abcam

Product datasheet

Anti-PCNA antibody ab18197

★★★★★ 51 Abreviews 371 References 画像数 11

製品の概要

製品名 Anti-PCNA antibody

製品の詳細 Rabbit polyclonal to PCNA

由来種 Rabbit

アプリケーション 適用あり: WB, ICC/IF, Flow Cyt (Intra), IHC-P

種交差性 交差種: Mouse, Rat, Human, Common marmoset

交差が予測される動物種: Sheep, Goat, Cow, Dog, Xenopus laevis, Monkey, Zebrafish 🔷

免疫原 Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

製品の特性

特記事項

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -

80°C. Avoid freeze / thaw cycle.

バッファー pH: 7.40

Preservative: 0.02% Sodium azide

Constituent: PBS

Batches of this product that have a concentration < 1mg/ml may have BSA added as a stabilising

agent. If you would like information about the formulation of a specific lot, please contact our

scientific support team who will be happy to help.

精製度 Immunogen affinity purified

ポリ/モノ ポリクローナル

アイソタイプ lgG

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The Abpromise guarantee <u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab18197の使用に適用されますアプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
WB	★★★★☆ (10)	Use a concentration of 1 µg/ml. Detects a band of approximately 29 kDa (predicted molecular weight: 29 kDa).
ICC/IF	★★★★ <u>(13)</u>	Use a concentration of 1 µg/ml.
Flow Cyt (Intra)		Use 0.05µg for 10 ⁶ cells.
IHC-P	★★★★★ (22)	Use at an assay dependent concentration.

ターゲット情報

機能 This protein is an auxiliary protein of DNA polymerase delta and is involved in the control of

eukaryotic DNA replication by increasing the polymerase's processibility during elongation of the

leading strand. Induces a robust stimulatory effect on the 3'-5' exonuclease and 3'-

phosphodiesterase, but not apurinic-apyrimidinic (AP) endonuclease, APEX2 activities. Has to

be loaded onto DNA in order to be able to stimulate APEX2.

配列類似性 Belongs to the PCNA family.

翻訳後修飾 Upon methyl methanesulfonate-induced DNA damage, mono-ubiquitinated by the UBE2B-RAD18

complex on Lys-164. This induces non-canonical polyubiquitination on Lys-164 through 'Lys-63' linkage of ubiquitin moieties by the E2 complex UBE2N-UBE2V2 and the E3 ligases, HLTF, RNF8 and SHPRH, which is required for DNA repair. 'Lys-63' polyubiquitination prevents genomic instability on DNA damage. Monoubiquitination at Lys-164 also takes place in

 $\ undamaged\ proliferating\ cells,\ and\ is\ mediated\ by\ the\ DCX(DTL)\ complex,\ leading\ to\ enhance$

 $\label{eq:pcna-dependent} \mbox{PCNA-dependent translesion DNA synthesis}.$

Acetylated in response to UV irradiation. Acetylation disrupts interaction with NUDT15 and

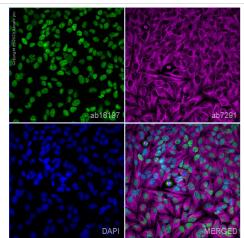
promotes degradation.

細胞内局在 Nucleus. Forms nuclear foci representing sites of ongoing DNA replication and vary in

morphology and number during S phase. Together with APEX2, is redistributed in discrete

nuclear foci in presence of oxidative DNA damaging agents.

画像



PCNA antibody (ab18197)

Immunocytochemistry/ Immunofluorescence - Anti-

250 kDa -150 kDa -100 kDa -75 kDa -50 kDa 🕳 Copyright (c) 2012 Abcam plo 37 kDa -25 kDa — 20 kDa — 15 kDa 🕳 10 kDa -

Western blot - Anti-PCNA antibody (ab18197)

ab18197 staining PCNA in HeLa cells. The cells were fixed with 100% methanol (5 min), permeabilized with 0.1% PBS-Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1%PBS-Tween for 1h. The cells were then incubated overnight at 4°C with ab18197 at 0.1µg/ml and ab7291, Mouse monoclonal [DM1A] to alpha Tubulin - Loading Control. Cells were then incubated with **ab150081**, Goat polyclonal Secondary Antibody to Rabbit IgG - H&L (Alexa Fluor® 488), pre-adsorbed at 1/1000 dilution (shown in green) and ab150120, Goat polyclonal Secondary Antibody to Mouse IgG - H&L (Alexa Fluor® 594), preadsorbed at 1/1000 dilution (shown in pseudocolour magenta). Nuclear DNA was labelled with DAPI (shown in blue).

