# antibodies -online.com

anti-IB2 antibody (C-Term)







# **Images**



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Overview	
Quantity:	100 μL
Target:	IB2 (JIP-2)
Binding Specificity:	C-Term
Reactivity:	Human, Rat, Mouse, Zebrafish (Danio rerio)
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This IB2 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 JIP2.
Specificity:	Recognizes endogenous levels of JIP2 protein.
Characteristics:	Rabbit polyclonal antibody to JIP2
Purification:	The antibody was purified by immunogen affinity chromatography.

## **Target Details**

Target:	IB2 (JIP-2)
Alternative Name:	JIP2 (JIP-2 Products)

### **Target Details**

Precaution of Use:

Storage Comment:

Storage:

Expiry Date:

l arget Details	
Background:	IB2, JIP2, PRKM8IPL, C-Jun-amino-terminal kinase-interacting protein 2, JIP-2, JNK-interacting protein 2, Islet-brain-2, IB-2, JNK MAP kinase scaffold protein 2, Mitogen-activated protein kinase 8-interacting protein 2
Gene ID:	23542, 60597
UniProt:	Q13387, Q9ERE9
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

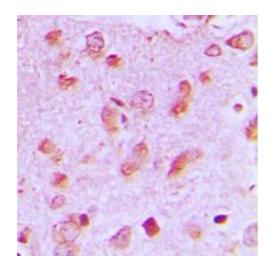
should be handled by trained staff only.

-20 °C

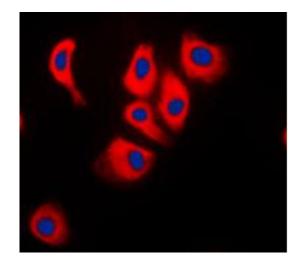
12 months

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.



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



### **Immunohistochemistry**

Image 1. Immunohistochemical analysis of JIP2 staining in human brain 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.

### **Western Blotting**

**Image 2.** Western blot analysis of JIP2 expression in Raw264.7 (A), mouse brain (B) whole cell lysates.

### Immunofluorescence

Image 3. Immunofluorescent analysis of JIP2 staining in Raw264.7 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).