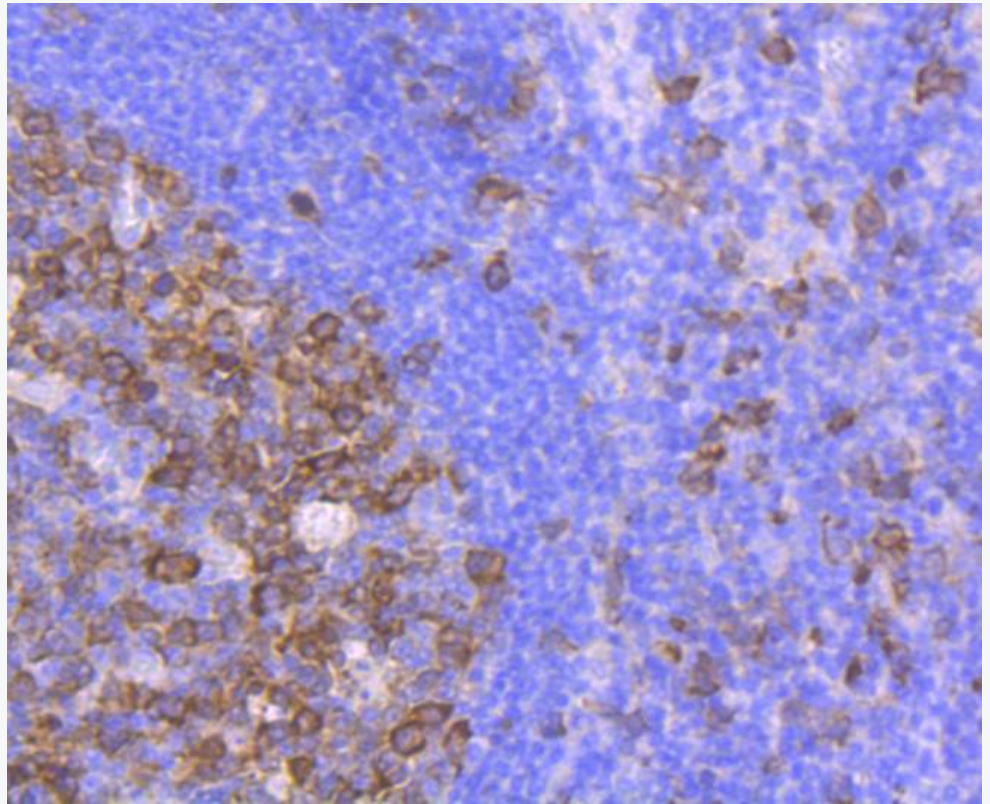
Lane 2: Jurkat (Human) Cell Lysate at 30 ug

Primary: Anti-CDK1 (SLM-52026R) at 1/500 dilution

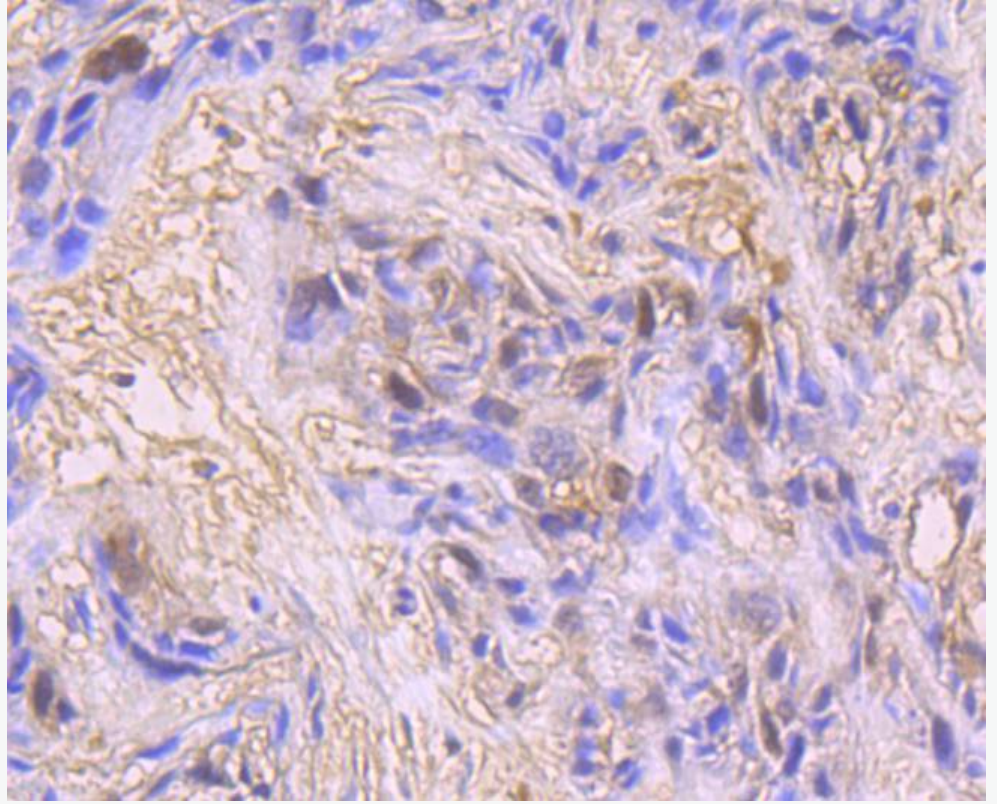
Secondary: Goat Anti-Rabbit IgG - HRP at 1/5000 dilution

