

Product Data Sheet

Anti-ABL1/2 Antibody

Catalog # Source Reactivity Applications

CPA1005 Rabbit H, M, R WB, IH

Description Rabbit polyclonal antibody to ABL1/2

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

region of human ABL1/2. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of ABL1/2 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 ABL1; ABL2

Alternative Names ABL1; ABL; JTK7; Tyrosine-protein kinase ABL1; Abelson murine leukemia viral

oncogene homolog 1; Abelson tyrosine-protein kinase 1; Proto-oncogene c-Abl;

p150; ABL2; ABLL; ARG; Abelson tyrosine-protein kinase 2; Abelson murine leukemia

viral oncogene homolog 2; Abelson-related gene protein; Tyrosine-protein kinase

ARG

Entrez Gene 25, 27 (Human); 11350 (Mouse)

SwissProt P00519, P42684 (Human); P00520, Q4JIM5 (Mouse)

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

freeze/thaw cycles.

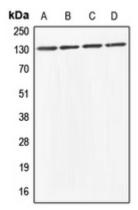
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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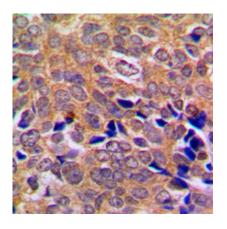
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Western blot analysis of ABL1/2 expression in HeLa (A), HEK293T (B), Raw264.7 (C), PC12 (D) whole cell lysates. (Predicted band size: 122; 128 kD; Observed band size: 135 kD)



Immunohistochemical analysis of ABL1/2 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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