### bsm-33035M

## [ Primary Antibody ]

# PCNA Mouse mAb, Nuclear Loading Control



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- DATASHEET -Host: Mouse Isotype: IgG Applications: WB (1:500-5000) IHC-P (1:100-500) Clonality: Monoclonal CloneNo.: 10E9 **IHC-F** (1:100-500) GenelD: 5111 SWISS: P12004 IF (1:100-500) Flow-Cyt (1:50) Target: PCNA Purification: affinity purified by Protein G Reactivity: Human, Mouse, Rat Concentration: 1mg/ml Storage: Size : 50ul/100ul/500ul Predicted 29 kDa 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol. MW.: Size : 200ug (PBS only) 0.01M PBS Subcellular Location: Nucleus Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles. Background: Proliferating cell nuclear antigen (PCNA) is a 28kDa nuclear protein associated with the cell cycle, a nuclear protein vital for cellular DNA synthesis. Proliferating cell nuclear antigen was originally identified by immunofluorescence as a nuclear protein whose appearance correlated with the proliferate state of the cell. PCNA is required for replication of DNA in vitro and has been identified as the auxiliary protein (cofactor) for DNA polymerase delta. The anti-PCNA antibodies react with the nuclei of proliferating cells. PCNA is essential for cellular DNA synthesis and is also required for the in vitro replication of simian virus 40 (SV40) DNA where it acts to coordinate leading and lagging strand synthesis at the replication fork. The PCNA protein may fulfil several separate roles in the cell

#### — VALIDATION IMAGES



25 ug total protein per lane of various lysates (see on figure) probed with PCNA monoclonal antibody, unconjugated (bsm-33035M) at 1:5000 dilution and 4°C overnight incubation. Followed by conjugated secondary antibody incubation at r.t. for 60 min.



nucleus associated with changes in its antigenic structure.

Paraformaldehyde-fixed, paraffin embedded Human Breast Cancer Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15 min; Antibody incubation with PCNA(Nuclear Loading Control) Monoclonal Antibody, Unconjugated(bsm-33035M) at 1:200 overnight at 4°C, followed by conjugation to the SP Kit (Mouse, sp-0024) and DAB (C-0010) staining.



Paraformaldehyde-fixed, paraffin embedded Human Duodenum; Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15 min; Antibody incubation with PCNA(Nuclear Loading Control) Monoclonal Antibody, Unconjugated(bsm-33035M) at 1:200 overnight at 4°C, followed by conjugation to the SP Kit (Mouse, sp-0024) and DAB (C-0010) staining.



Blank control:Jurkat. Primary Antibody (green line): Mouse Anti-PCNA antibody (bsm-33035M) Dilution: 1:50; Secondary Antibody : Goat antimouse IgG-FITC Dilution: 0.5ug/Test. Protocol The cells were fixed with 4% PFA (10min at room temperature) and then permeabilized with 90% ice-cold methanol for 20 min at -20°C. The cells were then incubated in 5%BSA to block nonspecific 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=3.99] Zhang, Zhongwei, et al. "Generation of glucagon like peptide 2 expressing Saccharomyces cerevisiae and its improvement of the intestinal health of weaned rats." Microbial Biotechnology (2016). IHC ;="Rat". 27641625
- [IF=3.3] Zhang et al. Hydrogen-rich water protects against acetaminophen-induced hepatotoxicity in mice. (2015) World.J.Gastroenterol. 21:4195-209 IHC ;Mouse. 25892869
- [IF=2.6] Xu Hongyan. et al. The New Roles of traf6 Gene Involved in the Development of Zebrafish Liver and Gonads. MAR BIOTECHNOL. 2024 Jun;:1-14 IF ;Zebrafish. 38861111
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