Predicted band size: 34 kD

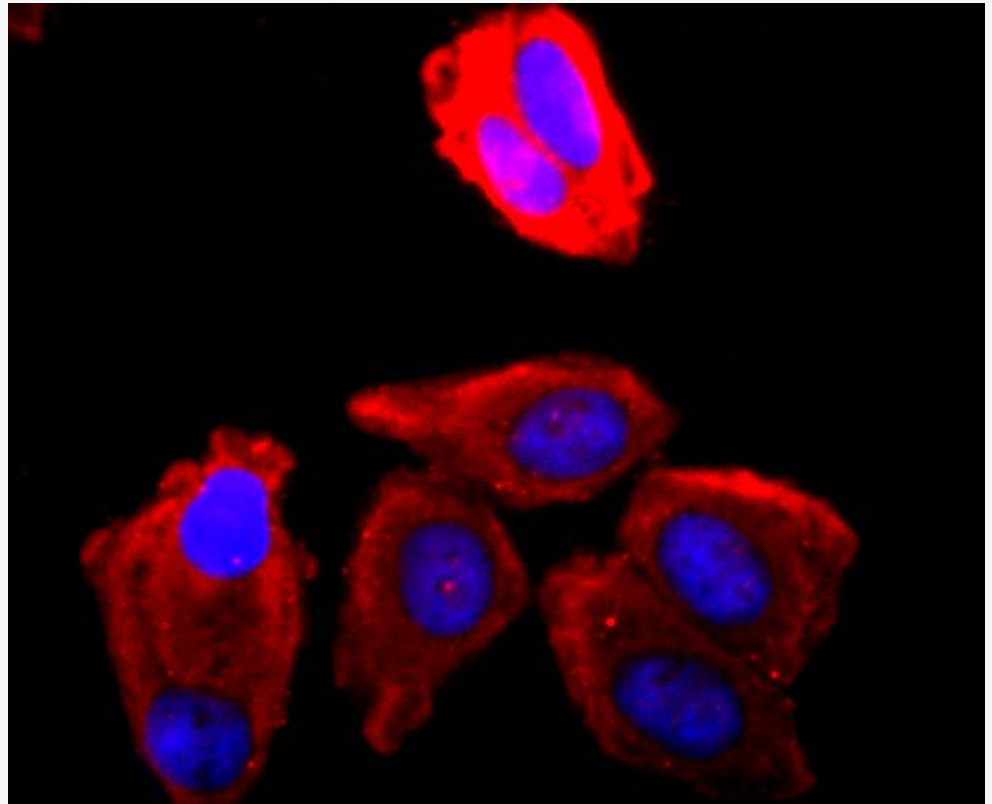
Observed band size: 34 kD



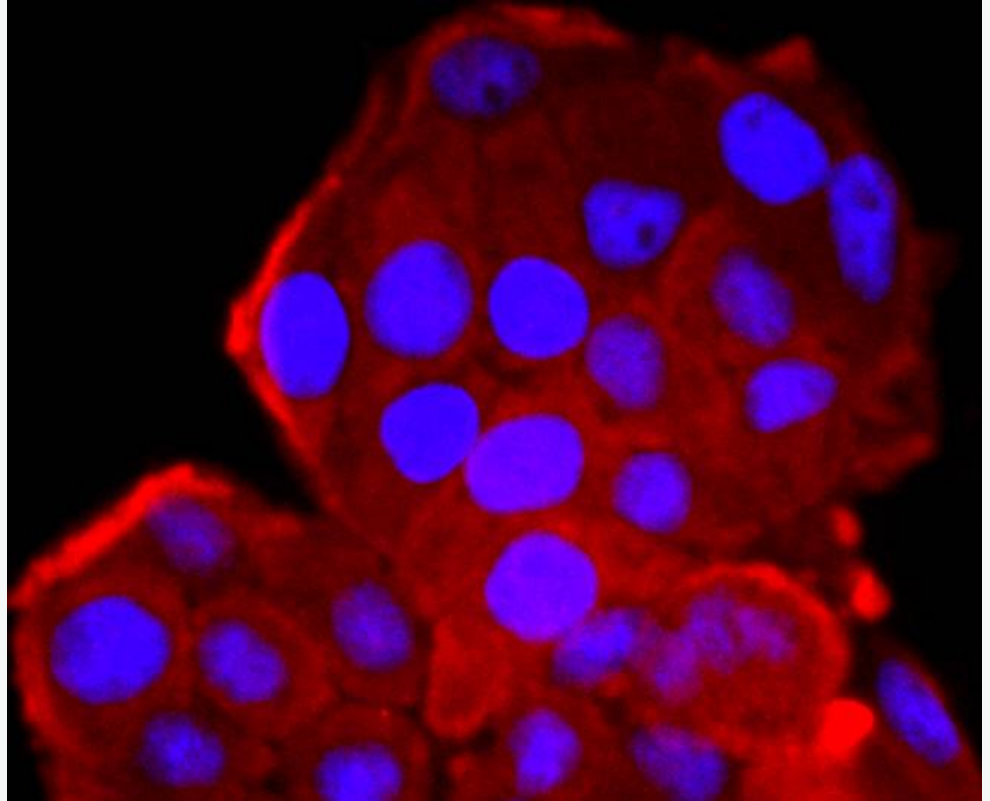
Paraformaldehyde-fixed, paraffin embedded (human tonsil); Antigen retrieval by boiling in so
citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 2
Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (CDK1) M
Antibody, Unconjugated (SLM-52026R) at 1:50 overnight at 4°C, followed by operating accor
