

Product Data Sheet

Anti-CADM1 Antibody

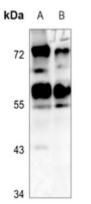
it H, M, R, B WB, IH, IF/IC Rabbit polyclonal antibody to CADM1			
KLH-conjugated synthetic peptide encompassing a sequence within the C-term			
region of human CADM1. The exact sequence is proprietary.			
The antibody was purified by immunogen affinity chromatography.			
Recognizes endogenous levels of CADM1 protein.			
Polyclonal			
Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
WB (1/500 - 1/1000), IH (1/50 - 1/100), IF/IC (1/50 - 1/200)			
oulin			
1;			
Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid			

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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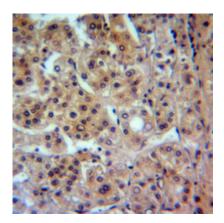
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For research purposes only, not for human use

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Western blot analysis of CADM1 expression in mouse brain (A), rat brain (B) whole cell lysates. (Predicted band size: 48 kD; Observed band size: 67 kD)



Immunohistochemical analysis of CADM1 staining in human liver 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.



Immunofluorescent analysis of CADM1 staining in NIH-3T3 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 AF594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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