## bs-0052R

## [ Primary Antibody ]

# Caspase 8 Rabbit pAb



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Applications: WB (1:500-2000) IHC-P (1:100-500) IHC-F (1:100-500) IF (1:50-200) Flow-Cyt (1µg /Test)

Reactivity: Human, Rat (predicted: Mouse, Pig, Cow, Dog, Horse)

Predicted MW.: 12/55 kDa

Subcellular Location: Cytoplasm

Host: Rabbit

- DATASHEET -

Clonality: Polyclonal GenelD: 54474

SWISS: Q9JHX4

Isotype: IgG

Target: Caspase 8

**Immunogen:** KLH conjugated synthetic peptide derived from rat Caspase-8 subunit p10: 411-482/482.

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

freeze/thaw cycles. Background: Caspases are cysteine proteases, expressed as inactive precursors,

that mediate apoptosis by proteolysis of specific substrates. Caspases have the ability to cleave after aspartic acid residues. There are two classes of caspases involved in apoptosis; initiators (activation by receptor cluster) and effectors (activation by mitochondrial permeability transition). Proapoptotic signals autocatalytically activate initiator caspases, such as Caspase 8 and Caspase 9. Activated initiator caspases then process effector caspases, such as Caspase 3 and Caspase 7, which in turn cause cell collapse.

## - VALIDATION IMAGES



Sample: Jurkat(Human) Cell Lysate at 30 ug K562(Human) Cell Lysate at 30 ug Primary: Anti-caspase-8 subunit p18 (bs-0052R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 55/18 kD Observed band size: 57 kD



Paraformaldehyde-fixed, paraffin embedded (rat liver); 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 8) Polyclonal Antibody, Unconjugated (bs-0052R) at 1:500 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



Blank control (blue line): U251 (blue). Primary Antibody (green line): Rabbit Anti-caspase-8 antibody (bs-0052R) Dilution: 1µg /10^6 cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody (white blue line): Goat anti-rabbit IgG-PE Dilution: 1µg /test. Protocol The cells were fixed with 70% ethanol overnight at 4°C and then permeabilized with 0.1% PBS-Tween for 20 min at room temperature. Cells stained with Primary Antibody for 30 min at room temperature. The cells were then incubated in 1 X PBS/2%BSA/10% goat serum to block non-specific protein-protein interactions followed by the antibody for 15 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=6.575] Anvita Gupta. et al. Androgen Receptor Activation Induces Senescence in Thyroid Cancer Cells. CANCERS. 2023 Jan;15(8):2198 IF ;Human. 37190127
- [IF=5.23] Zhao, Yong, et al. "Hydrogen Sulfide and/or Ammonia Reduces Spermatozoa Motility through AMPK/AKT Related Pathways." Scientific Reports 6 (2016): 37884. WB ;="Pig". 27883089
- [IF=5.4] Yueqi Yang. et al. A Compared Study of Eicosapentaenoic Acid and Docosahexaenoic Acid in Improving Seizure-Induced Cognitive Deficiency in a Pentylenetetrazol-Kindling Young Mice Model. MAR DRUGS. 2023 Sep;21(9):464 WB ;MOUSE. 10.3390/md21090464
- [IF=3.53] Fang C, Zhang J, Qi D, Fan X, Luo J, et al. (2014) Evodiamine Induces G2/M Arrest and Apoptosis via Mitochondrial and Endoplasmic Reticulum Pathways in H446 and H1688 Human Small-Cell Lung Cancer Cells. PLoS ONE 9(12): e115204. WB ;="Human". 25506932