# abcam

# Product datasheet

# Anti-Aurora A antibody [EP1008Y] ab52973

Recombinant RabMAb

★★★★★ 2 Abreviews 12 References

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#### Overview

**Product name** Anti-Aurora A antibody [EP1008Y]

**Description** Rabbit monoclonal [EP1008Y] to Aurora A

**Host species** Rabbit

**Tested applications** Suitable for: WB, IP, IHC-P

Unsuitable for: Flow Cyt or ICC/IF

Species reactivity Reacts with: Human

**Immunogen** Synthetic peptide within Human Aurora A aa 350-450 (C terminal). The exact sequence is

proprietary.

Positive control HepG2 nuclear cell lysate Human cervical carcinoma tissue IP: HepG2 whole cell lysate.

**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  $\mathsf{RabMAb}^{\texttt{®}}$  technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

Mouse, Rat: We have preliminary internal testing data to indicate this antibody may not react with

these species. Please contact us for more information.

#### **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.

Avoid freeze / thaw cycle.

Storage buffer pH: 7.20

Preservative: 0.05% Sodium azide

Constituents: 0.1% BSA, 40% Glycerol (glycerin, glycerine), 9.85% Tris glycine, 50% Tissue

culture supernatant

**Purity** Protein A purified

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Clonality Monoclonal
Clone number EP1008Y
Isotype IgG

### **Applications**

#### The Abpromise guarantee

Our Abpromise guarantee covers the use of ab52973 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
WB	<b>★★★★☆ (1)</b>	1/50000. Detects a band of approximately 46 kDa (predicted molecular weight: 46 kDa).
IP		1/50.
IHC-P		1/100 - 1/250. Perform heat mediated antigen retrieval before commencing with IHC staining protocol.

**Application notes** 

Is unsuitable for Flow Cyt or ICC/IF.

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Ьu	nction

Contributes to the regulation of cell cycle progression. Required for normal mitosis. Associates with the centrosome and the spindle microtubules during mitosis and functions in centrosome maturation, spindle assembly, maintenance of spindle bipolarity, centrosome separation and mitotic checkpoint control. Phosphorylates numerous target proteins, including ARHGEF2, BRCA1, KIF2A, NDEL1, PARD3, PLK1 and BORA. Regulates KIF2A tubulin depolymerase activity (By similarity). Required for normal axon formation. Plays a role in microtubule remodeling during neurite extension. Important for microtubule formation and/or stabilization.

### Tissue specificity

Highly expressed in testis and weakly in skeletal muscle, thymus and spleen. Also highly expressed in colon, ovarian, prostate, neuroblastoma, breast and cervical cancer cell lines.

### Sequence similarities

 $\label{thm:continuous} \textbf{Belongs to the protein kinase superfamily. Ser/Thr protein kinase family. Aurora subfamily.}$ 

Contains 1 protein kinase domain.

# Post-translational modifications

Activated by phosphorylation at Thr-288; this brings about a change in the conformation of the activation segment. Phosphorylation at Thr-288 varies during the cell cycle and is highest during M phase. Autophosphorylated at Thr-288 upon TPX2 binding. Phosphorylated upon DNA damage, probably by ATM or ATR.

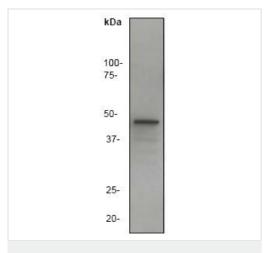
Ubiquitinated by CHFR, leading to its degradation by the proteasome (By similarity). Ubiquitinated by the anaphase-promoting complex (APC), leading to its degradation by the proteasome.

### **Cellular localization**

Cytoplasm > cytoskeleton > centrosome. Cytoplasm > cytoskeleton > spindle pole. Detected at the neurite hillock in developing neurons (By similarity). Localizes on centrosomes in interphase

cells and at each spindle pole in mitosis.

#### **Images**



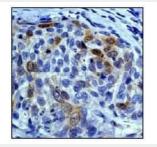
Western blot - Anti-Aurora A antibody [EP1008Y] (ab52973)

Anti-Aurora A antibody [EP1008Y] (ab52973) at 1/50000 dilution + HepG2 (Human liver hepatocellular carcinoma cell line) cell lysate at 10  $\mu g$ 

#### Secondary

Goat anti-rabbit HRP at 1/2000 dilution

**Predicted band size:** 46 kDa **Observed band size:** 46 kDa

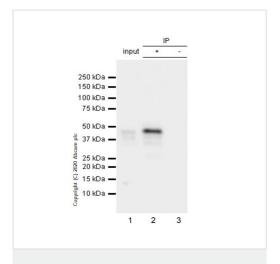


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Aurora A antibody
[EP1008Y] (ab52973)

Formalin-fixed, paraffin-embedded human cervical carcinoma tissue stained for Aurora A with ab52973 (1/100 dilution) in immunohistochemical analysis. Inset panel is a larger magnification of the image.

Inset panel is a larger magnification of the image.

Perform heat mediated antigen retrieval before commencing with IHC staining protocol.



Immunoprecipitation - Anti-Aurora A antibody [EP1008Y] (ab52973)

Purified ab52973 at 1/50 dilution ( $2\mu g$ ) immunoprecipitating Aurora A in HepG2 whole cell lysate.

Lane 1 (input): HepG2 (Human hepatocellular carcinoma epithelial cell) whole cell lysate 10µg

Lane 2 (+): ab52973 + HepG2 whole cell lysate.

Lane 3 (-): Rabbit monoclonal  $\lg G$  (<u>ab172730</u>) instead of ab52973 in HepG2 whole cell lysate.

VeriBlot for IP Detection Reagent (HRP) (<u>ab131366</u>) (1/1000 dilution) was used for Western blotting.

Blocking Buffer and concentration: 5% NFDM/TBST.

Diluting buffer and concentration: 5% NFDM/TBST.

Observed band size: 46 kDa



Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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