Kit(Rabbit) (sp-0023) instructions and DAB staining.



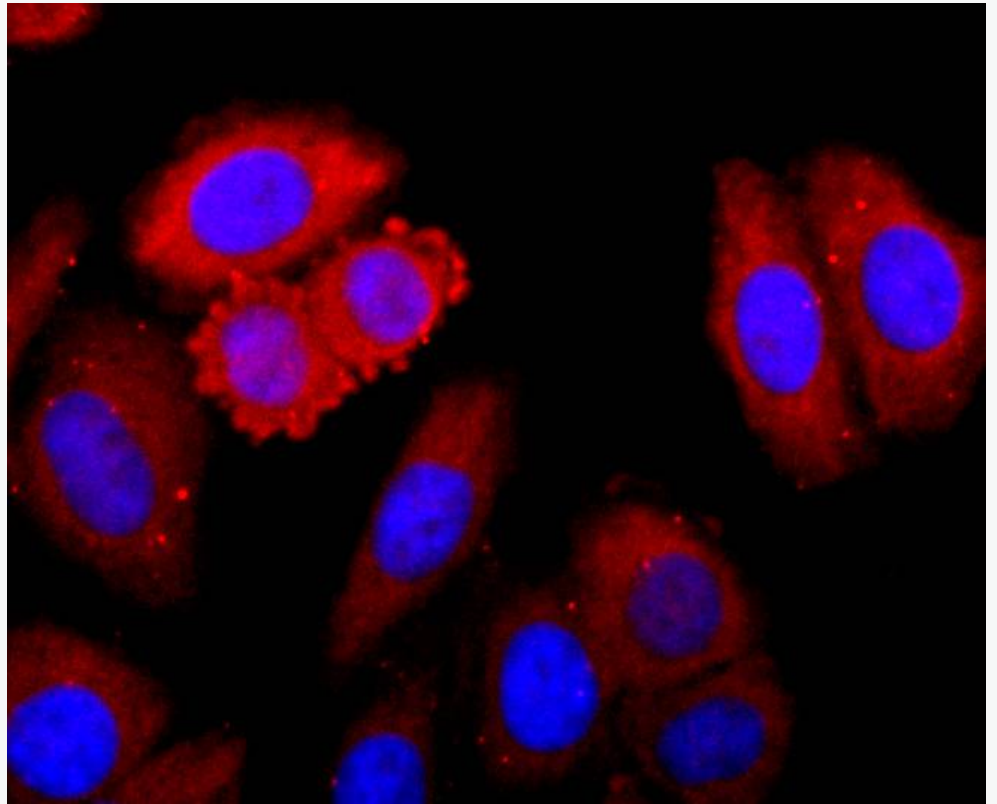
Paraformaldehyde-fixed, paraffin embedded (human breast carcinoma); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 30 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (C) Monoclonal Antibody, Unconjugated (SLM-52026R) at 1:50 overnight at 4°C, followed by op according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



