For Research Use Only

Phospho-CHEK2 (Thr68) Recombinant antibody

Catalog Number:81740-1-RR

1 Publications



Basic Information

Catalog Number: 81740-1-RR

Size: 1000 ug/ml Source: Rabbit

Isotype: IgG GenBank Accession Number: BC004207

GeneID (NCBI): 11200 UNIPROT ID: 096017

Full Name: CHK2 checkpoint homolog (S. pombe)

Calculated MW: 61 kDa Observed MW: 65 kDa

Applications

Tested Applications: WB, FC (Intra), ELISA Cited Applications:

WB

Species Specificity:

human
Cited Species:
human

Positive Controls:

WB: MMS treated PC-3 cells, UV treated HeLa cells

Purification Method:

CloneNo.:

Protein A purification

Recommended Dilutions:

WB 1:5000-1:50000

Background Information

Serine/threonine-protein kinase Chk2 (CHEK2) is a serine/threonine kinase which is activated upon DNA damage and is implicated in pathways that govern DNA repair, cell cycle arrest or apoptosis in response to the initial damage. ATM phosphorylates CHEK2 on T68. Phosphorylation on T68 and subsequent full activation of CHEK2 was shown to require priming phosphorylation on adjacent residues by Polo-like kinase 3 (PLK3) and the dualspecificity tyrosine and serine/threoninekinase TTK/hMPS1. Additionally TTK appears to phosphorylate T68. Phosphorylation of T68 promotes the binding of the N-terminal SQ/TQ-rich cluster of one CHEK2 molecule with the FHA domain of another CHEK2 molecule. (PMID: 28553140, PMID: 18004398, PMID: 33322746)

Notable Publications

Author	Pubmed ID	Journal	Application
Yukun Wang	39090319	EMBO Rep	WB

Storage

Storage:

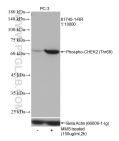
Store at -20°C. Stable for one year after shipment.

Storage Buffer

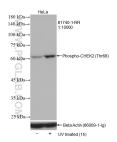
PBS with 0.02% sodium azide and 50% glycerol pH 7.3.

Aliquoting is unnecessary for -20°C storage

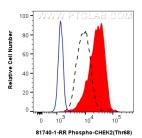
Selected Validation Data



Non-treated PC-3 and MMS treated PC-3 cells were subjected to SDS PAGE followed by western blot with 81740-1-RR (Phospho-CHEK2 (Thr68) antibody) at dilution of 1:10000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with Beta Actin antibody as loading control.



Non-treated HeLa and UV treated HeLa cells were subjected to SDS PAGE followed by western blot with 81740-1-RR (Phospho-CHEK2 (Thr68) antibody) at dilution of 1:10000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with Beta Actin antibody as loading control.



1X10^6 PC-3 cells untreated (dashed lines) or treated with MMS which intracellularly stained with 0.06 ug Phospho-CHEK2 (Thr68) Recombinant antibody (81740-1-RR, Clone:1L2) and CoraLite@488-Conjugated Goat Anti-Rabbit IgG(H+L) (\$A00013-2)(red), or 0.06 ug Rabbit IgG Isotype Control Recombinant Antibody (98136-1-RR, Clone: 240953C9) (blue). Cells were fixed with 4% PFA and permeabilized with 90% MeOH.