Also suitable in cells fixed with 4% paraformaldehyde (10 min). Image was acquired with a high-content analyser (Operetta CLS, Perkin Elmer) and a maximum intensity projection of confocal sections is shown.

All lanes: Anti-PCNA antibody (ab18197) at 1 µg/ml

Lane 1: HEK293 (Human embryonic kidney cell line) Whole Cell Lysate

Lane 2: NIH 3T3 (Mouse embryonic fibroblast cell line) Whole Cell Lysate

Lane 3: PC12 (Rat adrenal pheochromocytoma cell line) Whole Cell Lysate

Lane 4: HEK293 (Human embryonic kidney cell line) Whole Cell Lysate with Human PCNA peptide (ab18602) at 1 µg/ml

Lane 5: NIH 3T3 (Mouse embryonic fibroblast cell line) Whole Cell Lysate with Human PCNA peptide (ab18602) at 1 µg/ml

Lane 6: PC12 (Rat adrenal pheochromocytoma cell line) Whole Cell Lysate with Human PCNA peptide (ab18602) at 1 µg/ml

Lysates/proteins at 20 µg per lane.

Secondary

All lanes: Goat Anti-Rabbit IgG H&L (HRP) preadsorbed

(ab97080) at 1/5000 dilution

Developed using the ECL technique.

Performed under reducing conditions.

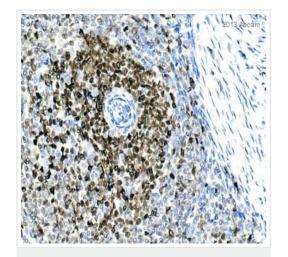
Predicted band size: 29 kDa Observed band size: 29 kDa

Additional bands at: 48 kDa, 52 kDa. We are unsure as to the

identity of these extra bands.

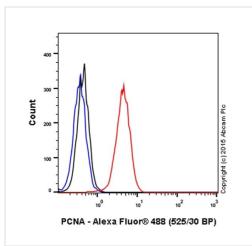
Exposure time: 1 minute

ab18197 staining PCNA in tissue sections of the goat spleen by Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections). Tissue was fixed with formaldehyde and a heat mediated antigen retrieval step was performed. Samples were then blocked with 1% B.S.A. for 10 minutes at 21°C followed by incubation with the primary antibody for 2 hours at 1/4000. A biotin-conjugated goat anti-rabbit polyclonal was used as secondary antibody at a 1/250 dilution.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-PCNA antibody (ab18197)

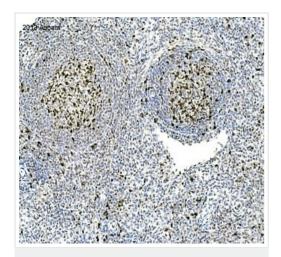
Image courtesy of Carl Hobbs, Kings College London, U.K.



Flow Cytometry (Intracellular) - Anti-PCNA antibody (ab18197)

Overlay histogram showing HeLa cells stained with ab18197 (red line). The cells were fixed with 80% methanol (5 min) and then permeabilized with 0.1% PBS-Tween 20 for 20 min. The cells were then incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (ab18197, 0.05µg/1x10⁶ cells) for 30 min at 22°C. The secondary antibody used was Alexa Fluor® 488 goat anti-rabbit lgG (H&L) (ab150081) at 1/4000 dilution for 30 min at 22°C. Isotype control antibody (black line) was rabbit lgG (polyclonal) (ab171870, 0.05µg/1x10⁶ cells) used under the same conditions. Unlabelled sample (blue line) was also used as a control.

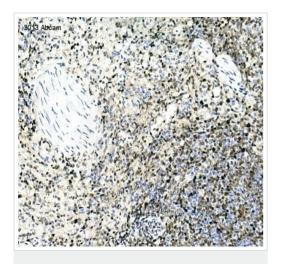
Acquisition of >5,000 events were collected using a 20mW Argon ion laser (488nm) and 525/30 bandpass filter.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-PCNA antibody (ab18197)

Image courtesy of Carl Hobbs, Kings College London, U.K.

