# abcam

# Product datasheet

# Anti-RPA32/RPA2 antibody [EPR2876(2)] - BSA and Azide free ab247754

יעלאעבע RabMAb

★★★★ 1 Abreviews

画像数 9

#### 製品の概要

製品名 Anti-RPA32/RPA2 antibody [EPR2876(2)] - BSA and Azide free

製品の詳細 Rabbit monoclonal [EPR2876(2)] to RPA32/RPA2 - BSA and Azide free

由来種 Rabbit

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

適用なし: №

種交差性 交差種: Rat, Human

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

ポジティブ・コントロール WB: HeLa, HUVEC, HepG2, and MCF7 cell lysates. IHC-P: Human colon and Human ovary

carcinoma tissues. ICC/IF: C6 and HeLa cells. Flow Cyt (intra): C6 and HeLa cells.

ChlC/CUT&RUN-Seq: HeLa cells.

特記事項 ab247754 is the carrier-free version of ab109084.

> Our carrier-free antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for increased conjugation efficiency.

This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cellbased assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our conjugation kits for antibody conjugates that are ready-to-use in as little as 20 minutes with <1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.

This product is compatible with the Maxpar® Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar<sup>®</sup> is a trademark of Fluidigm Canada Inc.

This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility
- Improved sensitivity and specificity
- Long-term security of supply
- Animal-free production

For more information see here.

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit

monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

Mouse: We have preliminary internal testing data to indicate this antibody may not react with this species. Please contact us for more information.

### 製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at +4°C. Do Not Freeze.

**バッファー** pH: 7.2

Constituent: PBS

キャリア・フリー はい

精製度 Affinity purified ポリ/モノ モノクローナル PDローン名 EPR2876(2)

アイソタイプ lgG

#### アプリケーション

**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ab247754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
Flow Cyt (Intra)		Use at an assay dependent concentration.
ChIC/CUT&RUN-seq		Use at an assay dependent concentration.
ICC/IF		Use at an assay dependent concentration.
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.  Antigen retrieval is recommended.
WB		Use at an assay dependent concentration. Predicted molecular weight: 29 kDa.

追加情報 Is unsuitable for IP.

#### ターゲット情報

機能

Required for DNA recombination, repair and replication. The activity of RP-A is mediated by single-stranded DNA binding and protein interactions.

Functions as component of the alternative replication protein A complex (aRPA). aRPA binds single-stranded DNA and probably plays a role in DNA repair; it does not support chromosomal

DNA replication and cell cycle progression through S-phase. In vitro, aRPA cannot promote efficient priming by DNA polymerase alpha but supports DNA polymerase delta synthesis in the presence of PCNA and replication factor C (RFC), the dual incision/excision reaction of nucleotide excision repair and RAD51-dependent strand exchange.

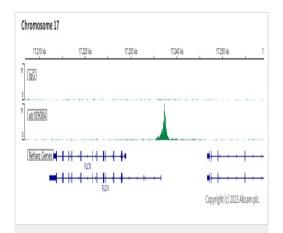
# 翻訳後修飾

Phosphorylated in a cell-cycle-dependent manner (from the S phase until mitosis). Phosphorylated by ATR upon DNA damage, which promotes its translocation to nuclear foci. Can be phosphorylated in vitro by PRKDC/DNA-PK in the presence of Ku and DNA, and by CDK1.

#### 細胞内局在

Nucleus. Nucleus > PML body. Also present in PML nuclear bodies. Redistributes to discrete nuclear foci upon DNA damage.

#### 画像

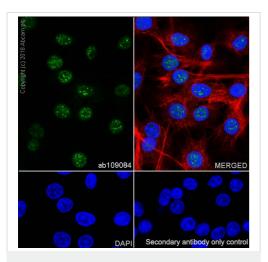


ChIC/CUT&RUN sequencing - Anti-RPA32/RPA2 antibody [EPR2876(2)] - BSA and Azide free (ab247754) ChIC/CUT&RUN was performed using a pAG-MNAse at a final concentration of 700 ng/mL, 2.5 x 10^5 HeLa (Human cervix adenocarcinoma epithelial cell line) cells and 5µg of <u>ab109084</u> [EPR2876(2)]. The resulting DNA was sequenced on the Illumina NovaSeq 6000 to a depth of 10 million reads. The negative IgG control <u>ab172730</u> is also shown.

Additional screenshots of mapped reads can be downloaded here.

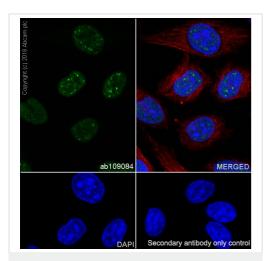
The University of Geneva owns patents relevant to ChlC (Chromatin Immuno-Cleavage) methods.

This data was developed using <u>ab109084</u>, the same antibody clone in a different buffer formulation.



Immunocytochemistry/ Immunofluorescence - Anti-RPA32/RPA2 antibody [EPR2876(2)] - BSA and Azide free (ab247754) This data was developed using <u>ab109084</u>, the same antibody clone in a different buffer formulation.

