

Product Data Sheet

Anti-GIT1 Antibody

Catalog # Source Reactivity Applications

CPA2635 Rabbit H, M WB, IH

Description Rabbit polyclonal antibody to GIT1

Immunogen KLH-conjugated synthetic peptide encompassing a sequence within the C-term

region of human GIT1. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of GIT1 protein.

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 - 1/100)

Gene Symbol GIT1

Alternative Names ARF GTPase-activating protein GIT1; ARF GAP GIT1; Cool-associated and

tyrosine-phosphorylated protein 1; CAT-1; CAT1; G protein-coupled receptor

kinase-interactor 1; GRK-interacting protein 1

Entrez Gene 28964 (Human); 216963 (Mouse)

SwissProt Q9Y2X7 (Human); Q68FF6 (Mouse)

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

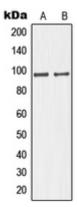
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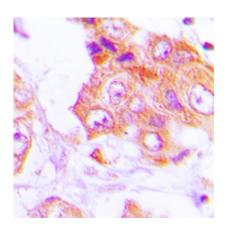




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Western blot analysis of GIT1 expression in HuvEc (A), HeLa (B) whole cell lysates. (Predicted band size: 84 kD; Observed band size: 95 kD)



Immunohistochemical analysis of GIT1 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.

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