

Anti-ABL1 (Phospho-Y204) Antibody

Catalog #	Source	Reactivity	Applications
CPA4326	Rabbit	H, M, R, Mk, Z	WB, IH
Description	Rabbit polyclonal antibody to ABL1 (Phospho-Y204)		
Immunogen	KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding Y204 of human ABL1 protein. The exact sequence is proprietary.		
Purification	The antibody was purified by immunogen affinity chromatography.		
Specificity	Recognizes endogenous levels of ABL1 protein only when phosphorylated at Y204.		
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		
Alternative Names	ABL; JTK7; Tyrosine-protein kinase ABL1; Abelson murine leukemia viral oncogene homolog 1; Abelson tyrosine-protein kinase 1; Proto-oncogene c-Abl; p150		
Entrez Gene	25 (Human); 11350 (Mouse)		
SwissProt	P00519 (Human); P00520 (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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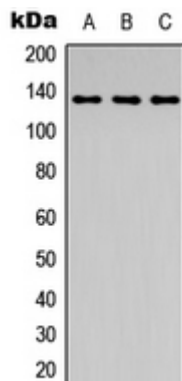
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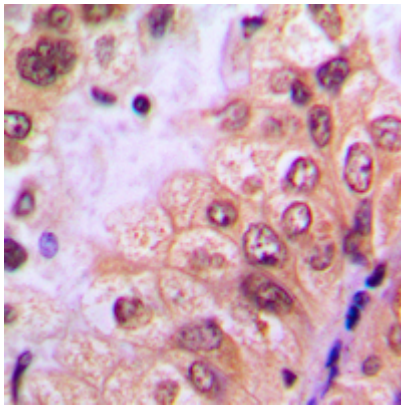
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Product Data Sheet



Western blot analysis of ABL1 (Phospho-Y204) expression in HeLa colchicine-treated (A), mouse kidney (B), PC12 colchicine-treated (C) whole cell lysates. (Predicted band size: 122 kD; Observed band size: 135 kD)



Immunohistochemical analysis of ABL1 (Phospho-Y204) 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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