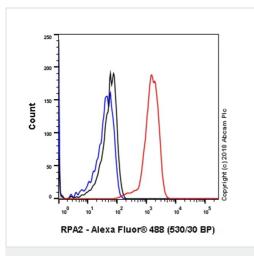
Immunocytochemistry/ Immunofluorescence analysis of C6 (rat glial tumor glial cell) cells labeling RPA32/RPA2 with purified  $\underline{ab109084}$  at 1:50 (9.34 µg/ml). Cells were fixed in 100% Methanol and permeabilized with None. Cells were counterstained with Ab195889 Anti-alpha Tubulin antibody [DM1A] - Microtubule Marker (Alexa Fluor 594) 1:200 (2.5 µg/ml). Goat anti rabbit lgG (Alexa Fluor 488,  $\underline{ab150077}$ ) was used as the secondary antibody at 1:1000 (2 µg/ml) dilution. DAPI nuclear counterstain. PBS instead of the primary antibody was used as the secondary antibody only control.



Immunocytochemistry/ Immunofluorescence - Anti-RPA32/RPA2 antibody [EPR2876(2)] - BSA and Azide free (ab247754)

This data was developed using <u>ab109084</u>, the same antibody clone in a different buffer formulation.

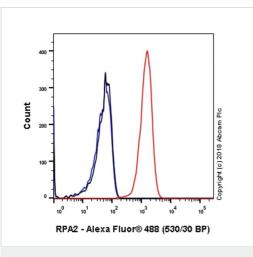
Immunocytochemistry/ Immunofluorescence analysis of HeLa (human cervix adenocarcinoma epithelial cell) cells labeling RPA32/RPA2 with purified  $\underline{ab109084}$  at 1:50 (9.34 µg/ml). Cells were fixed in 100% Methanol and permeabilized with None. Cells were counterstained with Ab195889 Anti-alpha Tubulin antibody [DM1A] - Microtubule Marker (Alexa Fluor® 594) 1:200 (2.5 µg/ml). Goat anti rabbit lgG (Alexa Fluor® 488,  $\underline{ab150077}$ ) was used as the secondary antibody at 1:1000 (2 µg/ml) dilution. DAPI nuclear counterstain. PBS instead of the primary antibody was used as the secondary antibody only control.



Flow Cytometry (Intracellular) - Anti-RPA32/RPA2 antibody [EPR2876(2)] - BSA and Azide free (ab247754)

This data was developed using <u>ab109084</u>, the same antibody clone in a different buffer formulation.

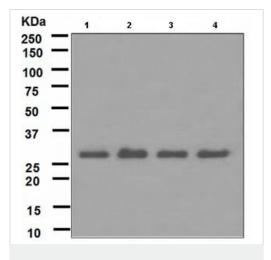
Intracellular Flow Cytometry analysis of C6 (rat glial tumor glial cell) cells labeling RPA32/RPA2 with purified  $\underline{ab109084}$  at 1/460 dilution (1.01 µg/ml) - Red. Cells were fixed with 4% paraformaldehyde . A Goat anti rabbit lgG (Alexa Fluor® 488,  $\underline{ab150077}$ ) secondary antibody was used at 1/2000 dilution. Isotype control - Rabbit monoclonal lgG ( $\underline{ab172730}$ ) - Black. Unlabeled control - Blue. Untreated cells - Green



Flow Cytometry (Intracellular) - Anti-RPA32/RPA2 antibody [EPR2876(2)] - BSA and Azide free (ab247754)

This data was developed using <u>ab109084</u>, the same antibody clone in a different buffer formulation.

Intracellular Flow Cytometry analysis of HeLa (human cervix adenocarcinoma epithelial cell) cells labeling RPA32/RPA2 with purified  $\underline{ab109084}$  at 1/460 dilution (1.01 µg/ml) - Red. Cells were fixed with 4% paraformaldehyde . A Goat anti rabbit lgG (Alexa Fluor 488,  $\underline{ab150077}$ ) secondary antibody was used at 1/2000 dilution. Isotype control - Rabbit monoclonal lgG ( $\underline{ab172730}$ ) - Black. Unlabeled control - Blue. Untreated cells - Green



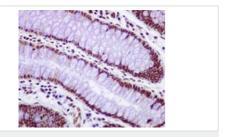
Western blot - Anti-RPA32/RPA2 antibody
[EPR2876(2)] - BSA and Azide free (ab247754)

**All lanes :** Anti-RPA32/RPA2 antibody [EPR2876(2)] (<u>ab109084</u>) at 1/1000 dilution

Lane 1 : HeLa cell lysate
Lane 2 : HUVEC cell lysate
Lane 3 : HepG2 cell lysate
Lane 4 : MCF7 cell lysate

Lysates/proteins at 10 µg per lane.

Predicted band size: 29 kDa

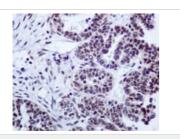


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-RPA32/RPA2 antibody

[EPR2876(2)] - BSA and Azide free (ab247754)

This data was developed using <u>ab109084</u>, the same antibody clone in a different buffer formulation.

This data was developed using <u>ab109084</u>, the same antibody clone in a different buffer formulation.<u>ab109084</u> at 1/250 dilution staining RPA32/RPA2 in Human colon by Immunohistochemistry, Paraffin-embedded tissue. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-RPA32/RPA2 antibody
[EPR2876(2)] - BSA and Azide free (ab247754)

This data was developed using <u>ab109084</u>, the same antibody clone in a different buffer formulation.<u>ab109084</u> at 1/250 dilution staining RPA32/RPA2 in Human ovary carcinoma by Immunohistochemistry, Paraffin-embedded tissue. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



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