# abcam

# Product datasheet

# Anti-Cyclin A2 antibody [EPR17351] ab181591



★★★★★ 3 Abreviews 79 References 12 Images

#### Overview

**Product name** Anti-Cyclin A2 antibody [EPR17351]

**Description** Rabbit monoclonal [EPR17351] to Cyclin A2

**Host species** Rabbit

**Tested applications** Suitable for: ICC/IF, IHC-P, WB Species reactivity Reacts with: Mouse, Rat, Human

**Immunogen** Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.

Positive control WB: Jurkat, C6, RAW 264.7 and PC-12 whole cell lysates; HeLa Untreated, asynchronous cells,

> HeLa G1/S arrested cells (thymidine treatment) and HeLa G2/M arrested cells (sequential thymidine and nocodazole treatments) cell lysates; Human tonsil and fetal kidney lysates; Rat spleen lysate. IHC-P: Human tonsil, Human cervix carcinoma, rat colon and mouse endometrium

tissues. ICC/IF: HeLa cells.

General notes 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.

#### **Properties**

**Form** Liquid

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long

term. Avoid freeze / thaw cycle.

Storage buffer Preservative: 0.01% Sodium azide

Constituents: 59% PBS, 40% Glycerol, 0.05% BSA

**Purity** Protein A purified

Clonality Monoclonal Clone number EPR17351

#### **Applications**

The Abpromise guarantee Our Abpromise guarantee covers the use of ab181591 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
ICC/IF		1/500.
IHC-P		1/500. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
WB	****(1)	1/2000. Detects a band of approximately 50 kDa (predicted molecular weight: 47 kDa).

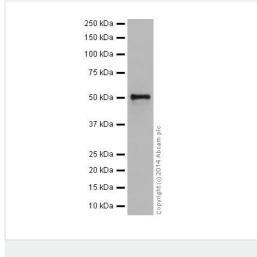
# **Target**

**Function** Essential for the control of the cell cycle at the G1/S (start) and the G2/M (mitosis) transitions.

**Sequence similarities**Belongs to the cyclin family. Cyclin AB subfamily.

**Developmental stage**Accumulates steadily during G2 and is abruptly destroyed at mitosis. **Cellular localization**Nucleus. Cytoplasm. Cytoplasmic when associated with SCAPER.

#### **Images**



Western blot - Anti-Cyclin A2 antibody [EPR17351] (ab181591)

Anti-Cyclin A2 antibody [EPR17351] (ab181591) at 1/20000 dilution + Jurkat (Human T cell leukemia cell line from peripheral blood) whole cell lysate at 20 µg

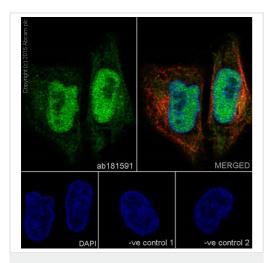
#### **Secondary**

Goat Anti-Rabbit IgG, (H+L),Peroxidase conjugated at 1/1000 dilution

**Predicted band size:** 47 kDa **Observed band size:** 50 kDa

Exposure time: 10 seconds

Blocking/Dilution buffer: 5% NFDM/TBST.



Immunocytochemistry/ Immunofluorescence - Anti-Cyclin A2 antibody [EPR17351] (ab181591)

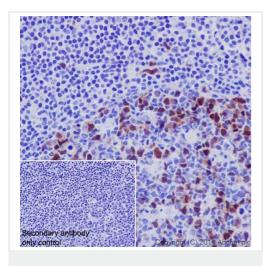
Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized HeLa (Human epithelial cell line from cervix adenocarcinoma) cells labeling Cyclin A2 with ab181591 at 1/500 dilution, followed by Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) (ab150077) secondary antibody at 1/500 dilution (green). Confocal image showing nuclear and weakly cytoplasmic staining on HeLa cells. The nuclear counter stain is DAPI (blue).

