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## Chk1 Ab

Cat.#: AF5014 Concn.: 1mg/ml Mol.Wt.: 56kDa Size: 100ul.200ul Source: Rabbit Clonality: Polyclonal

Application: WB 1:500-1:2000. IF/ICC 1:100-1:500

Reactivity: Human

Purification: The antiserum was purified by peptide affinity

chromatography using SulfoLink™ Coupling Resin (Thermo

Fisher Scientific).

Specificity: Chk1 Ab detects endogenous levels of total Chk1.

Immunogen: A synthesized peptide derived from human Chk1.

Uniprot: 014757

Description: DNA damage induced protein phosphorylation; regulation of

> mitotic centrosome separation; regulation of S phase; peptidyl-threonine phosphorylation; DNA repair; chromatinmediated maintenance of transcription; negative regulation

of mitosis:

Subcellular Location: Cytoskeleton; Nucleus;

Tissue Specificity: Expressed ubiquitously with the most abundant expression

in thymus, testis, small intestine and colon.

Similarity: The autoinhibitory region (AIR) inhibits the activity of the

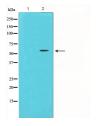
> kinase domain. Belongs to the protein kinase superfamily. CAMK Ser/Thr protein kinase family. NIM1 subfamily.

Storage Condition and

Buffer:

Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Store at -20

°C.Stable for 12 months from date of receipt.



Western blot analysis of Chk1 expression in HeLa whole cell lysates, The lane on the left is treated with the antigen-specific

peptide.



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AF5014 staining HeLa by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25°C. The primary Ab was diluted at 1/200 and incubated with the sample for 1 hour at 37°C. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary Ab.

<code>IMPORTANT:</code> For western blot, incubate membrane with diluted primary Ab in 5% w/v milk , 1X TBS, 0.1% Tween020 at  $4^{\circ}$ C with gentle shaking, overnight.

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