# antibodies -online.com





# anti-Adenosine A2a Receptor antibody (Internal Region)





Go to Product page

$\sim$				
( )\	10	K\ / I	01	A /
1 1	ve.	I \/ I	-1	/\/

Quantity:	100 μL
Target:	Adenosine A2a Receptor (ADORA2A)
Binding Specificity:	Internal Region
Reactivity:	Human, Mouse, Rat
Host:	Rabbit
Clonality:	Polyclonal
Application:	Western Blotting (WB), ELISA, Immunohistochemistry (IHC), Immunofluorescence (IF), Immunocytochemistry (ICC)
Product Details	

Immunogen:	A synthesized peptide derived from human ADORA2A, corresponding to a region within the internal amino acids.
Isotype:	IgG
Specificity:	ADORA2A Antibody detects endogenous levels of total ADORA2A.
Purification:	The antiserum was purified by peptide affinity chromatography using SulfoLink <sup>TM</sup> Coupling Resin (Thermo Fisher Scientific).

## Target Details

Target:	Adenosine A2a Receptor (ADORA2A)	
Alternative Name:	ADORA2A (ADORA2A Products)	
Background:	Description: Receptor for adenosine. The activity of this receptor is mediated by G proteins	

## **Target Details**

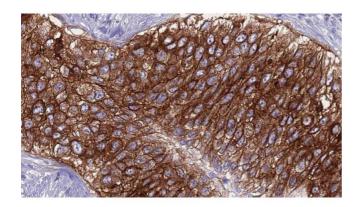
	which activate adenylyl cyclase.  Gene: ADORA2A	
Molecular Weight:	37 kDa	
Gene ID:	135	
UniProt:	P29274	
Pathways:	Neurotrophin Signaling Pathway, cAMP Metabolic Process, Synaptic Membrane, Feeding Behaviour, Cancer Immune Checkpoints	

## Application Details

Application Notes:	WB 1:500-1:1000, IHC: 1:50-1:200, IF/ICC 1:100-1:500, ELISA(peptide) 1:20000-1:40000
Restrictions:	For Research Use only

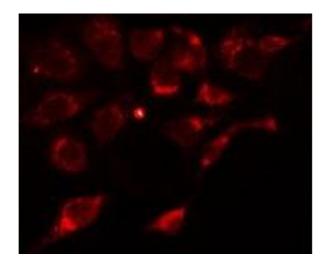
# Handling

Format:	Liquid
Concentration:	1 mg/mL
Buffer:	Rabbit IgG in phosphate buffered saline , pH 7.4, 150 mM NaCl, 0.02 % sodium azide and 50 % glycerol.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C
Storage Comment:	Store at -20 °C. Stable for 12 months from date of receipt.
Expiry Date:	12 months



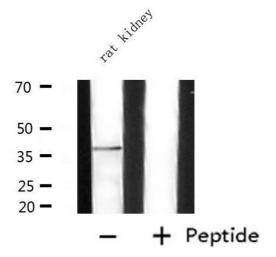
#### **Immunohistochemistry**

**Image 1.** ABIN6275913 at 1/100 staining Human urothelial cancer tissue by IHC-P. The sample was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The sample was then blocked and incubated with the antibody for 1.5 hours at 22¡ãC. An HRP conjugated goat anti-rabbit antibody was used as the secondary



#### Immunofluorescence (fixed cells)

Image 2. ABIN6275913 staining Hela cells by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25¡ãC. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at 37¡ãC. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) antibody(Cat.# S0006), diluted at 1/600, was used as secondary antibod



#### **Western Blotting**

**Image 3.** Western blot analysis of extracts from rat kidney, using ADORA2A Antibody.