Tubulin is detected with Anti-alpha Tubulin antibody [EPR17351]-Loading Control (<u>ab7291</u>) at 1/1000 dilution, followed by Goat Anti-Mouse IgG H&L (Alexa Fluor<sup>®</sup> 594) preadsorbed (<u>ab150120</u>) at 1/1000 dilution (red).

The negative controls are as follows:

-ve control 1: ab181591 at 1/500 dilution, followed by Goat Anti-Mouse IgG H&L (Alexa Fluor<sup>®</sup> 594) preadsorbed (**ab150120**) at 1/500 dilution.

-ve control 2: Anti-alpha Tubulin antibody [EPR17351]- Loading Control (ab7291) at 1/1000 dilution, followed by Goat Anti-Rabbit lgG H&L (Alexa Fluor<sup>®</sup> 488) (ab150077) secondary antibody at 1/500 dilution.

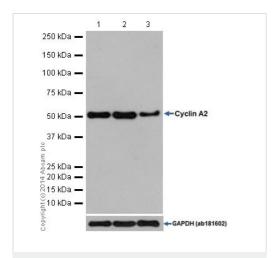


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Cyclin A2 antibody
[EPR17351] (ab181591)

Immunohistochemical analysis of paraffin-embedded Human tonsil tissue labeling Cyclin A2 with ab181591 at 1/500 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Nuclear and weak cytoplasmic staining on germinal center cells of Human tonsil tissue is observed. Counter stained with Hematoxylin. Secondary antibody only control: Used PBS instead of primary

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) (ab97051) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Western blot - Anti-Cyclin A2 antibody [EPR17351] (ab181591)

**All lanes :** Anti-Cyclin A2 antibody [EPR17351] (ab181591) at 1/20000 dilution

Lane 1 : HeLa (Human epithelial cell line from cervix adenocarcinoma) lysate; Untreated, asynchronous cells (ab136811)

Lane 2 : HeLa (Human epithelial cell line from cervix adenocarcinoma) lysate; G1/S arrested cells (thymidine treatment) (ab136811)

Lane 3: HeLa (Human epithelial cell line from cervix adenocarcinoma) lysate; G2/M arrested cells (sequential thymidine and nocodazole treatments) (ab136811)

Lysates/proteins at 1 µg per lane.

#### **Secondary**

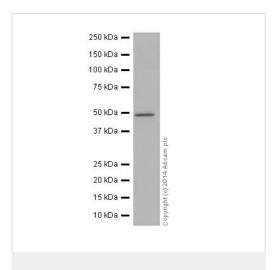
**All lanes :** Goat Anti-Rabbit lgG, (H+L),Peroxidase conjugated or Anti-Rabbit lgG (HRP), specific to the non-reduced form of lgG at 1/1000 dilution

**Predicted band size:** 47 kDa **Observed band size:** 50 kDa

Exposure time: 30 seconds

Blocking/Dilution buffer: 5% NFDM/TBST.

Cyclin A2 is down regulated at the G2/M phase.



Western blot - Anti-Cyclin A2 antibody [EPR17351] (ab181591)



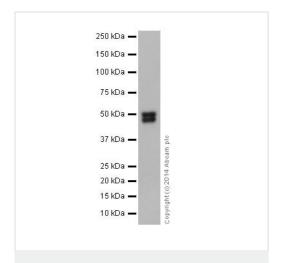
#### **Secondary**

Anti-Rabbit lgG (HRP), specific to the non-reduced form of lgG at 1/1000 dilution

**Predicted band size:** 47 kDa **Observed band size:** 50 kDa

Exposure time: 1 minute

Blocking/Dilution buffer: 5% NFDM/TBST.



Western blot - Anti-Cyclin A2 antibody [EPR17351] (ab181591)

Anti-Cyclin A2 antibody [EPR17351] (ab181591) at 1/2000 dilution + Human fetal kidney lysate at 10 µg

#### **Secondary**

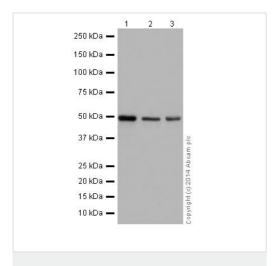
Anti-Rabbit lgG (HRP), specific to the non-reduced form of lgG at 1/1000 dilution

**Predicted band size:** 47 kDa **Observed band size:** 50 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFDM/TBST.

