

Product Data Sheet

Anti-GRP75 Antibody

Catalog # Source Reactivity Applications

CPA1556 Rabbit H, M, R WB, IH, IF/IC

Description Rabbit polyclonal antibody to GRP75

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

region of human GRP75. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of GRP75 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 (1/100 - 1/200), IF/IC (1/100 - 1/500)

Gene Symbol HSPA9

Alternative Names HSPA9; GRP75; HSPA9B; Stress-70 protein mitochondrial; 75 kDa glucose-regulated

protein; GRP-75; Heat shock 70 kDa protein 9; Mortalin; MOT; Peptide-binding

protein 74; PBP74

Entrez Gene 3313 (Human); 15526 (Mouse); 291671 (Rat)

SwissProt P38646, Q8N1C8 (Human); P38647 (Mouse); P48721 (Rat)

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

freeze/thaw cycles.

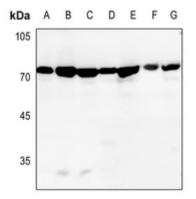
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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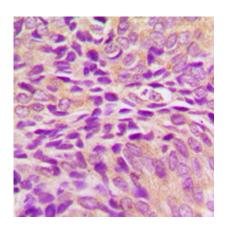
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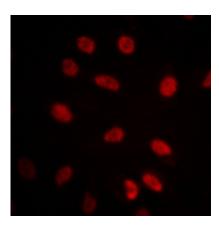
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Western blot analysis of GRP75 expression in HEK293T (A), Jurkat (B), MCF7 (C), mouse brain (D), mouse liver (E), rat brain (F), rat liver (G) whole cell lysates. (Predicted band size: 73 kD; Observed band size: 75 kD)



Immunohistochemical analysis of GRP75 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.



Immunofluorescent analysis of GRP75 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 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.

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