

## **Product Data Sheet**

# Anti-RhoGDI alpha (Phospho-S174) Antibody

Catalog # Source Reactivity Applications

CPA4567 Rabbit H, Mk WB, IH

Description Rabbit polyclonal antibody to RhoGDI alpha (Phospho-S174)

Immunogen KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding

S174 of human RhoGDI alpha protein. The exact sequence is proprietary.

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

**Specificity** Recognizes endogenous levels of RhoGDI alpha protein only when phosphorylated at

S174.

**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 (1/100 - 1/200)

Gene Symbol ARHGDIA

Alternative Names GDIA1; Rho GDP-dissociation inhibitor 1; Rho GDI 1; Rho-GDI alpha

Entrez Gene 396 (Human)

SwissProt P52565 (Human)

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

freeze/thaw cycles.

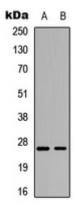
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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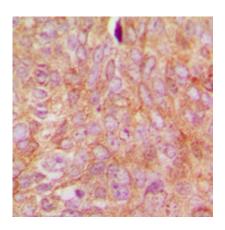
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Western blot analysis of RhoGDI alpha (Phospho-S174) expression in HeLa (A), Jurkat (B) whole cell lysates. (Predicted band size: 23 kD; Observed band size: 26 kD)



Immunohistochemical analysis of RhoGDI alpha (Phospho-S174) staining in human breast 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.

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