

# Datasheet for ABIN7299472 anti-Aurora A antibody (C-Term)





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Quantity:	100 μL	
Target:	Aurora A (AURKA)	
Binding Specificity:	C-Term	
Reactivity:	Human, Mouse, Rat, Pig, Monkey	
Host:	Rabbit	
Clonality:	Polyclonal	
Conjugate:	This Aurora A antibody is un-conjugated	
Application:	Western Blotting (WB), Immunohistochemistry (IHC), Immunofluorescence (IF),	
	Immunochromatography (IC)	
Product Details		
Immunogen:	KLH-conjugated synthetic peptide encompassing a sequence within the C-term region of	
	human Aurora A.	
Specificity:	Recognizes endogenous levels of Aurora A protein.	
Characteristics:	Rabbit polyclonal antibody to Aurora A	
Purification:	The antibody was purified by immunogen affinity chromatography.	
Target Details		
Target:	Aurora A (AURKA)	
Alternative Name:	Aurora A (AURKA Products)	

Target Details		
Background:	AIK, AIRK1, ARK1, AURA, AYK1, BTAK, IAK1, STK15, STK6, Aurora kinase A, Aurora 2, Aurora/IPL1-related kinase 1, ARK-1, Aurora-related kinase 1, hARK1, Breast tumor-amplified kinase, Serine/threonine-protein kinase 15, Serine/threonine-protein kinase 6, Serine/threonine-protein kinase aurora-A	
Gene ID:	6790, 20878	
UniProt:	O14965, P97477, P59241	
Pathways:	Cell Division Cycle, Asymmetric Protein Localization	
Application Details		
Application Notes:	WB (1:500 - 1:1000), IH (1:100 - 1:200), IF/IC (1:100 - 1:500)	
Restrictions:	For Research Use only	
Handling		
Format:	Liquid	
Buffer:	Liquid in 0.42 % Potassium phosphate, 0.87 % Sodium chloride, pH 7.3, 30 % glycerol, and 0.01 % sodium azide.	
Preservative:	Sodium azide	

should be handled by trained staff only.

-20 °C

12 months

This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which

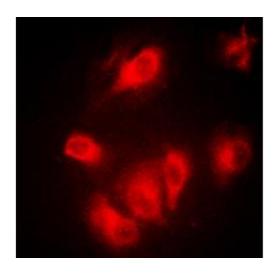
Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid freeze/thaw cycles.

Precaution of Use:

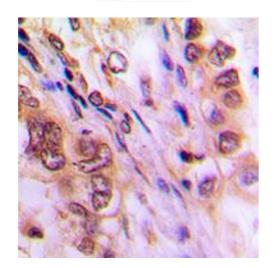
Storage Comment:

Storage:

Expiry Date:



# kDa A B C 200 140 100 80 60 50 40 30 20



### **Immunofluorescence**

Image 1. Immunofluorescent analysis of Aurora A staining in MCF7 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

### **Western Blotting**

Image 2. Western blot analysis of Aurora A expression in MCF7 (A), SP2/0 (B), H9C2 (C) whole cell lysates.

## **Immunohistochemistry**

**Image 3.** Immunohistochemical analysis of Aurora A staining in human lung cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.