For Research Use Only

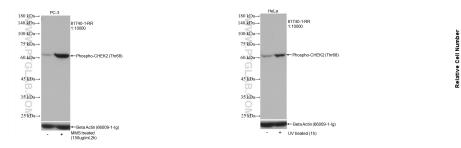
Phospho-CHEK2 (Thr68) Recombinant antibody, PBS Only Catalog Number:81740-1-PBS

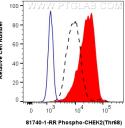


Basic Information	Catalog Number: 81740-1-PBS	GenBank Accession Number: BC004207	Purification Method: Protein A purification
	Size: 1mg/ml	GenelD (NCBI): 11200	CloneNo.: 1L2
	Source: Rabbit	UNIPROT ID: 096017	
	Isotype:Full Name:IgGCHK2 checkpoint homolog (S. por		mbe)
		Calculated MW: 61 kDa	
		Observed MW: 65 kDa	
Applications	Tested Applications: WB, FC (Intra), Indirect ELISA		
	Species Specificity: human		
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)		
Storage	Storage: Store at -80°C. The product is shipped with ice	packs. Upon receipt, store it immediatel	ly at -80°C

For technical support and original validation data for this product please contact: T: 4006900926 E: Proteintech-CN@ptglab.com W: ptgcn.com This product is exclusively available under Proteintech Group brand and is not available to purchase from any other manufacturer.

Selected Validation Data





Non-treated PC-3 and MMS treated PC-3 cells were subjected to SDS PACE 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. This data was developed using the same antibody clone with 81740-1-PBS in a different storage buffer formulation. Non-treated HeLa and UV treated HeLa cells were subjected to SDS PACE 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. This data was developed using the same antibody clone with 81740-1-PBS in a different storage buffer formulation. 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 Isotype Control Recombinant Antibody (98136-1-RR, Clone: 240953C9) (blue). Cells were fixed with 4% PFA and permeabilized with 90% MeOH. This data was developed using the same antibody