For research purposes only, not for human use

Product Data Sheet

Anti-ADRM1 Antibody

Catalog #	Source	Reactivity	Applications
CQA3016	Rabbit	H, M, R	WB, IH, IF/IC

Description	Rabbit polyclonal antibody to ADRM1
Immunogen	Recombinant full length protein of human ADRM1
Purification	The antibody was purified by immunogen affinity chromatography.
Specificity	Recognizes endogenous levels of ADRM1 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 - 1/2000), IH (1/50 - 1/200), IF/IC (1/50 - 1/200)
Gene Symbol	ADRM1
Alternative Names	GP110; Proteasomal ubiquitin receptor ADRM1; 110 kDa cell membrane glycoprotein; Gp110; Adhesion-regulating molecule 1; ARM-1; Proteasome regulatory particle non-ATPase 13; hRpn13; Rpn13 homolog
Entrez Gene	11047 (Human); 56436 (Mouse); 65138 (Rat)
SwissProt	Q16186 (Human); Q9JKV1 (Mouse); Q9JMB5 (Rat)
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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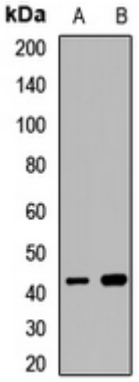
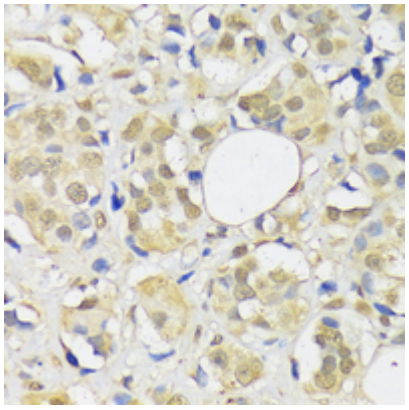
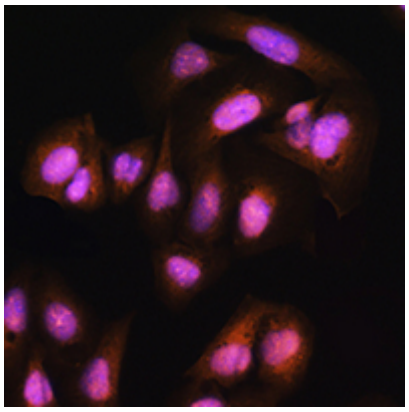


Western blot analysis of ADRM1 expression in Hela (A), MCF7 (B) whole cell lysates. (Predicted band size: 42 kD; Observed band size: 42 kD)



Immunohistochemical analysis of ADRM1 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.



Immunofluorescent analysis of ADRM1 staining in U2OS 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 humidified chamber. Cells were washed with PBST and incubated with a AF594-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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