The smaller band is due to a tissue-specific splice variant, PMID 22745723.



Western blot - Anti-Cyclin A2 antibody [EPR17351] (ab181591)

**All lanes :** Anti-Cyclin A2 antibody [EPR17351] (ab181591) at 1/2000 dilution

Lane 1: C6 (Rat glial tumor cell line) whole cell lysate

**Lane 2**: RAW 264.7 (Mouse macrophage cell line transformed with Abelson murine leukemia virus) whole cell lysate

Lane 3 : PC-12 (Rat adrenal gland pheochromocytoma cell line) whole cell lysate

Lysates/proteins at 10 µg per lane.

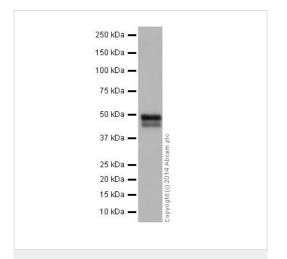
### Secondary

**All lanes :** Goat Anti-Rabbit IgG, (H+L),Peroxidase conjugated at 1/1000 dilution

**Predicted band size:** 47 kDa **Observed band size:** 50 kDa

Exposure time: 3 seconds

Blocking/Dilution buffer: 5% NFDM/TBST.



Western blot - Anti-Cyclin A2 antibody [EPR17351] (ab181591)

Anti-Cyclin A2 antibody [EPR17351] (ab181591) at 1/2000 dilution + Rat spleen lysate at 10  $\mu g$ 

# Secondary

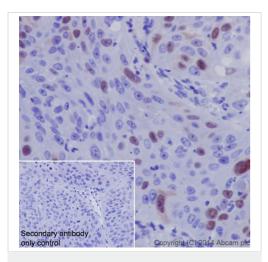
Goat Anti-Rabbit lgG, (H+L),Peroxidase conjugated at 1/1000 dilution

**Predicted band size:** 47 kDa **Observed band size:** 50 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFDM/TBST.

The smaller band is due to a tissue-specific splice variant, PMID 22745723.

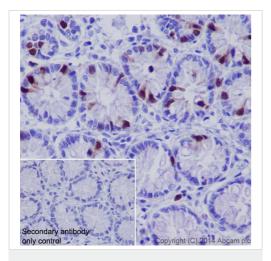


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Cyclin A2 antibody
[EPR17351] (ab181591)

Immunohistochemical analysis of paraffin-embedded Human cervix carcinoma tissue labeling Cyclin A2 with ab181591 at 1/500 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Nuclear and weak cytoplasmic staining on some tumor cells in Human cervix carcinoma tissue is observed. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) (ab97051) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

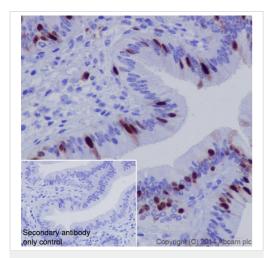


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Cyclin A2 antibody
[EPR17351] (ab181591)

Immunohistochemical analysis of paraffin-embedded rat colon tissue labeling Cyclin A2 with ab181591 at 1/500 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Nuclear and weak cytoplasmic staining on a proportion of epithelial cells in rat colon tissue is observed. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) (ab97051) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

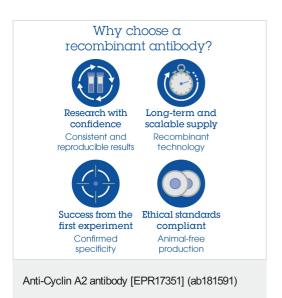


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Cyclin A2 antibody
[EPR17351] (ab181591)

Immunohistochemical analysis of paraffin-embedded mouse endometrium tissue labeling Cyclin A2 with ab181591 at 1/500 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Nuclear and weak cytoplasmic staining on a proportion of epithelial cells in mouse endometrial tissue is observed. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) (ab97051) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



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