

Product Data Sheet

Anti-CD109 Antibody

| Catalog # | Source | e Reactivity | Applications | |
|----------------|--------|---|---|--|
| CPA6024 | Rabbit | | WB, IH | |
| Description | | Rabbit polyclonal antibody | to CD109 | |
| Immunogen | | KLH-conjugated synthetic p | eptide encompassing a sequence within the center | |
| | | region of human CD109. Th | e exact sequence is proprietary. | |
| Purification | | The antibody was purified l | by immunogen affinity chromatography. | |
| Specificity | | Recognizes endogenous lev | els of CD109 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, | /50 - 1/200) | |
| Gene Symbol | | CD109 | | |
| Alternative Na | ames | CPAMD7; CD109 antigen; 1 | 50 kDa TGF-beta-1-binding protein; C3 and PZP-like | |
| | | alpha-2-macroglobulin don | nain-containing protein 7; Platelet-specific Gov antigen; | |
| | | p180; r150; CD109 | | |
| Entrez Gene | | 135228 (Human) | | |
| SwissProt | | Q6YHK3 (Human) | | |
| Storage/Stabi | lity | Shipped at 4°C. Upon delive | ery 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

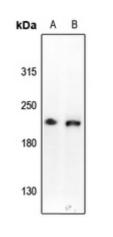
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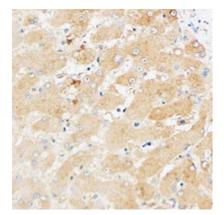
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Western blot analysis of CD109 expression in HepG2 (A), A549 (B) whole cell lysates. (Predicted band size: 161 kD; Observed band size: 200 kD)



Immunohistochemical analysis of CD109 staining in human liver 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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