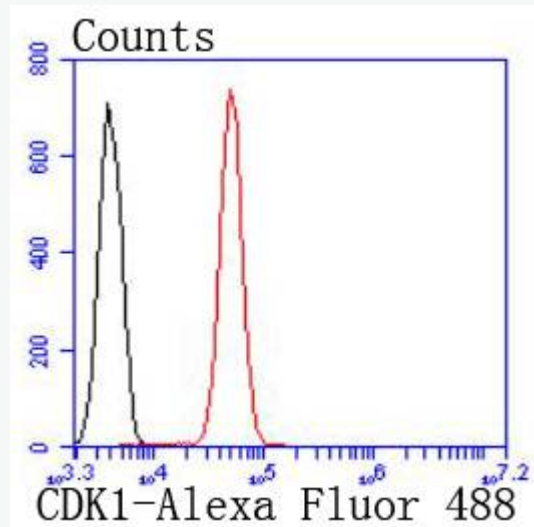
MCF-7 cell; 4% Paraformaldehyde-fixed; Triton X-100 at room temperature for 20 min; Block (normal goat serum, C-0005) at 37°C for 20 min; Antibody incubation with (CDK1) monoclonal Antibody, Unconjugated (SLM-52026R) 1:50, 90 minutes at 37°C; followed by a conjugated Anti-Rabbit IgG antibody at 37°C for 90 minutes, DAPI (blue, C02-04002) was used to stain nuclei.



HeLa cell; 4% Paraformaldehyde-fixed; Triton X-100 at room temperature for 20 min; Blocking (normal goat serum, C-0005) at 37°C for 20 min; Antibody incubation with (CDK1) monoclonal Antibody, Unconjugated (SLM-52026R) 1:50, 90 minutes at 37°C; followed by a conjugated Anti-Rabbit IgG antibody at 37°C for 90 minutes, DAPI (blue, C02-04002) was used to stain nuclei.



HepG2 cell; 4% Paraformaldehyde-fixed; Triton X-100 at room temperature for 20 min; Block (normal goat serum, C-0005) at 37°C for 20 min; Antibody incubation with (CDK1) monoclonal Antibody, Unconjugated (SLM-52026R) 1:50, 90 minutes at 37°C; followed by a conjugated Anti-Rabbit IgG antibody at 37°C for 90 minutes, DAPI (blue, C02-04002) was used to stain nuclei.



Blank control: Jurkat.

Primary Antibody (green line): Rabbit Anti-CDK1 antibody (SLM-52026R)

Dilution: 1:50;

Isotype Control Antibody (orange line): Rabbit IgG .

Secondary Antibody : Goat anti-rabbit IgG-AF488

Dilution: 1:1000.

Protocol

The cells were fixed with 4% PFA (10min at room temperature)and then permeabilized with 95% ice-cold methanol for 20 min at-20°C. The cells were then incubated in 5%BSA to block non-specific protein-protein interactions for 30 min at room temperature .Cells stained with Primary Antibody for 30 min at room temperature. The secondary antibody used for 40 min at room temperature. Acquisition of 20,000 events was performed.