

Product Data Sheet

Anti-VAP-B/C Antibody

Catalog # Source Reactivity Applications

CQA1199 Rabbit H, M, R WB, IH, IF/IC

Description Rabbit polyclonal antibody to VAP-B/C

Immunogen Recombinant full length protein of human VAP-B/C

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of VAP-B/C 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 VAPB

Alternative Names Vesicle-associated membrane protein-associated protein B/C; VAMP-B/VAMP-C;

VAMP-associated protein B/C; VAP-B/VAP-C

Entrez Gene 9217 (Human); 60431 (Rat)

SwissProt 095292 (Human); Q9QY76 (Mouse); Q9Z269 (Rat)

Storage/Stability Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid

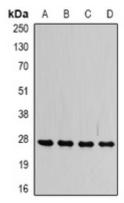
freeze/thaw cycles.

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

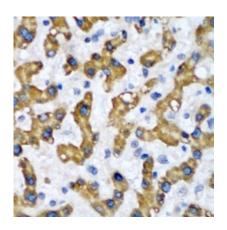
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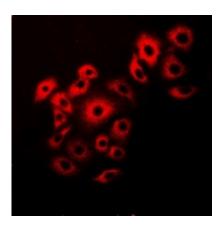
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Western blot analysis of VAP-B/C expression in SW480 (A), HepG2 (B), Raji (C), mouse brain (D) whole cell lysates. (Predicted band size: 11; 27 kD; Observed band size: 27 kD)



Immunohistochemical analysis of VAP-B/C 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 VAP-B/C 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.

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