

Product Data Sheet

Anti-WNK1 Antibody

Catalog # Source Reactivity Applications

CPA4313 Rabbit H, M, R, P WB, IH

Description Rabbit polyclonal antibody to WNK1

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

region of human WNK1. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of WNK1 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)

Gene Symbol WNK1

Alternative Names HSN2; KDP; KIAA0344; PRKWNK1; Serine/threonine-protein kinase WNK1;

Erythrocyte 65 kDa protein; p65; Kinase deficient protein; Protein kinase

lysine-deficient 1; Protein kinase with no lysine 1; hWNK1

Entrez Gene 65125 (Human); 232341 (Mouse); 116477 (Rat)

SwissProt Q9H4A3 (Human); P83741 (Mouse); Q9JIH7 (Rat)

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

freeze/thaw cycles.

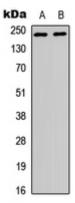
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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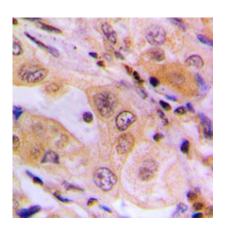
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Western blot analysis of WNK1 expression in HEK293T (A), HeLa (B) whole cell lysates. (Predicted band size: 250 kD; Observed band size: 230 kD)



Immunohistochemical analysis of WNK1 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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