[Primary Antibody]

phospho-P53 (Ser15) Rabbit pAb



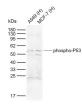
www.bioss.com.cn sales@bioss.com.cn techsupport@bioss.com.cn 400-901-9800

(predicted: Rabbit, Pig, Sheep, Cow, Dog,

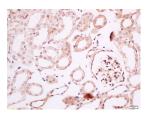
- DATASHEET		400-901-9800	
Host: Rabbit	lsotype: IgG	Applications: WB (1:500-2000)	
Clonality: Polyclonal		IHC-P (1:100-500) IHC-F (1:100-500)	
GenelD: 7157	SWISS: P04637	IF (1:100-500)	
Target: P53 (Ser15)		Flow-Cyt (1µg /test) ICC/IF (1:100)	
Immunogen: KLH conjugated Synthesised phosphopeptide derived from human P53 around the phosphorylation site of Ser15: PL(p-S)QE.			
Purification: affinity purified by Protein A			
Concentration: 1mg/ml			
Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol. Shipped at 4°C. Store at -20°C for one year. Avoid repeated		Predicted MW.: ^{43 kDa}	
freeze/thaw cycles.		Subcellular Location: Cytoplasm ,Nucleus	
transcriptional activ domains. The encod stresses to regulate o cell cycle arrest, apo metabolism. Mutatio human cancers, inclu	tumor suppressor protein containing ation, DNA binding, and oligomerization ed protein responds to diverse cellular expression of target genes, thereby inducing ptosis, senescence, DNA repair, or changes in ons in this gene are associated with a variety of uding hereditary cancers such as Li-Fraumeni ve splicing of this gene and the use of alternate		

promoters result in multiple transcript variants and isoforms. Additional isoforms have also been shown to result from the use of alternate translation initiation codons (PMIDs: 12032546, 20937277). [provided by RefSeq, Feb 2013].

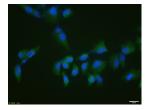
- VALIDATION IMAGES



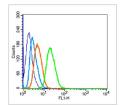
Sample: Lane 1: Human A549 cell lysates Lane 2: Human MCF-7 cell lysates Primary: Antiphospho-P53 (Ser15) (bs-3702R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 43 kDa Observed band size: 53 kDa



Tissue/cell: Mouse kidney tissue; 4% Paraformaldehyde-fixed and paraffinembedded; Antigen retrieval: citrate buffer (0.01M, pH 6.0), Boiling bathing for 15min; Block endogenous peroxidase by 3% Hydrogen peroxide for 30min; Blocking buffer (normal goat serum,C-0005) at 37°C for 20 min; Incubation: Anti-p53 Polyclonal Antibody, Unconjugated(bs-3702R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining



A549 cell; 4% Paraformaldehyde-fixed; Triton X-100 at room temperature for 20 min; Blocking buffer (normal goat serum, C-0005) at 37°C for 20 min; Antibody incubation with (phospho-P53 (Ser15)) polyclonal Antibody, Unconjugated (bs-3702R) 1:100, 90 minutes at 37°C; followed by a conjugated Goat Anti-Rabbit IgG antibody at 37°C for 90 minutes, DAPI (blue, C02-04002) was used to stain the cell nuclei.



Blank control (blue line): MCF 7(fixed with 70% ethanol (Overnight at 4°C) and then

permeabilized with 90% ice-cold methanol for 30 min on ice). Primary Antibody (green line): Rabbit Anti-phospho-P53 (Ser15) antibody (bs-3702R),Dilution: 1µg /10^6 cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody (white blue line): Goat antirabbit IgG-FITC,Dilution: 1µg /test.

- SELECTED CITATIONS -

- [IF=8] Hao, X. et al. TC2N, a novel oncogene, accelerates tumor progression by suppressing p53 signaling pathway in lung cancer.(2018) Cell Death & Differentiation. WB ;. 30254375
- [IF=6.1] Jianfang Wang. et al. Knockdown of NFIC Promotes Bovine Myoblast Proliferation through the CENPF/CDK1 Axis. J AGR FOOD CHEM. 2024;72(22):12641–12654 WB ;Bovine. 38780097