## [ Primary Antibody ]

## Caspase-1 P10 Rabbit pAb



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– DATASHEET –––––		400-901-9800
Host: Rabbit	Isotype: IgG	Applications: IHC-P (1:100-500)
Clonality: Polyclonal		<b>IHC-F</b> (1:100-500)
GenelD: 834	SWISS: P29466	Flow-Cyt (lug/Test)
Target: Caspase-1 P.	10	Reactivity: Human Mouse Rat
Immunogen: KLH conjuga 320-404/404	ted synthetic peptide derived from human Caspase-1:	neuerrey maintail, mouse, rat
Purification: affinity purif	ed by Protein A	
Concentration: 1mg/ml		Predicted MW.: <sup>10/45</sup> kDa
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 freeze/thaw cycles.		Subcellular Location: Cytoplasm
Background: This gene en aspartic acic caspases pla apoptosis. C proteolytic p subunits, lar This gene wa activate the in the proces healing. This may function gene in mou disease. Alte variants enc	codes a protein which is a member of the cysteine- protease (caspase) family. Sequential activation of ys a central role in the execution-phase of cell aspases exist as inactive proenzymes which undergo rocessing at conserved aspartic residues to produce 2 ge and small, that dimerize to form the active enzyme. Is identified by its ability to proteolytically cleave and inactive precursor of interleukin-1, a cytokine involved sees such as inflammation, septic shock, and wound gene has been shown to induce cell apoptosis and in various developmental stages. Studies of a similar se suggest a role in the pathogenesis of Huntington rnative splicing of this gene results in five transcript oding distinct isoforms. [provided by RefSeq].	

## VALIDATION IMAGES



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 (Caspase-1 P10) Polyclonal Antibody, Unconjugated (bs-0169R) at 1:500 overnight at 4°C, followed by a conjugated secondary (sp-0023) for 20 minutes and DAB staining.



Paraformaldehyde-fixed, paraffin embedded (Rat 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 (Caspase-1 P10) Polyclonal Antibody, Unconjugated (bs-0169R) at 1:500 overnight at 4°C, followed by a conjugated secondary (sp-0023) for 20 minutes and DAB staining.



Blank control:HL-60. Primary Antibody (green line): Rabbit Anti-Caspase-1 P10 antibody (bs-0169R) Dilution: 1µg /10^6 cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody : Goat anti-rabbit IgG-AF488 Dilution: 1µg /test. Protocol The cells were fixed with 4% PFA (10min at room temperature) and then permeabilized with 0.1% PBST for 20 min at room temperature. The cells were then incubated in 5%BSA to block non-specific protein-protein interactions for 30 min at room temperature .Cells stained with Primary Antibody for 30 min at room temperature. The secondary antibody used for 40 min at room temperature. Acquisition of 20,000 events was performed.

## - SELECTED CITATIONS -

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- [IF=12.779] Marques-da-Silva, Camila. et al. AIM2 sensors mediate immunity to <i>Plasmodium</i> infection in hepatocytes. P NATL ACAD SCI USA. 2023 Jan;120(2):e2210181120 WB ;Human. 36595704
- [IF=10.7] Lin Gan. et al. Chondroitin sulfate modulates oxidative stress and inflammation in the substantia nigra via gut microbiota regulation: Mechanistic insights into Parkinson's disease treatment. CARBOHYD POLYM. 2024 Oct;:122874 WB ;MOUSE. 10.1016/j.carbpol.2024.122874
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