

Product Data Sheet

Anti-Connexin 43 (Phospho-S368) Antibody

Catalog # Source Reactivity Applications

CPA3961 Rabbit H, M, R WB, IH

Description Rabbit polyclonal antibody to Connexin 43 (Phospho-S368)

Immunogen KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding

S368 of human Connexin 43 protein. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of Connexin 43 protein only when phosphorylated at

S368.

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 GJA1

Alternative Names GJAL; Gap junction alpha-1 protein; Connexin-43; Cx43; Gap junction 43 kDa heart

protein

Entrez Gene 2697 (Human); 14609 (Mouse); 24392 (Rat)

SwissProt P17302 (Human); P23242 (Mouse); P08050 (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

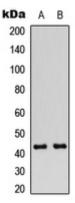
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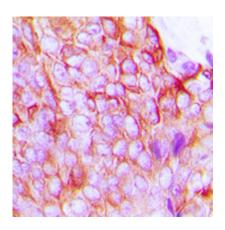




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Western blot analysis of Connexin 43 (Phospho-S368) expression in Jurkat (A), A549 (B) whole cell lysates. (Predicted band size: 43 kD; Observed band size: 43 kD)



Immunohistochemical analysis of Connexin 43 (Phospho-S368) 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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