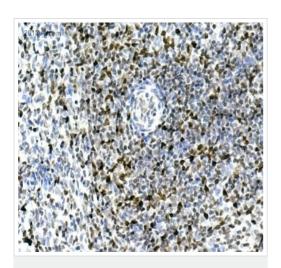
ab18197 staining PCNA in tissue sections of the marmoset spleen by Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections). Tissue was fixed with formaldehyde and a heat mediated antigen retrieval step was performed. Samples were then blocked with 1% B.S.A. for 10 minutes at 21°C followed by incubation with the primary antibody for 2 hours at 1/6000. A biotin-conjugated goat anti-rabbit polyclonal was used as secondary antibody at a 1/250 dilution.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-PCNA antibody (ab18197)

Image courtesy of Carl Hobbs, King College London, U.K.

ab18197 staining PCNA in tissue sections of the cow spleen by Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections). Tissue was fixed with formaldehyde and a heat mediated antigen retrieval step was performed. Samples were then blocked with 1% B.S.A. for 10 minutes at 21°C followed by incubation with the primary antibody for 2 hours at 1/4000. A biotin-conjugated goat anti-rabbit polyclonal was used as secondary antibody at a 1/250 dilution.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-PCNA antibody (ab18197)

Image courtesy of Carl Hobbs, King College London, U.K.

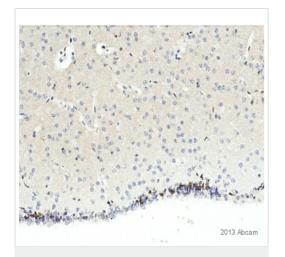
ab18197 staining PCNA in tissue sections of the sheep spleen by Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections). Tissue was fixed with formaldehyde and a heat mediated antigen retrieval step was performed. Samples were then blocked with 1% B.S.A. for 10 minutes at 21°C followed by incubation with the primary antibody for 2 hours at 1/6000. A biotin-conjugated goat anti-rabbit polyclonal was used as secondary antibody at a 1/250 dilution.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-PCNA antibody (ab18197)

Image courtesy of Carl Hobbs, Kings College London, U.K.

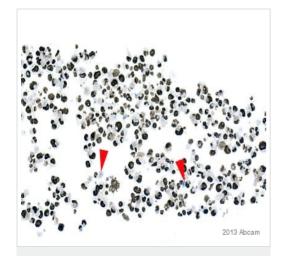
ab18197 staining PCNA in tissue sections of the rat brain by Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections). Tissue was fixed with formaldehyde and a heat mediated antigen retrieval step was performed. Samples were then blocked with 1% B.S.A. for 10 minutes at 21°C followed by incubation with the primary antibody for 2 hours at 1/10000. A biotin-conjugated goat anti-rabbit polyclonal was used as secondary antibody at a 1/250 dilution.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-PCNA antibody (ab18197)

Image courtesy of Carl Hobbs, Kings College London, U.K.

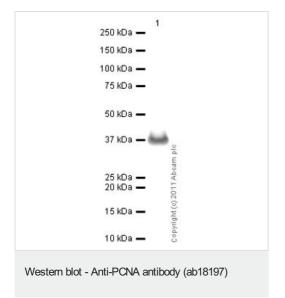
ab18197 staining PCNA in tissue sections of the mouse brain by Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections). Tissue was fixed with formaldehyde and a heat mediated antigen retrieval step was performed. Samples were then blocked with 1% B.S.A. for 10 minutes at 21°C followed by incubation with the primary antibody for 2 hours at 1/6000. A biotin-conjugated goat anti-rabbit polyclonal was used as secondary antibody at a 1/250 dilution.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-PCNA antibody (ab18197)

Image Courtesy of Carl Hobbs, Kings College London, U.K.

ab18197 staining PCNA in monkey COS cell pellet by Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections). Tissue was fixed with formaldehyde and a heat mediated antigen retrieval step was performed. Samples were then blocked with 1% B.S.A. for 10 minutes at 21°C followed by incubation with the primary antibody for 2 hours at 1/4000. A biotin-conjugated goat anti-rabbit polyclonal was used as secondary antibody at a 1/250 dilution.



Anti-PCNA antibody (ab18197) at 1 μ g/ml + Recombinant Human PCNA protein (ab85651) at 0.01 μ g

Secondary

Goat Anti-Rabbit lgG H&L (HRP) preadsorbed (ab97080) at 1/5000 dilution

Developed using the ECL technique.

Performed under reducing conditions.

Predicted band size: 29 kDa

Exposure time: 10 seconds

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