

## **Product Data Sheet**

# **Anti-JIP1 (Phospho-T103) Antibody**

Catalog # Source Reactivity Applications

CPA6510 Rabbit H, M, R WB, IH, IF/IC

**Description** Rabbit polyclonal antibody to JIP1 (Phospho-T103)

Immunogen KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding

T103 of human JIP1 protein. The exact sequence is proprietary.

**Purification** The antibody was purified by immunogen affinity chromatography.

**Specificity** Recognizes endogenous levels of JIP1 protein only when phosphorylated at T103.

**Clonality** Polyclonal

Conjugation

Form Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,

and 0.01% sodium azide.

Dilution WB (1/500 - 1/1000), IH (1/50 - 1/200), IF/IC (1/50 - 1/200)

Gene Symbol MAPK8IP1

Alternative Names IB1; JIP1; PRKM8IP; C-Jun-amino-terminal kinase-interacting protein 1; JIP-1;

JNK-interacting protein 1; Islet-brain 1; IB-1; JNK MAP kinase scaffold protein 1;

Mitogen-activated protein kinase 8-interacting protein 1

**Entrez Gene** 9479 (Human); 19099 (Mouse); 116457 (Rat)

SwissProt Q9UQF2 (Human); Q9WVI9 (Mouse); Q9R237 (Rat)

Storage/Stability Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid

freeze/thaw cycles.

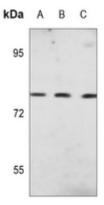
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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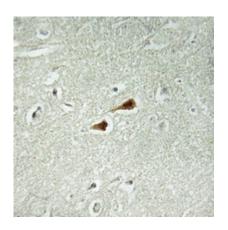
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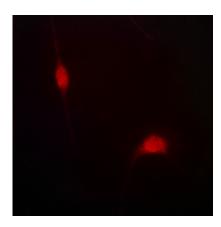
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Western blot analysis of JIP1 (Phospho-T103) expression in mouse brain (A), mouse kidney (B), rat brain (C) whole cell lysates. (Predicted band size: 77 kD; Observed band size: 77 kD)



Immunohistochemical analysis of JIP1 (Phospho-T103) 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.



Immunofluorescent analysis of JIP1 (Phospho-T103) staining in NIH3T3 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 hidified chamber. Cells were washed with PBST and incubated with a AF594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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