

Product Data Sheet

Anti-Paxillin Antibody

| Catalog # | Source | Reactivity | Applications | |
|----------------|---------|---|--|--|
| CPA4185 | Rabbit | Н, Р | WB, IH, IF/IC | |
| Description | R | abbit polyclonal antibody | to Paxillin | |
| Immunogen | K | LH-conjugated synthetic | peptide encompassing a sequence within the N-term | |
| | re | egion of human Paxillin. T | he exact sequence is proprietary. | |
| Purification | T | he antibody was purified | by immunogen affinity chromatography. | |
| Specificity | R | ecognizes endogenous le | vels of Paxillin protein. | |
| Clonality | P | olyclonal | | |
| Form | Li | Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, | | |
| | a | nd 0.01% sodium azide. | | |
| Dilution | W | /B (1/500 - 1/1000), IH (1/1 | 100 - 1/200), IF/IC (1/100 - 1/500) | |
| Gene Symbol | P | XN | | |
| Alternative Na | ames Pa | axillin | | |
| Entrez Gene | 58 | 829 (Human); 19303 (Mo | use); 360820 (Rat) | |
| SwissProt | P | 49023 (Human); Q8VI36 | (Mouse); Q66H76 (Rat) | |
| Storage/Stabi | lity Sl | hipped at 4°C. Upon deliv | ery aliquot and store at -20°C for one year. Avoid | |
| | fr | eeze/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

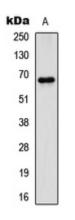
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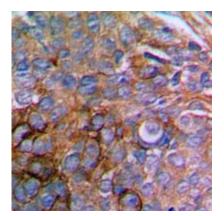


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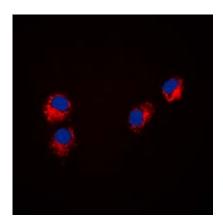
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Western blot analysis of Paxillin expression in HeLa (A) whole cell lysates.



Immunohistochemical analysis of Paxillin staining in human prostate 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 conjugad compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of Paxillin staining in Hela 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 hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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