– DATASHEET –

[Primary Antibody]

phospho-IRAK1 (Thr387) Rabbit pAb

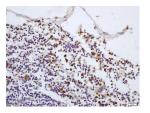


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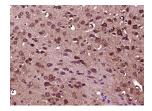
- DATASHEEL -		
Host: Rabb	it Isotype: IgG	Applications: WB (1:500-2000)
Clonality: Polyc	clonal	IHC-P (1:100-500)
GenelD: 3654	SWISS: P51617	IHC-F (1:100-500) IF (1:100-500)
Target: IRAK		Flow-Cyt (0.2µg /Test) ELISA (1:5000-10000)
Immunogen: KLH conjugated Synthesised phosphopeptide derived from human IRAK1 around the phosphorylation site of Thr387: RG(p-T)LA.		
Purification: affinity purified by Protein A		
Concentration: 1mg/	'ml	,,
Glyce Ship	/ TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% erol. ped at 4°C. Store at -20°C for one year. Avoid repeated e/thaw cycles.	Predicted MW.: ^{78 kDa} Subcellular Location: ^{Cytoplasm} ,Nucleus
Background: This point of assocration of the term of term	1, on. of cript	
- VALIDATION IN	IAGES	



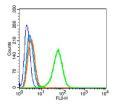
Sample: Lane1: Liver (Mouse) Cell Lysate at 30 ug Lane2: Colon carcinoma (Human) Lysate at 30 ug Primary: Anti-Phospho-IRAK1 (Thr387)(bs-3194R) at 1:200 dilution; Secondary: HRP conjugated Goat Anti-Rabbit IgG(bs-0295G-HRP) at 1: 3000 dilution; Predicted band size : 70kD Observed band size : 70kD



Tissue/cell: mouse spleen 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-Phospho-IRAK1 (Thr387) Polyclonal Antibody, Unconjugated(bs-3194R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining



Paraformaldehyde-fixed, paraffin embedded (Mouse brain); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (Phospho-IRAK1 (Thr387)) Polyclonal Antibody, Unconjugated (bs-3194R) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



Blank control(blue): RSC96 cells(fixed with 2% paraformaldehyde (10 min) , then permeabilized with 90% ice-cold methanol for 30 min on ice). Primary Antibody: Rabbit Anti-Phospho-IRAK1 (Thr387) antibody(bs-3194R), Dilution: 0.2µg in 100 µL 1X PBS containing 0.5% BSA; Isotype

Control Antibody: Rabbit IgG(orange) ,used under the same conditions); Secondary Antibody: Goat anti-rabbit IgG-PE(white blue), Dilution: 1:200 in 1 X PBS containing 0.5% BSA.

- SELECTED CITATIONS -

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- [IF=6.8] Mengyuan Li. et al. NLRP12 attenuates ozone-induced pulmonary inflammation by regulating canonical NF-κB Pathway. ECOTOX ENVIRON SAFE. 2023 Sep;262:115275 WB ;Mouse. 37531929
- [IF=6.43] Wen et al. Positive Feedback Regulation between Transglutaminase 2 and Toll-Like Receptor 4 Signaling in Hepatic Stellate Cells Correlates with Liver Fibrosis PostSchistosoma japonicumInfection. (2018) Front.Immunol. 8:1808 IHC ;Mouse. 29321784
- [IF=3.8] Wei Wang. et al. Sulforaphane inhibits the migration and invasion of BPDE-induced lung adenocarcinoma cells by regulating NLRP12. TOXICOL APPL PHARM. 2024 Mar;:116916 WB ;Human. 38537874