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AM10035FC-N Monoclonal

Monoclonal Antibody to PCNA FITC

Alternate names:	Cyclin, Proliferating Cell Nuclear Antigen
Quantity:	0.1 mg
Concentration:	0.1 mg/ml
Background:	Proliferating Cell Nuclear Antigen, commonly known as PCNA, is a protein that acts as a processivity factor for DNA polymerase delta in eukaryotic cells. The protein encoded by this gene is found in the nucleus and is a cofactor of DNA polymerase delta. The encoded protein acts as a homotrimer and helps increase the processivity of leading strand synthesis during DNA replication. In response to DNA damage, this protein is ubiquitinated and is involved in the RAD6-dependent DNA repair pathway. Two transcript variants encoding the same protein have been found for this gene. Pseudogenes of this gene have been described on chromosome 4 and on the X chromosome. PCNA was originally identified as an antigen that is expressed in the nuclei of cells during the DNA synthesis phase of the cell cycle It is increased during late G1 phase and S phase of the cell cycle and declines during G2 and M phases.
Uniprot ID:	<u>P12004</u>
NCBI:	<u>9606</u>
GenelD:	<u>5111</u>
Host / Isotype:	Mouse / IgG2a
Clone:	PC10
Immunogen:	Rat PCNA made in the protein A expression vector pR1T2T.
Format:	State: Liquid purified IgG fraction Purification: Affinity Chromatography on Protein G Buffer System: PBS Preservatives: 0.09% Sodium Azide Stabilizers: 1% BSA Label: FITC – Fluorescein Isothiocyanate Isomer 1
Applications:	Flow Cytometry: Use 10 μ l of Neat-1/10 of dilutued antibnody to label 10 ⁶ cells in 100 μ l. Cell permeabilisation is required for this application. Other applications not tested. Optimal dilutions are dependent on conditions and should be determined by the user.

For research and in vitro use only. Not for diagnostic or therapeutic work. Material Safety Datasheets are available at www.acris-antibodies.com or on request.

	AM10035FC-N: Monoclonal Antibody to PCNA FITC
Specificity:	This antibody recognises the proliferating cell nuclear antigen (PCNA), a nuclear protein vital for cellular DNA synthesis. PCNA is highly conserved between mammalian species and other vertebrates. Mouse anti Human PCNA, clone <i>PC10</i> has been used for the detection of PCNA in a number of species including Rat (Elsässer et al. 1994), Mouse (Park et al. 2008), Chicken (Franz- Odendaal 2008) and Abalone (Harris et al. 2006). Species: Human, Insects, Vertebrates, Rat, Ferret, Chicken, Rabbit, Xenopus, Rhesus Monkey, Hamster, Atlantic Salmon, Mouse, Horse, Sheep, Dog, Cat, Cynomolgus monkey.
	Other species not tested.
Storage:	Store undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer. This product is photosensitive and should be protected from light. Avoid repeated freezing and thawing. Shelf life: one year from despatch.
General Readings:	 Mathews MB, Bernstein RM, Franza BR, Garrels JI. Identity of the proliferating cell nuclear antigen and cyclin. Nature. 1984 May 24-30;309(5966):374-6. PubMed PMID: 6145097. Garcia RL, Coltrera MD, Gown AM. Analysis of proliferative grade using anti- PCNA/cyclin monoclonal antibodies in fixed, embedded tissues. Comparison with flow cytometric analysis. Am J Pathol. 1989 Apr;134(4):733-9. PubMed PMID: 2565087. Landberg G, Tan EM, Roos G. Flow cytometric multiparameter analysis of proliferating cell nuclear antigen/cyclin and Ki-67 antigen: a new view of the cell cycle. Exp Cell Res. 1990 Mar;187(1):111-8. PubMed PMID: 1967582. Wilson, G.D. et al. (1992) Flow cytometric characterisation of proliferating cell nuclear antigen using monoclonal antibody PC10. Eur. J. Cancer 28A: 2010-2017. Elsässer HP, Biederbick A, Kern HF. Growth of rat pancreatic acinar cells quantitated with a monoclonal antibody against the proliferating cell nuclear antigen. Cell Tissue Res. 1994 Jun;276(3):603-9. PubMed PMID: 7914831. Prosperi E, Stivala LA, Sala E, Scovassi Al, Bianchi L. Proliferating cell nuclear antigen complex formation induced by ultraviolet irradiation in human quiescent fibroblasts as detected by immunostaining and flow cytometry. Exp Cell Res. 1993 Apr;205(2):320-5. PubMed PMID: 8097724. Harris, L. et al. (2006) Characterisation of cell types in abalone (Haliotis spp.) tissues using immunohistochemical techniques Aquaculture 261: 1413-21 8. Buggins AG, Milojkovic D, Arno MJ, Lea NC, Mufti GJ, Thomas NS, et al. Microenvironment produced by acute myeloid leukemia cells prevents T cell activation and proliferation by inhibition of NF-kappaB, c-Myc, and pRb pathways. J Immunol. 2001 Nov 15;167(10):6021-30. PubMed PMID: 11698483. Kapitonova MY, Kuznetsov SL, Khlebnikov VV, Zagrebin VL, Morozova ZCh, Degtyar YV. Immunohistochemical characteristics of the hypophysis in normal conditions and chronic stress. Neurosci Behav P

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Pictures: Staining of KM-H2 cells (permeabilised) with MOUSE ANTI PCNA:FITC (SM1421F)



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